UNITED STATES OF AMERICA CENTERS FOR DISEASE CONTROL

NATIONAL INSTITUTE FOR OCCUPATIONAL SAFETY AND HEALTH

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ADVISORY BOARD ON RADIATION AND WORKER HEALTH

69th MEETING

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WEDNESDAY, MAY 19, 2010

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The meeting convened at 8:15 a.m., Eastern Daylight Savings Time, in the Crowne Plaza Hotel, 300 3rd Street Niagara Falls, NY, James M. Melius, Chairman, presiding.

PRESENT:

JAMES M. MELIUS, Chairman
HENRY ANDERSON, Member
JOSIE BEACH, Member
BRADLEY P. CLAWSON, Member
R. WILLIAM FIELD, Member
MICHAEL H. GIBSON, Member
MARK GRIFFON, Member
RICHARD LEMEN, Member
JAMES E. LOCKEY, Member
WANDA I. MUNN, Member
JOHN W. POSTON, SR., Member
ROBERT W. PRESLEY, Member
DAVID B. RICHARDSON, Member*
GENEVIEVE S. ROESSLER, Member

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COURT REPORTERS AND TRANSCRIBERS 1323 RHODE ISLAND AVE., N.W. WASHINGTON, D.C. 20005-3701 PRESENT: (continued)

PHILLIP SCHOFIELD, Member

PAUL L. ZIEMER, Member*

TED KATZ, Designated Federal Official

REGISTERED AND/OR PUBLIC COMMENT PARTICIPANTS

ADAMS, NANCY, NIOSH Contractor

ALLEN, DAVID, DCAS

ALLAN, DONALD

BEYERLEIN, TOM, Dayton Daily News*

BONSIGNORE, ANTOINETTE, Linde Petitioner

BRADFORD, SHANNON, DCAS

BREYER, LAURIE, DCAS

BUCHALSKI, IAN

BUDREWICZ, VICTORIA

BURGOS, ZAIDA, NIOSH Contractor

CINELLI, JUDITH

CIVILETTO, SAM

CORBETT, MEGAN, Congressman Higgins' Office

COLUCCI, EUGENE

CRUZ, RUBEN, CDC

DYSTER, PAUL, Mayor of Niagara Falls

EVASKOVICH, ANDREW, LANL Petitioner

FITZGERALD, JOE, SC&A

FRANCO, TINO, Bethlehem Steel Petitioner

FRATELLO, MELISSA, Senator Kirsten

Gillibrand's Office

GIRARDO, MARY, Hooker Electrochemical Petitioner

GLOVER, SAM, DCAS

GREELEY, BILL, Congressman Higgins' Office

HINNEFELD, STUART, DCAS

HOWELL, EMILY, HHS

HINNEFELD, STU, DCAS

JACQUEZ-ORTIZ, MICHELE, Senator Tom Udall's Office

KERN, CATHY

KOHRER, FRED

KOTSCH, JEFFREY, DOL

REGISTERED AND/OR PUBLIC COMMENT PARTICIPANTS

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MCFEE, MATT, ORAU Team

MCGOLERICK, ROBERT, HHS

MENDOLA-HAUG, NANCY

MEYERS, CODY

MIDDLEBROOKS, LARRY

MONTE, LAURA, Senator Schumer's Office

MORT, DORATHEA

NOONAN, FRAN

NOONAN, KAREN MORTENSEN

NOWICKI, RICK

OSTROW, STEVE, SC&A

PRESLEY, LOUISE

RAMSPOTT, JOHN

RYKIEL, SANDY

ROLFES, MARK, DCAS

RUTHERFORD, LAVON, DCAS

SARDINA, JOSEPH

SCREMMIN, P.

SHAFFER, KATHLEEN

SWEENEY, THERESA

SWIFT, JACKIE

TORNABENE, ELEANOR

ULSH, BRANT, DCAS

VENTURA, MARGARET

VENTURA, SAM

WADE, LEW, DCAS

WALKER, JOYCE

WERNER, JIM

WHIPPLE, STEVE

WITRYOL, AMY

*Participating via telephone

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1 P-R-O-C-E-E-D-I-N-G-S 2 (8:26 a.m.)3 WELCOME Why don't we 4 CHAIRMAN MELIUS: 5 get started. Ted, do you want to give your 6 speech? 7 MR. KATZ: Sure. I don't have a long speech. 8 CHAIRMAN MELIUS: The phone 9 10 speech. Yes, I'll do the phone 11 MR. KATZ: bit, but first of all let me just say welcome 12 13 to everyone who is here and welcome to everyone who is on the line on behalf as well 14 of the Secretary of Health and Human Services 15 16 and the Director of NIOSH, we are happy to be here in western New York and we have a very 17 full agenda. 18 19 Let me just note for people on the 20 line, the agenda has two public comment sessions, one at the end of the day today at 21 22 4:30, beginning at 4:30,

tomorrow

and

beginning at 6:00 p.m. And let me also note some of the agenda items may move around a bit as we get things done sooner than we expect or they take longer. But the SEC petitions that are listed on the agenda, the start times for those are fairly much time-certain, just so you are aware of that.

Okay, and then let me just -several things to note. For people listening
on the phone, please mute your phones except
when you are addressing the group and to mute
your phone, if you don't have a mute button,
use *6, and then *6 again, pressing it again,
will un-mute your phone when you do want to
speak to the group. And please do not put the
call on hold at any point. Hang up and dial
back in. Hold will disrupt the call and the
meeting.

And then let me just note also for the public comment sessions, people listening, I'm not sure I see any people, local people here right now, but I'll say it now and repeat

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it later. There is a sign-in outside if you wish to address the Board in one of the public comment sessions, outside on a table, so please sign in, and for the folks on the line, we'll just take you as we go after the people have commented locally.

And last thing just to mention for Board members, we have a lot of SECs. A number of these, some Board members have conflicts. Just keep that in mind as we get to those sessions. And that's it, thank you.

CHAIRMAN MELIUS: And we will get started. Just to note for the record we have a full complement of our Board members with Paul Ziemer and David Richardson on the telephone, and Paul, David, I will try to remember all the time to recognize you when we are having discussions and so forth, but feel free to speak up if I do forget.

With that we will start with the NIOSH program update, Stu Hinnefeld.

NIOSH PROGRAM UPDATE AND PROGRAM EVALUATION

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MR. HINNEFELD: Good morning, everyone. I'm here to present our customary program update status and a little bit about things that are going on in the program, keep everybody abreast.

A few news items to cover before we get into our statistics. Between the last meeting in February and now, the Office of Compensation Analysis and Support's name was Division changed the οf Compensation This Analysis Support. essentially and removed our organization from the office of the director, and gave it division status similar to other divisions within NIOSH. Uр to this, the Office is a somewhat smaller, I guess, political designation than a division, and it just made I believe the Institute's org chart look a little cleaner not having this huge office of the Director. That is my own personal opinion on why they did it.

In the phone call after this became effective, Dr. Howard congratulated me

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on my elevation to division director, and so I went back and increased the setting on my office chair by a couple of inches so I was elevated, because I have noticed no other difference associated with this name change, except that we spent a lot of effort changing stationery and things like that.

Also in March of this year the Government Accountability Office issued its latest report about the EEOICPA program. It addressed the activities of both NIOSH and the Department of Labor. We mainly read those things with an eye toward NIOSH in what they say. They had no recommendations in their report for NIOSH, so we had no recommendations to take under advisement and decide what kind of actions we could take in response.

They commented - essentially what they said was dose reconstructions take a long time and costs a fair amount of money, but Congress told NIOSH to do dose reconstructions, and that's what it takes to

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do dose reconstructions; it's a difficult technical process. So that is essentially paraphrasing what they said about the NIOSH part. They did make a few recommendations to the Department of Labor. I'm not privy to anything that may follow from those.

And finally I want to comment very quickly about a data review that was done very -- on a very short turn-around within the last -- well, actually the data, the documents were looked at last week Thursday. We identified, contractor identified or our fairly recently, about two or three weeks ago, a finding aide from the National Archives and Records Administration facility, essentially a federal records warehouse, essentially, in College Park, Maryland, there finding aide classified of some was information that included quite a long list of sites that are covered in our program, and one of the names on the list was Chapman Valve. Chapman Valve is on the agenda for this

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meeting, and so we wanted to see what did that
tell us about Chapman Valve before this
meeting. And so with the cooperation of the
Department of Energy and participation by
NARA, we did manage to have one of our
staffers one of our dose reconstruction
staffers and one of the staffers from Sanford
Cohen & Associates to get there to see the
documents in the box that were associated with
Chapman Valve. And Joe Fitzgerald, who is
here at the meeting, was one of the people who
attended. Mark Rolfes, who is at the meeting
although I don't know if he is in the room
right now was our staffer who attended. And
they concluded that the information about
Chapman Valve didn't really inform us any more
than we already are: didn't provide any
explanation to that two-percent uranium
sample. It didn't give any more information
about the work that was done there in terms of
the machining of the natural uranium that we
already know about. So there was nothing new

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in that. So at least we did manage to get there. And the Department of Energy helped us out a lot there in getting clearances verified and over to NARA so that people could go look at those documents.

We will be making additional reviews back to that. We weren't able, of course, to look at all the documents we were interested in from all the sites on that finding aide during that one-day visit before this meeting. But we wanted to make sure we got the Chapman Valve documents reviewed.

Now our statistics, these numbers climb up a little bit every time. We are now up to almost 32,000 cases -- this is as of April 30 -- that have been sent to us for dose reconstruction. Some 88 percent of those have been returned to DOL either with the completed dose reconstruction or through -- because they were pulled by the Department of Labor. Of those pulled by the Department of Labor, about 2,550 of them were pulled for SEC Classes that

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had been added by our processes. Those were not the statutory SECs, but SECs that had been added through the Board's recommendation, and the Secretary's designation.

So that leaves some 3,200 that still have to do for dose we reconstruction, and then there are about 605 that were administratively closed. generally happens when the claimant declines to sign the OCAS-1 form and return that to us. They can be reopened at any time that the provides the OCAS-1 if claimant or provide us additional information that is relevant to the dose reconstruction. Theoretically, that might cause it to reopened as well.

This is just the same information about current case status in a pie chart. The -- it's reassuring I guess to see a big chunk that is completed, but -- and then some of these other smaller chunks are also things that we are essentially completed with,

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anything that's pulled.

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So now of those 3,266 cases that are still out for dose reconstruction, several dose of them are in the reconstruction process; some of the newer ones, we're still in development, getting all the case information together that we need in order to do the dose reconstruction. And then there are 555 where the draft dose reconstruction is in the hands of the claimants. In other completed the draft words, we have reconstruction, and the claimant now received that draft dose reconstruction and the OCAS-1 form for them to either say, okay I I have no more information to provide and send the OCAS-1 back, or for them to say, no wait, I have more information that might be relevant to dose reconstruction and we want you to look at that. So there are some 555 of those that we expect, a fair portion of those we expect we are probably done with as well, expect for the paperwork of sending the --

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getting the form back and then sending the final dose reconstruction.

One of the categories, broad categories we have of cases are what we call pended cases, and that means there is some reason why we need additional information before we can complete that claim, either additional research on our part or information from some other agency.

The top five categories of the pends are listed here. The majority, you see this is over the half of the cases that are pended are pended because they are associated with SECs that have not yet had a final designation and that would probably include cases that we are recommending today, or at this meeting, not necessarily today but at this meeting, but would include Classes that we are recommending to the Board during this meeting.

MR. KATZ: Excuse me, there is someone on the line who doesn't have their

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phone muted, and there is sort of a scrabbling sound coming through the line. Thank you.

*6, if you don't have a mute bottom.

HINNEFELD: So then we have MR. COI issue, or close-out interview issues, and those are instances where the claimant has the draft dose reconstruction in their hands, they have the OCAS-1, they say, wait a minute, here is some additional information we think is And so they raise that to us, and then we have to go investigate that additional information. Sometimes it's information, maybe it's additional medical information which then has to be verified by the go Department of Labor. There various are categories that goes into.

The category non-SEC, pending DR methodology, that is for Classes, cases that are from sites where there is an SEC but these don't qualify usually because they don't have one of the listed cancers. And so we have to determine what it is we can do. We have

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reached a determination that we can't do every bit of the radiation dose; we can't do all the radiation dose in the reconstruction. want to and so once we qet to that conclusion we will bring the recommendation forward in order to get those SEC Classes And on occasion we still need to do moving. some additional research to figure out, well, what is it we can do. What is the entirety of the doses we can do in order to do this partial, to do as much of a partial dose reconstructions for the non-SEC cancers as we can?

We have a number of cases where we have made additional data requests to DOE, something for data beyond just the individual exposure record that we receive at first request. And then there are some 24 where a Technical Basis Document issue is something we need to finalize the approach on where dose, for that some type of dose We need additional research reconstruction.

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on how we can do that.

Here is the breakdown of the cases that have been sent back in terms of their PoC score, and about 31 percent of the cases where dose reconstructions have been done have PoC greater than 50 percent, and a little over 16,000, 16 and a half thousand cases had a PoC less than 50 percent. So that is about 31 percent success on SEC -- on dose reconstruction.

Recall though that there were some 2,500 cases, 2,550 cases that had been pulled and returned to DOL for SEC which appeared to be compensable through SECs. So that actually makes some 10,000 cases that were sent to us for dose reconstruction. It looks like they will have a compensable outcome when you combine the dose reconstructions above 50 percent and the SEC numbers.

This is a chart that we've shown for a long time. It shows the distribution of cases according to their Probability of

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Causation score. It's skewed a little bit at the right side because everything greater than 50 percent is lumped together in one large bar as opposed to all of the others which are just 10-percent intervals of dose reconstruction. You can see there are quite a large number at very low dose reconstructions, and then it kind of, you have a valley until you get up to the 50, where I said it's sort of artificially goes back up.

couple of meetings For а we presented a graph that we intended to show the improving timeliness of cases as the program has matured, and the original slide we showed on this was flawed because it was sort of incomplete. We would run these dose reconstruction groupings in groups of 5,000. These numbers across the bottom are the NIOSH tracking numbers for claims, and those are assigned chronologically. So the first that came in was assigned case number 1, and we are now assigning cases up close to 32,000,

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think, as they come in. And what the average presented before was time complete a dose reconstruction for each block, for the 5,000 cases in one through 5,000, and then for the 5,000 cases in 5,001 to 10,000, and the number went down. The average dose reconstruction time went down. It was pointed out in a couple of meetings that that data was incomplete because for any dose reconstruction in any of those populations that was not yet done, it's not yet included in the average, so that will have a higher number than what you have, and so you are representing sort of artificially low numbers there.

So in order to give a complete view of the dataset, we went back, we decided, well, let's see how we are doing in completing our initial drafts within one year of the time we received the claim, which is of course our current management objective, which is to get cases done within a year from when we get them. And for that reason we dropped off the

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25,001 to 30,000, because at the end of that period, the claims at the end of that period are not yet a year old. So again we would have incomplete data on the dataset. So we dropped off the last grouping of 5,000. And this shows the improving timeliness, or the improving percentage of claims that we are able to complete in a timely manner. And we would expect not necessarily for 25,001 to 30,000, but for the next grouping beyond that, we would expect that number to be very close to 100 percent of those getting done within one year.

Just a little more statistics. This is DOE response to requests for exposure records. We get good service out of the Department of Energy. We still get about 200 new claims a month, almost. Over the last 12 months we've averaged somewhere around 190 new claims from DOL a month, people who have newly filed in the program. We send those, if they are at a site that has exposure records, we

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send those to DOE. DOE responds very well.

We have a few stragglers. We work

consistently with DOE to try to make sure we

get those cleaned up and back to us.

Special Exposure Cohort status: we've had 171 petitions we have received. But when this slide was made up on May 6th we didn't have any undergoing qualification review. I think that might be different today almost two weeks have gone by, and we may have gotten a Form BN since then.

A hundred and three have been qualified for evaluation. We have six under evaluation. I believe this next is eight, I believe we are presenting eight today, or at this meeting? Bomber is nodding his head, so we are presenting eight SEC Evaluation Reports at this meeting. We have 16 that have been presented or are under discussion. You all know that we oftentimes will have technical discussions about the logic of it following recommendations, and those go on with the Work

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Groups and the Board's contractor.

One petition with the Secretary awaiting final decision on May $6^{\rm th}$, I believe that final decision was sent since May $6^{\rm th}$, and that was the Canoga facility which the Board voted to recommend at their phone call at the end of March.

and then the 61 petitions have resulted in additions of Classes, 56 Classes. The reason for that is petitions sometimes come in covering the same time period or a very similar time period, and we will combine petitions, and so evaluate them essentially with two, and with a single class. Five have been denied; 63 did not qualify; and there are 5 that we did include in the 63 that were received before the rule became final. I mean those people were allowed to re-petition once there were rules for petitioning.

Of the 56 Classes that have been added since May of 2005, they split exactly equally between 83.13 process and the 83.14

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process. For anyone who doesn't know, the dot 13 process is a petition received by a petitioner and interested party on their own, says, I don't think you can do my dose reconstruction. The 83.14 process is when we are researching a site, and we arrive at the conclusion from our research that we don't have enough information here to complete all aspects of the dose, and so we initiate those largely on our own.

And it represents, these come from 40 different sites, and represent over 2,500 people.

I spoke a little bit earlier about our management objective to improve timeliness. This slide has been up here a few times. We adopted this almost one year ago, trying to get to the point where claims would be done within a year of when they were sent to us. We had some claims that were getting pretty old. We issued that objective last June with the objective of being in place by

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this June, which is of course about two weeks And we originally published the policy to address initials, in other words when a case first came to us from the Department of Labor, and we expanded the population during the course of the year to say, why don't we just try to get any case that comes whether it's an initial one, or one that has come back to us for rework because perhaps the medical information changed or something. Let's try to get them all so they are not here more than a year.

Any of them that we don't get done, the policy calls for us to critically evaluate the relevant obstacles and write a memo to the claim file that recommends the best way to proceed.

And there was a lot of work done in the last year to try to accomplish this. I'm not 100 percent sure of these items, but I know there were a lot of 83.14 SEC petitions that have been prepared in the past year. I

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think the Board will recognize for about the last three meetings we have had a very heavy agenda in terms of SEC petitions; a large part of that is a result of this effort to finish up the research for sites that have not yet been researched, and reach conclusions on whether dose reconstruction is feasible or not.

have also a large number of Technical Basis Documents to revise, incorporate the information we needed. As we completed research there were cases where we figured out, okay, well, we do have enough information to do dose reconstruction, and so would write the technical document we supporting it, summarizing the research we And with that time period we had an made. average of about 70 dose reconstructions per week for the year. And we had to continue to qualify and evaluate new 83.13 SEC petitions, and we received 11 of those since June of last So that work didn't stop, that had to year.

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continue. And we have provided some support to the Advisory Board, Subcommittees and Work Groups, although I think you will recognize that may have suffered a little bit because of all the emphasis on these other things.

In terms of how we are doing over time, this is for the initial cases, it shows the decrease in those numbers. This is the reworks, and then this is the combined, so I'll talk about this a little bit.

Today we are down to about 450 or 430 that remaining number, since 31st, so we are down quite a lot more since then. You all recognize that that is an awful lot to get done in a few weeks. As a matter of fact we are not going to get every one of those done. There are certain categories of claims that will not be done by the end of June. For instance, we are presenting today several 83.14 SEC recommendations in that we have completed our research, we don't feel the dose reconstruction is feasible, and

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we are going to bring a recommendation to add those Classes.

Those claims will not be done by June 1st. First of all, the process for getting them, the process of the SEC takes 2 to 2-1/2 months after the Board's recommendation before those claims -- before it's actually effective, and so those won't be done.

And similarly we haven't done the non-presumptive, the non-SEC cases, because we don't feel like we have a formal finding that the dose reconstruction isn't feasible.

I think I actually have a couple of slides that talk about this. Here are some SEC Classes that we know won't be done. GE Evendale, which you know we are researching, we are doing additional research in to try to define the size of the class. These remaining five are being presented at this Board meeting in terms of our recommendation to add -- a recommendation essentially to add, to amend

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the Class Definition that was provided that was added earlier.

And there are a handful of claims that we cannot complete because we need additional information from either DOE or DOL.

The DOL are questions about employment; the DOE ones are usually supplementary Requests for Additional Information. I don't believe any of those are the initial requests that we made for the individual exposure records. But at times we will make supplementary requests that are more difficult to fulfill, and there is some number of those.

And then there are a handful of cases where theoretically we could, and in fact this first claim, this 83.13 petitioners where feasibility of dose reconstruction is under consideration. This slide is a couple of weeks old, and in the meantime, that one should be scratched off. We are proceeding with those dose reconstructions, but we are going to put a modified letter on the dose

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reconstruction. We are going to send them their standard letter. The purpose of the modified letter is to reassure that particular claimant or petitioner that the fact that we are doing a dose reconstruction and we are sending a draft dose reconstruction does not mean that we have decided definitely that be reconstructed. their dose can petition is still being evaluated; it may in fact turn out not to be feasible. But we are idea of giving them where the dose an reconstruction would come out under the dose reconstruction process. But don't worry that this has no effect on the deliberations on petition, and we are still open your SEC mindedly approaching those discussions and it may in fact be true that their petition may be successful.

We have just a few sites, this amounts to some 20-some odd claims, that use surrogate data, and we are evaluating our uses of surrogate data as it pertains to these

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sites against our own criterion IG-004. And those evaluations aren't yet complete. And so we have elected not to send these dose reconstructions yet until we are comfortable with those uses of surrogate data based on our own criteria.

recognize the Board is doing their own deliberation on that. At that time when understanding reach а common whether surrogate is acceptable and what type is acceptable, it looks like we may in fact have some revisiting of our decision. decisions are what we can do now with our criteria. Of course any Board advice will be taken as it normally is when we receive it, and may modify work that was done earlier just as other Board decisions do on occasion, too.

And there was one claim that was returned to us because of unresolved findings on Site Profile that was not yet resolved. I believe it's the Savannah River Site Profile. It was, we thought we had completed dose

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reconstruction. The petitioner asked DOL to reopen it based on the findings in the Site Profile. They did and returned it to us. Those findings are not yet resolved, and so we've not yet gone ahead with the dose reconstruction.

If we were to redo the dose reconstruction today; we have not made any changes to the Site Profile, it would be the same dose reconstruction as the fire one.

So there are -- other than those of claims associated with those categories is on the order of 300. We really wanted to get categories it appears we'll be successful in meeting the June 1st target. The total number done. We recognized probably a few months ago that we would be able to get all the SEC Classes recommended -- researched, recommended by the February meeting which would have been the meeting where it had to get done for those cases to get done by June 1st. So we were not to able do that; we recognized that

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months ago. But other than these categories of exclusion for those reasons and for waiting for data for others we feel like we have at least done a creditable job of getting those cases done. All those have a pathway, all those 300 claims. Their pathway is determined, when we get the additional information from the Agency and you finish it, you get the Board's recommendation one way or the other.

Ιf declines the Board add to we have recommended, then we would have to do some sort of I guess additional research to do the dose reconstruction. But That would take a little while. so be it. the Board agrees with us and recommends those Classes, those cases will then go according to the prescribed pathway. So we think that all those can be done, and their pathway is set, so the letters to the file will essentially say that the obstacles remaining, there don't really seem to be any obstacles remaining.

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1	It's just a matter of process to get through
2	this, essentially.
3	Okay, that's the end of my slide
4	show. Does anybody have any questions?
5	Okay, if there are no question
6	oh, I'm sorry. I was trying.
7	CHAIRMAN MELIUS: Oh, Wanda.
8	MEMBER MUNN: This is a simple
9	process question more than anything else.
10	These status reports for some of us are a
11	crucial part of accumulation of information
12	that we maintain for the Board. My current
13	electronic file does not include that
14	presentation. I'm assuming that the
15	presentation will be added to our file for
16	this meeting?
17	MR. HINNEFELD: So what you are
18	saying is you didn't get this presentation
19	electronically?
20	MEMBER MUNN: That is correct, I
21	did not get it.
22	MR. HINNEFELD: That's news to

1	me.
2	MEMBER ZIEMER: Dr. Melius?
3	CHAIRMAN MELIUS: Yes.
4	MEMBER ZIEMER: Paul Ziemer here.
5	Let me just verify the same thing that Wanda
6	said, because I was looking for the file in
7	the we have a Niagara Falls list of
8	documents that Zaida provided, and I just
9	emailed Zaida while we were talking and Zaida
10	didn't have that available. So I suspect it
11	can be easily transmitted to all the Board
12	members.
13	MR. HINNEFELD: We'll make sure
14	we'll send this to all the Board members.
15	MEMBER ZIEMER: And we may have a
16	similar issue on the upcoming presentations,
17	as far as the PowerPoints are concerned.
18	MR. HINNEFELD: None of the
19	PowerPoints got to the members?
20	CHAIRMAN MELIUS: It's funny
21	because Jim Melius it's on the one I
22	have. I have the following is Stu

1	MEMBER ZIEMER: Is it on the O:
2	drive?
3	CHAIRMAN MELIUS: Not on the O:
4	drive, on the stick. Okay, I see it. Okay.
5	MEMBER ZIEMER: What is it under?
6	CHAIRMAN MELIUS: Mine is on a
7	separate file, sorry.
8	MR. KATZ: Nancy is saying, Paul,
9	it's under NIOSH updates on the O: drive.
10	MEMBER ZIEMER: I am looking at
11	the Niagara Falls section of the O: drive and
12	there is nothing called NIOSH updates.
13	MEMBER MUNN: Some of you may
14	have a later version of Niagara Falls.
15	MR. HINNEFELD: It's not a
16	problem. We can put it on the O: drive and
17	that would be no problem in transmitting this
18	via email. And we may do both, to make sure
19	all the Board members have it. And I
20	apologize for it; I thought it was there. I
21	thought it would be on the stick.
22	MEMBER MUNN: We don't get a

stick.

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MR. HINNEFELD: You don't get a stick anymore? It just goes on the O: drive?

Okay, I thought it would be there. I thought it would be wherever it was supposed to be.

CHAIRMAN MELIUS: Any other questions?

MR. HINNEFELD: Okay, now one thing I did not mention was the program review which is being done of our program by the Office of the Director, Director of NIOSH, not my office, and Lew Wade wants to say just a couple of words about that.

DR. Thank you. WADE: Good morning, Board members. I will be very brief and give you a brief update on the status of the 10-year program review. As you know, the review was going to focus on five topical Those areas are the quality of science areas. practiced by the program, the timing accomplishment of NIOSH's program tasks, the individual activities, the dose SEC

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reconstruction activities, and then the services provided to claimants and petitioners.

The review will take place in two The first phase is to focus on data phases. and facts surrounding those five topical That phase one review will conclude by observations and comments by the authors. me briefly remind you of the authors of these various pieces. The quality of science piece will be offered by Professor Henry Spitz of the University of Cincinnati and Doug Daniels of NIOSH's staff. The timing piece by Nancy The SEC piece by Randy Rabinowitz who Adams. is with us here. The dose reconstruction piece, I'll author. And then the claimant and petitioner quality of service will be done by Denise Brock, Nancy Adams a team of myself.

Once that phase one report is complete, then there will be a phase two. That's where John Howard, the NIOSH director,

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and a group of senior NIOSH leaders will look at those phase one reports, and formulate thoughts as to modifications of the program, be they policy, legislative modifications, again, based upon the findings of fact.

What I've shared with you now, I hope, I know it's in your book, is a preliminary draft of the phase one report that deals with individual dose reconstructions. Again it's dated May 12th and should be in your book.

The reason why I've given you that is really threefold. One is it lays out a format that we intend to follow in the phase one reports. Secondly, it uses Board work. I refer to Board work products in this document, and therefore thought it appropriate that I get it to you as quickly as possible.

I also wanted to talk to you briefly about the topics that will be explored in that document. And let me read them for you briefly. Again, you should have this in

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your book.

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The first topic explored in the DR piece is the Advisory Board's review of completed dose reconstructions. Secondly, we are looking at the Advisory Board's review of Site Profiles and procedures used to accomplish individual dose reconstructions. Third, statistics concerning the number and time complete individual dose to reconstructions. Fourth, statistics concerning the number and time to complete individual dose reconstructions as evaluated by dose reconstruction types, dose estimation Fifth, statistics concerning types. reason for, and time to complete number, reconstructions. partial dose Six, the dose reconstructions that percent of resulted in PoCs greater than 50 percent. last, individual dose reconstruction compensation results based on the cancer model used.

Now in the document I've given you

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COURT REPORTERS AND TRANSCRIBERS 1323 RHODE ISLAND AVE., N.W. WASHINGTON, D.C. 20005-3701 I've written up all but two of those sections.

I don't have the data yet to complete the dose estimation technique used or the partials. But I wanted to get this to you for you to look at and consider and comment upon either as individual Board members, as you choose, or as a Board as a whole, particularly since I'm making use of Board work products here. I think it is important that you have a number of opportunities to comment upon them.

The schedule that I see unfolding at your August meeting I would intend to have the complete draft of the phase one report to you prior to that meeting, so you could look at it and react to it. Once you have a chance to react to it and the document is modified, then John Howard will start his review from a policy point of view.

I will be available on your phone call on the $14^{\rm th}$ of August -- $14^{\rm th}$ of July, I'm sorry -- if the Board has any comments that we would like to make collectively as a Board at

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1	that point.
2	So again, comments by the Board as
3	you would like, individual comments from Board
4	members any time you would like. That's all I
5	really have.
6	CHAIRMAN MELIUS: Questions or
7	comments for Lew. Josie.
8	MEMBER BEACH: Just a quick
9	question. You kept saying, it's available in
10	our book. And I guess I need to know what
11	book you are referring to.
12	CHAIRMAN MELIUS: It's the O:
13	drive.
14	MEMBER BEACH: What is it listed
15	under?
16	MR. KATZ: So it apparently
17	they are trying to sort this out out there. I
18	don't know exactly what the situation is.
19	MEMBER ZIEMER: This is Ziemer.
20	I can shed some light on that. This was
21	distributed by Zaida just a few days ago. The
22	email is dated May 11 th . It's under Zaida's

1	email address but is from Dr. Wade. And it
2	has two documents attached, the one document
3	is entitled, 10 Year Review of the NIOSH
4	Radiation Dose Evaluation Program, Phase I
5	Report. And the second document is called 10
6	Year Review Phase I Dose Reconstruction.
7	Those are just attachments to the email. And
8	I'm looking at this distribution. It went to
9	every Board member and was sent on 5/11.
10	DR. WADE: And there are hard
11	copies on the table. So when you have an
12	opportunity take a look at that and comments
13	as you would like.
14	CHAIRMAN MELIUS: I for one
15	didn't get that. So I guess we have some
16	distribution problems here.
17	MEMBER ZIEMER: Dr. Melius,
18	Ziemer again. I notice used your CDC address
19	on this. So that sometimes is an issue still.
20	MEMBER RICHARDSON: This is David
21	Richardson. I had a question about the
22	dosimetry, dose reconstruction report that you

shared, this is in response to an area of focus which is described here as the appropriateness and consistency of individual dose reconstructions. Is that right?

DR. WADE: Correct.

MEMBER RICHARDSON: I mean I just read the report quickly. But it seemed to me that there was a lot of information in the organization report under the current headings that were related to the timing of dose reconstructions, tables so one, two, three, four, maybe, are kind of -- it seemed that they are overlapping with some of the other topics that are in that NIOSH review, topic two, the timing accomplishment, topic five, the timing of service provided. And I was wondering whether that needed to be there, or whether the focus could be reoriented a little bit toward issues of consistency of dose reconstruction. That was the sort of information I was looking for there.

DR. WADE: Thank you. The

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approach we are taking now is to let each of the authors assemble their document completely and fully as they would like, realizing that there is likely to be overlaps, and then deal with those overlaps on editing of the full document. So you will find overlaps in the drafts as we Hopefully those edits will -- those forward. overlaps will be removed through the editing process.

MEMBER RICHARDSON: And one other question I had was, I don't tend to think about evaluations of dose reconstruction as being tied into Probability of Causation determinations. Ι think that those largely a function of the risk estimate that you associate with the doses. And I was wondering how that fit in again to this issue of the quality of dose reconstruction and its consistency.

DR. WADE: I understand your point of view. I do think that to track the

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issue completely it's, in my opinion, worth looking at what the dose reconstructions bring about in terms of Probability of Causation and compensation decision to see if there is anything in there that piques one's curiosity. But I do understand the boundary you are building, and at least at this point I'm inclined to include it in an early draft and then possibly remove it if appropriate.

MEMBER RICHARDSON: Okay.

CHAIRMAN MELIUS: Thanks, David.

Ted has an O: drive update.

MR. KATZ: I just had an update on the computer-access issue. I think if you are in the O: drive right now and you go up a level and then back into it, you will find documents now, and it's -- I gather what happened is that there were no controls put on that O: drive folder. It was put out there so that any Board member that moved it or did anything to the folder, it disappeared for everybody, including -- they had to track it

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1	down and find where it was. But in the future
2	they'll lock that down so it can't go
3	wandering.
4	CHAIRMAN MELIUS: Wanda stole the
5	folder.
6	(Laughter.)
7	MEMBER MUNN: Never even found
8	the folder.
9	(Laughter.)
10	MEMBER MUNN: It's there now.
11	CHAIRMAN MELIUS: It's hiding
12	from you. It heard you coming.
13	MR. KATZ: So all those documents
14	now should be there for you if you are in the
15	O: drive.
16	MEMBER MUNN: Thank you.
17	CHAIRMAN MELIUS: Can I suggest
18	that, since we are just finding the
19	information Lew was referring to the
20	documents, let's take a look at them in the
21	next couple of days. Maybe we can come back
22	and talk, but maybe the way to go, given the

timing, would be that we provide individual comments between now and the July call, and then we can put on the agenda for the July call a time to discuss those, and if there are common comments we want to make as a Board, that would be helpful. Maybe if you'd copy me in your comments to Lew then I will try to see if there are common themes or common issues that we want to address as a Board. But I tend to think the individual comments would be the best way to go. Think about it once you look at the document, once you find it and look at it, and then we'll gather information and then we can decide.

Any other comments or questions on that? If not, we'll move on to DOL program update.

DOL PROGRAM UPDATE

MR. KOTSCH: Good morning. I have to apologize up front. I managed to get through an entire winter without picking up a cold or flu, and now this past weekend we had

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some pretty wild temperature oscillations back in D.C. and somehow I picked up something.

This is the standard update that I present every time. And there are only a few people I think that probably haven't seen this at least five or six times. So we will proceed semi-quickly through it. This is the of background the Energy Employees Occupational Illness Compensation Program Act. There are two pieces to it, or two parts, Part B, which is the part that is mostly of interest here because of the NIOSH reconstructions. That became effective on July 31st, 2001. As of May 6^{th} , 2010, we've had filed 70,599 cases or 105,761 claims. I always mention, the number of claims always higher because cases with survivors generally will have one or more survivors. it's greater than a one-to-one relationship; 31,931 cases have been referred to NIOSH for dose reconstruction.

The other part of the program is

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the Part E program, the toxic chemical portion of the program, which was effective for the Department of Labor on October 28, 2004. There have been 61,917 cases filed. That's 87,691 claims. And initially when the program started 25,000 cases were transferred over from Department of Energy.

The overall breakdown as far as compensation as of again May 6^{th} is \$5.7 billion in total compensation, breaking down to be \$3.29 billion for Part B, \$1.95 billion for Part E, and \$488 million for medical benefits that are associated with the awards.

And then this is just the breakdown of the paid cases under the Act.

There have been 43,000, almost 44,000 Part B and E cases, 40,000 -- almost 500 Part E payees and roughly 26,500 cases. And roughly 18,500 Part E Payees, and about 17,500 cases.

So the breakdown is roughly 40 percent Part E, 60 percent Part B.

Again, just quickly, for Part B it

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covers or addresses radiation-induced cancers, chronic beryllium disease and beryllium sensitivity; silicosis for the miners at the Nevada test site and Amchitka in Alaska. And the supplement for the RECA Section 5 uranium workers, which the RECA portion is adjudicated by the Department of Justice. That is Radiation Exposure Compensation Act.

Again this is just the eligibility under Part B. DOE employees, the feds, the DOE contractors and subcontractors. The Atomic Weapons Employers, beryllium vendors, the survivors that are listed there, and again the RECA Section 5 uranium workers. Again, just the presumptive cancers, the SECs, the four legislative statutory sites that were in the Act, the three gaseous diffusion plants, Smith, Paducah, 825, plus the Amchitka test site.

And the benefits, the \$150,000 lump sum payment; the medical benefits for covered conditions; and the medical treatment

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and monitoring only for beryllium sensitivity.

CBD and cancers are covered in the upper part.

And the breakdown of the final decisions as of May 6th, 28,214 final decisions approved; 21,140 final decisions denied. And the bars to the right on the right side, the bulk of the denials resulting from about 14,900 Probability of Causations less than 50, and about 5,700 where there was insufficient medical evidence.

Part E, just quickly, it's a federal entitlement program, like Part B, lump sum payments, up to \$250,000, usually on top of the Part B payment, if there is a cancer. Plus medical benefits for the accepted conditions.

Part E is a little bit different.

A couple of the things are, it only applies to the DOE contractors and subcontractors, not the AWE folks or the beryllium vendors. Survivors, there is some differences in the survivors. For the deceased workers, you can

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compare those. But it does deal in Part E with any occupational disease from any toxic exposure. That includes again the Part B people.

It addresses in Part E impairment, the percentages of award for permanent whole body impairment is \$2,500 per point. It addresses in Part E wage loss, and there you see the distribution of those. Again based on decreased capacity to work.

And just in summary, the Part E cases, the final decisions, 23,504 approved; 19,394 denied, and again, actually the bulk of those is 13,342, is medical information insufficient to support the claim.

Getting back to the NIOSH -referrals to NIOSH, we are indicating as of
May 6th 31,931 cases referred to NIOSH for
dose reconstruction. That breaks down to
27,783 cases returned by NIOSH that are
currently at DOL, 24,100 roughly of those had
dose reconstructions, and the rest did not

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have dose reconstructions. They were probably returned for SEC Classes and things like that, or insufficient information to proceed with the adjudication.

We are showing 4,148 cases at NIOSH which breaks down to about 2,750 for initial referrals and about 1,400 for reworks.

Cases that generally back to NIOSH for -- after the initial dose reconstruction generally for, the biggest reasons are additional employment or evidence of additional cancers.

As far as the new SEC-related cases about -- well, not about, 2,887 cases were withdrawn from NIOSH for review; 2,544 have had final decisions issues; and 2,480 of those final approvals; 43 have have finals; recommended but no 91 cases pending at Labor; and 209 cases were closed because of problems with information.

Again a quick overview of the process at DOL, a recommended decision is

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rendered at the district-office level; that goes to the claimant, if it's obviously not a compensable case, the claimant has the opportunity to provide objections or additional information. Then the final adjudication board within Labor renders final decision based on a review of that information, and renders what's up there as the final decision or the final approval.

Status of the dose reconstruction is 24,102 have been returned by NIOSH with a dose reconstruction; 21,540 cases have a dose reconstruction and a final decision. Again you see basically the breakdown as far as from the Labor statistics: 66 percent final denials; 34 percent final approvals.

Part B, cancer cases with final decisions to accept. There have been 6,872 dose-reconstructed cases that have been \$1 billion accepted, resulting in in compensation. There have been 10,684 accepted SEC cases, which resulted in \$1.6 billion in

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compensation. Cases accepted both on SEC status and Probability of Causation 50 percent or greater were 386, or \$57.4 million in compensation. Which totals out for all accepted SEC and DR cases, dose-reconstructed cases, 17,942 or \$2.6 billion in compensation.

The next series are just monthly Part B cases sent to NIOSH, running about, I guess in the upper 200s. Nationwide the new Part B cases that the DOL receives monthly which include obviously the beryllium and the silicosis as well as the cancer has ticked up a little bit. I think that may be a result, as we'll see later, may be a result of some of the new SEC, say at Hanford or something, that has prompted some additional submittals of claims.

Last couple of months, 552 in March, April, 480. Just a quick note about the top four worksites that are generating Part B claims or cases for us. Hanford, Y-12, Savannah River site, Oak Ridge Diffusion

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plant, K-25. Again there's numbers for those. Like I said I think we are seeing that uptick for Hanford I think as a result of the Hanford Class, the SEC Classes, and the people just responding to that. Y-12 is pretty much probably level; these are just normal fluctuations. Savannah River, those are those numbers. And K-25.

These are just the percentages, this and the next slide are the percentages of the new Part B cases. These are the DOE cases received monthly by the Department of Labor by percentage. And it usually runs about the low 90s, the split between that and this slide which is the AWEs, which are generally running in the upper single digits.

And then these, I'm not going to go through all these, the next few sets of slides are the information that we provide for both the SEC Classes that are being presented at this meeting just as far as background information goes, and some of the local

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facilities, I mean facilities that are local to Niagara Falls-Buffalo area. So that's what the next few slides are. You see here, Mound, just a few quick ones, 1,712 cases, both Part B and E for Mound. There have been 484 cases returned by NIOSH with a dose reconstruction; 665 final Part B decisions; 246 Part B approvals; 270 Part E approvals for total compensation and medical bills paid, \$56.5 million.

As you go across, you see University of Rochester. You see since that is an AWE it's not affected by or not covered under Part E, so there's only Part B numbers there. Same with BWX Technologies, Hooker Electrochemical.

Just a few Linde Ceramics, \$27.4 million in compensation for 144 Part B approvals and 62 Part E approvals. There is the St. Louis Airport site, Weldon Spring, Blockson Chemical, Chapman Valve, again it's an AWE, so it's only Part B, \$6.8 million in

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compensation. Bethlehem Steel, \$61.7 million in compensation with 406 approvals. \$199 million with 855 Part B approvals, 890 Part E approvals. De Soto Avenue, Downey facility, Bliss & Laughlin, Carborundum, Linde Air Products, Ashland Oil, Seaway Industrial Park, Lake Ontario Ordnance, and Electro Metallurgical. And then just the pie chart of all the B cases files, the breakdown of those. Six percent ended up with NIOSH, 36 percent others includes in for the our space silicosis, the beryllium claims, things like that.

Now Dr. Melius had asked for a bit of an update or an overview I guess of how we implement SEC Classes at Labor. And I'll give you a quick one, and then you can ask any questions. The process starts with NIOSH submitting a letter to the Department of Labor, sharing the draft language for the new Class Definition. I said a month. It varies, the time varies, but sometimes we get quite a

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bit of lead time for the ones that are coming up essentially for scheduled for the next Monday.

And then we review that and we send it back, send comments back. The easy ones are the ones that say all employees that are, whether it's AWE or DOE, at a facility, those are fairly straightforward, and don't require a lot of head-scratching to come up with the fact that we probably don't have a problem with the Class Definition.

But the ones that do have, that cause us to review them a little more closely are the ones where there is some type of restriction based on the Class, whether it's a particular building is cited, or an area of a plant like Area 4 at Santa Susana Field Laboratory or buildings at a particular site, or in the early days, there were a lot of monitored or should have been monitored. Things like that caused us at Labor a little more, required us to provide a little more

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scrutiny. But what we often did, we'd talk to the District office that was affected by that site and say, hey, pull some cases, tell us if you can really implement that Class because of employment restrictions or that building restriction or that monitoring restriction, and can we actually do that, can we get that information?

The whole intent of this thing is to improve the consistency and the fairness of the claims adjudication process; ultimately to speed the process of determining which cases can be considered as part of the new SEC Class. But we do not at any time comment on whether the new SEC Class is indeed necessary.

There's a point on the next slide which should probably be up here, but at this point, too, not only is there management and technical review; there is also review by our solicitor's office, primarily to the point of sometimes we get into facilities question as to whether -- and I can't think of specific

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examples now. One of them was I think maybe MIT or something. Anyway, where there was a discussion of building put in that Class, and we looked at it from a facility definition standpoint, had an issue with that, and talked to NIOSH about that, got that resolved. So things like that are going on at that point in the process, trying to first of all make sure that that definition fits within the facility definitions that already exist and exist, and whether we can actually put people into that Class. And that is the whole goal when we review that draft proposal language.

this for bullet, for Okay, Classes that do not cover all the workers, DOE strives to describe as many ways as possible individuals in place the the covered buildings or areas or what have you. All cases, whether they are really an open class, like all workers, or some kind of restricted Class where they are defined more by buildings or something else, they are all done on a

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case-by-case basis. You don't always -- you always all the have employment information you need to determine either the location or the 250 day requirement or things like number of pieces that. So а information are used. Obviously all the employment information from the site reviewed. We receive information, request and receive information from the Social Security Administration. Pay records essentially are summaries. There be department may affiliation records that we can look at, or monitoring records that are often very useful. There may be medical records that give some indications of the buildings or the time period that a person has worked in. There may be security records, access records. And as always, the claimant has the option providing affidavits from other workers, coworkers, or other individuals that can place people on sites or in buildings or things like that.

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And DOL includes in the bulletin a list of records that can be used when establishing a membership in the SEC class. But again done on a case-by-case basis, and that can encompass a wide range of documents if necessary. Once the HHS letter is submitted to Congress, that is really the bell to our procedure folks to move on the final approval of the bulletin. Ιt goes through our management process, and the Energy Employees Occupational Illness Compensation the Offices of the Solicitor Division, then current office, the Office our Workers Compensation Programs.

Unlike NIOSH, offices are higher in our organization than divisions, so.

But anyway the DOL goal is to have the bulletin finalized by the SEC Class effective date, and in the early days -- well, maybe even not that early -- but we didn't always achieve that. But hopefully we will move close. Nowadays we have lists, NIOSH

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provides us with lists of cases. We generate our own lists of what we think are the effective cases, and we start to share those with our district offices during this process so they can start basically queuing up those cases so that when the gun goes off on the net effective date, we are now, at least over the last few times we've done this we have been pretty good about getting out of the blocks pretty quick.

And the last part, after the Class is officially effective, DOL is working through its review of the cases that are in the SEC Class to determine whether they meet the requirements of the class, the 250 days, have the specified cancer, if there are additional requirements of the Class meet those.

We get into the outreach phase.

If it's a small Class usually we just contact

-- if it's a few people or one person we'll

just contact the claimant by phone and discuss

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the Class with them. If it's a larger Class like the Nevada Test Sites or the Hanfords or even the Lawrence Livermores which are going to be coming up, I forget the dates, people and often with representatives NIOSH will go out and basically do townhall meetings or meetings with -- if the -- in the area of the facility to discuss the new SEC with that, class. Concurrent there are whatever efforts are made to try to get word through union newsletters out or newspapers or however our outreach program has decided is the best that way get information out.

Any questions?

CHAIRMAN MELIUS: Questions?
Yes, Mark.

MEMBER GRIFFON: Just my regular question, Jeff, on the Rocky Flats review of the Ruttenber Data, I'm hoping that you will have something to tell us at this point. It's been several meetings.

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1	MR. KOTSCH: Yes, I understand.
2	I have something to say but it's not what you
3	want to hear, unfortunately. It is still with
4	management as far as how they want to handle
5	the use of the Ruttenber data, because it
6	would be used in conjunction with the
7	information that we are already using for that
8	class.
9	MEMBER GRIFFON: And are you at
10	any point going to like report back to us on
11	just what you found in your review and why you
12	are ending up with a decision I guess once
13	you have a final decision?
14	MR. KOTSCH: Yes, once the
15	Department has decided which way they want to
16	go with the use of it or how they want to use
17	that data, then we'll provide some background
18	on the thought process.
19	MEMBER GRIFFON: Okay.
20	CHAIRMAN MELIUS: Anybody else?
21	If not, could you comment on we've been
22	going through this process recently of re-

looking at old Classes. Could you comment on sort of how this has arisen? Why are we looking at these old Classes, like some of the restricted Classes we are now talking about making them more generalized, and sort of what would be the rationale for that and the process for that?

MR. KOTSCH: I think those issues are -- I mean, NIOSH is implementing those issues. I mean we had -- and I don't know if LaVon wants to discuss it, but the things that spring to my mind are the Y-12 class, maybe Lawrence Livermore where there were some -- I don't know, do you want to talk, LaVon?

MR. RUTHERFORD: Yes, I think it's more critical from our end that -- in retrospect after looking at the claims -- I'm LaVon Rutherford by the way -- in retrospect after looking at the claims we felt that some of those claims should have ended up in the class. And when you look at that situation, if there is a question as to whether a claim

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should end up in a class, maybe we haven't defined it appropriately, so that's why we've gone back and done that.

Lawrence Livermore National Lab, that's one that we've done. We are getting ready to do Los Alamos for the same reason.

MEMBER GRIFFON: And have you looked at Rocky Flats?

MR. RUTHERFORD: Not yet. That whole process of reviewing all the Classes is really going to occur mainly after June 1 to a period of time, and I'll have a better update. But if you look at the Los Alamos one, we defined it as technical areas. And ultimately we went back, and we said, okay here's another technical area we missed, and we revised that list, and even in that process of defining all these technical areas where there potential radioactive material we still ended up with a few claims that came back to us that were concerned from our end that probably should have been included in the

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class. And from -- so it was our approach that if there are claims that are not being included in a Class that we felt they should have been, then we probably did not define the Class appropriately. And so that's why we are going back and revising some of this.

CHAIRMAN MELIUS: How do we know that that's a widespread problem? There is always going to be -- given how complicated these sites are and given some of the vagaries of personnel records and how people were assigned, there's always going to be people that may not fit. It may be hard for DOL to administer. So I'm getting two things. Sometimes it's starting with NIOSH. Sometimes it's issues of how can DOL administer it based on the records that are available to DOL. just trying understand where to different -it may be both in different situations, but how this applies, how it comes about.

MR. RUTHERFORD: I don't think

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there is a widespread problem, first of all. think the approach have with the we Department of Labor, I think we do give that opportunity to the Department of Labor front because of lessons learned early on in the process to see if they can implement the class. I think their implementing of class, even with Los Alamos and Lawrence Livermore, the ones that we have changed now, I think we learned over time that initially they felt they could implement the Class as But we didn't written, and they probably are. actually -- we probably missed some of the boundaries involved, I should say. And that is how Class claims ended up coming back to us that should have been included in the class.

CHAIRMAN MELIUS: Then who determines when you miss the boundaries what the new boundaries are? It seems like on Los Alamos you are talking about well there are a couple of technical areas you missed. And then you like sort of it becomes a very

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generalized Class that covers everybody. Hanford became the entire site. The buildings, the office buildings. And I guess I'm just trying to understand how that process There may be situations where comes about. there aren't adequate personnel records which DOL would be the one handling those and trying to make the individual determinations. then does NIOSH make their how own determinations?

Well, when we MR. RUTHERFORD: review and initially define an infeasibility, put boundaries around that we to infeasibility. We put years, and then we have look at, okay, where was the exposure potential for that infeasibility? How widespread was it? Once we define if it's limited to a building, then we determine, okay, there potentially individuals, are maintenance workers, firefighters, individuals that would have went into that building, that would have walked into that building, security

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guards that need to be defined included in that class? We try to define if we can limit that Class to only workers that were in a specific building, then we have to go to the Department of Labor and say, okay, we plan on — in fact Mound is coming up, it's going to be a lot of discussion — we say that our infeasibility is limited to this building. Can you identify workers specifically that worked in this building for 250 days, and can you administer this Class basically?

And then the Department of Labor reviews that Class Definition, reviews their information available, employment records and so on, to determine if they can administer the class. The difficulty you get into is are there other workers that were potentially from our end that may be missed in that process? So that is a collaboration we have to work with the Department of Labor and come to an agreement on that.

CHAIRMAN MELIUS: And where does

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DOE fit in? Do you -- DOL interacts with DOE, that was on one of your slides, Jeff, in terms of your setting up the Class and so forth. They apparently at least review the Class Definition.

MR. KOTSCH: I don't know if they always get the Class Definition. have issues with think we are going to employment, or putting people in buildings, or something, we will talk to Greg or somebody at DOE and try to determine whether we really have that kind of information available to us to proceed with putting people in a building or were they monitored or whatever the issue is, or can we get employment records or the completeness of those records.

So there is communications -- I won't say it happens on every class, but certainly on the ones that we have issues with or that we think we may have issues with, where we do talk to the Department of Energy.

CHAIRMAN MELIUS: So there is no

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formal -- they are not formally involved in any of these?

MR. KOTSCH: I know that sometimes -- it's more like when we are getting into facilities, like definitions of things. I know we exchange letters on those kinds of things. On this process I don't think it's as formal.

CHAIRMAN MELIUS: Okay.

MR. KOTSCH: Like I said at the beginning, when we have issues like that, we try to sample -- we ask the district office to try to sample some cases, and that is semirandom. We'd hope, just to see, kind of get some general feel for whether that information exists in the records, or is available in a record. I suppose sometimes that backfires on us.

MR. HINNEFELD: If I can offer one thing, when we go through this process of defining a Class now -- we haven't always done this -- we are looking for documentary

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evidence of people in the plants, what is the document, rather than write a Class where something has to be generated, or the claimant has to provide, the burden of proof is based on the claimant that said they spent 250 days in that building. So we want to avoid placing that burden on the claimants to have to provide additional evidence. We are looking for some sort of documentary evidence defines or includes this Class of people. if that exists then we may have restrictive Class Definition. Absent that kind of evidence that allows you to define it, we tend now to go to the larger, say we can't provide evidence of who actually was in that area or not.

CHAIRMAN MELIUS: That's helpful, Stu. I think there's been two issues where this has been raised. One is something like the General Electric site where the Class is so broad that it raises issues about equity and so forth given that particular facility.

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are going back to revising the Classes that you are doing, so it reflects the justification for that. We are suddenly generally expanding a Class pretty significantly, and how do we capture what we are doing in a report. In some of the reports you do well, and there is documentation. In others it's less clear why we're doing it. Los Alamos, at least I found, it's not quite as clear why you are doing it, making this
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as clear why you are doing it, making this
change now. And I think that is the other
thing. I'm trying to understand the process
so we sort of understand where it's coming
from and what information to look for without
making more work for DOL necessarily, or
generate more reports or anything like that.

MR. HINNEFELD: I personally don't care how much work we make for DOL. But the idea is not to put a lot of work on the claimants, and would document a record.

CHAIRMAN MELIUS: And then the

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third is, I guess I'm trying to understand where DOE could fit in. At our last meeting when we discussed it, I believe it was Regina or Pat Worthington jumped in and said that they thought that there may be a role for them to be more helpful with this also.

MR. HINNEFELD: To the extent

that they may know more about the entirety of the record set that is available, that would be I think where they would come in.

CHAIRMAN MELIUS: No, when you go to your field offices, I mean I have visited your field offices. And you had a few cases to work from, limited information. And then you had some staff that were often from the facility, so they knew the facility. But even they would not know the records for every contractor or every employee. These are very complicated facilities and it's difficult.

Paul or David, do you have any questions?

MEMBER RICHARDSON: Yes?

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1	CHAIRMAN MELIUS: Yes, hi, any
2	questions, David or Paul?
3	MEMBER RICHARDSON: No.
4	CHAIRMAN MELIUS: Okay. Well,
5	we'll move on.
6	MEMBER ZIEMER: This is Ziemer, I
7	have no questions. Sorry, I had the mute in
8	the wrong position.
9	CHAIRMAN MELIUS: Okay, I just
10	don't want to be ignoring you.
11	MEMBER ZIEMER: Thank you.
12	CHAIRMAN MELIUS: Okay, we'll
13	hear from Department of Energy.
14	DOE PROGRAM UPDATE
15	MR. LEWIS: Good morning,
16	everybody. Again, I'm Greg Lewis. I'm the
17	program manager for the EEOICPA program at the
18	Department of Energy. The EEOICPA program at
19	DOE is run out of the Office of Health and
20	Safety, and the director of the Office of
21	Health and Safety, Dr. Pat Worthington, was

going to come and present to you today but was

unable to make it last minute. So here I am.

just to address Dr. Melius' And question from Jeff's presentation, I know we have had involvement in these types of issues in terms of whether or not the records are able to place people in certain buildings or locations, or they are able to meet the needs of the various SEC Classes. I know in the past it's been requested by DOL and NIOSH, although Ι believe since the last meeting I think we've been more involved. think DOL is sending us every proposed Class Definition before it goes out and allowing us So we have done so on a few, and to comment. we have been involved.

And again at DOE our primary role in the EEOICPA program is to provide records. Our core mandate is to work on behalf of program claimants to ensure that all available worker and facility records and data are provided to DOL, NIOSH and the Advisory Board. So again we provide records.

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We have three primary responsibilities under the program; we respond to individual requests for information from DOL and NIOSH on a per claimant basis. We also provide support and assistant to DOL, NIOSH and the Advisory Board on large-scale records research projects and data retrieval at DOE sites. And the third responsibility is to conduct research in coordination with DOL and NIOSH on issues related to the covered facility list.

As far as individual claims, we do about 6,500 employment verifications per year, about 3,000 dose requests from NIOSH, and about 6,500 what we call DARs from DOL, and those are requests for exposure information, HR, IH, medical information, the various things that DOL needs to adjudicate these.

In FY 2009 we did just under 16,000 individual records requests, and in FY 2010 we anticipate the number to be about the same. And again it's somewhat difficult

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because our numbers don't always match up with the Department of Labor's and NIOSH's that's because we do it on a per-site basis, so I think typically within the DOE world many of the workers worked at two and facilities over their career, and if that is the case, we still have to do two and three searches for records, whereas it may just be one case or one claim. So the numbers don't quite always match up, but those number of our first requests that we completed.

The primary role in our program is taken by the local EEOICPA site point of These are the folks at the various contacts. active DOE sites throughout the country that manage the program at that site. They have a responsibility and really huge are backbone of our ability to provide records. These folks attend local public meetings, they set up site visits and tours for the NIOSH and DOL staff on large-scale records requests.

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They facilitate interviews for these records research projects. They provide -- locate and provide subject matter experts that can interview and talk with the various research They also provide information to the groups. current workers, so they conduct some outreach and work with some of the current workers that may be involved are claimants or in And of course then these are program. people that manage the 16,000 records requests Their response is at the site, per year. gathering the records from the various active provisions and responding to DOL and NIOSH within the 60-day timeframe.

So again our -- in addition to the individual records requests, the most work we do for large-scale records research is of course the SEC evaluations. And currently these are a number of SEC projects that we are either working on or have recently completed our supporting NIOSH's efforts. So Hanford, Mound, Savannah River, Pantex, Weldon Springs,

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BWXT, Los Alamos, Brookhaven, Nevada, Santa Susannah Area 4, and St. Louis Airport Storage Site and Linde.

I'll go through a few of these.

I'll get more into detail on a few of them,
but if there are questions at the end just
ask. So for Mound we've facilitated meetings
where members of NIOSH and the Advisory Board
have had classified discussions about Mound
activities. We have also had subject matter
experts on our end in to talk to them about
various classification concerns and how they
could articulate their various issues.

We have facilitated over 40 worker interviews, and because Mound is a closure site and we don't have an active DOE facility or location for classified interviews, we have worked with the FBI office in Dayton for use of their facilities to conduct these interviews.

We have also set up Mound document review visits at numerous locations. Our NSA

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service center in Albuquerque and College Park, Oak Ridge, Pantex, Los Alamos, Denver, and of course primarily at our Mound View facility in Dayton that is run by our Office of Legacy Management.

At Pantex we are continuing to facilitate worker interviews. There is actually an onsite visit happening this week, so they are down at Pantex. I think members of SC&A are down there doing interviews. I think somewhere around 30 just this week alone, I believe.

We are also working to set up a second site tour for members of NIOSH, the Advisory Board and their contractors, while the SC&A group is down there this week, I think they were talking with the Pantex folks about how to set up that tour, what is going to be on the agenda and the timeframe. So we will continue to work on that.

At Linde we've supported research for Linde Ceramics documents. Most of the

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Linde documents are held in the DOE Office of Legacy Management which handles the records for the closure sites or former DOE sites. LM has worked extensively with NIOSH to provide copies, and we have also supported record review visits for Linde at various federal record centers, and additional smaller sources within the DOE.

Hanford, this work is primarily completed, although I know we do continue to work with SC&A and NIOSH on various follow-up actions, but I won't go through all of these, but these are some of the efforts that we took for Hanford.

Savannah River again we -thousands of documents and hundreds of
thousands of pages were produced and reviewed
for the research effort.

Again at Brookhaven, multiple data capture visits were supported, hundreds of boxes of records. We facilitated a site subject matter experts to talk to and

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interview with the various groups, and that's similar to the process at most of the sites.

As far document reviews, as recently with the creation of our security NIOSH security plan plan and the and the Advisory Board's security plan, we have had a amount of work completing document for the various reviews source documents requested, as well as the NIOSH and SC&A and Board reports that we are reviewing. are committed to provide documents to DOL, NIOSH and the Advisory Board, but we must do so in a responsible manner to protect national security interest.

We just make sure to comply with existing DOE and NIOSH security plans. We follow those to a T, and if there are any issues, we can always go back and look at those security plans to see if they need to be changed. And our security plan is at that link, and I believe the NIOSH and Board plans are on their website as well.

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Since March of 2010, so roughly since the last Board meeting, NIOSH, the Advisory Board and the contractors have submitted 74 documents for DOE classification review.

The average turn-around time for these 74 documents was eight working days, and as always in certain cases where an expedited review is necessary we have returned documents in two to three days, when necessary.

And again just to clarify these are the reviews of documents that come to DOE headquarters. We also review documents at the field sites. We have a little bit less control over the field sites and depending on their workload, it's not necessarily eight days, especially because the documents that go to the DOE field sites are typically source documents, and where a report can be 50 to 100 pages, a source document can be five, six, 700 times, pages at SO those can be difficult. But we make every effort to return

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those documents to NIOSH, and as soon as we can in the timeframe that meets their needs.

And then third our major responsibility at DOE is to research and maintain the covered facility database, which is a database of over 300 facilities covered under EEOICPA. It includes DOE facilities, AWEs and beryllium vendors and a full listing is at that link and we again work closely with DOL and NIOSH to identify any inconsistencies additions that should be making, or we those research and make the change, necessary.

Office of Legacy Our Management supports us heavily in this records research. Legacy Management is the group within DOE that handles the records and management of closure facilities. So records are one of their primary roles within DOE. They have records experts. We currently have five staff that we work with over there that have an average οf 20 years records management

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experience within the DOE complex, so they are very knowledgeable about the history of DOE, the records history specifically within DOE, and where to go to locate records when the various requests come in from NIOSH and DOL.

Currently these are just a few of the sites we are working: GE Vallecitos, and then I selected a couple of local ones, Simonds Saw and Steel from Lockport and then recently we actually completed this one but for the Carborundum Company in Niagara Falls, we added a second distinct work location and time period of the work was changed to 43 to 44 and 59 to 67. I don't know what it was before, but I believe we added time in that case based on new information that we located through our research.

We have a number of initiatives going on. Some of these you may have seen before, but some of them are new since the last Board meeting. Again we continue to emphasize cooperation with NIOSH and DOL with

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our various sites. One of the questions that we had from the Board was, you know, they had been hearing concerns about workers and being hesitant to interview because of fear retribution or reprisal. And I think we told you at the last Board meeting we released a memo from the DOE Deputy Secretary encouraging workers to participate in these interviews. We have been continuing to make that available to workers, when they are It's available to site staff interviewing. just so they know why and what is going on with these NIOSH and DOL interviewers that are coming in and wanting to talk to workers. And this is also, these memos are available for NIOSH and DOL to hand to the workers when they are conducting the interviews as well.

We hold routine conference calls with members of NIOSH, the Board, SC&A, ORAU, the various contractors, make sure that we know what their needs are, what they are working on, and how we can support that.

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We have DOE subject matter experts that participate and contribute Advisory Board, Working Group and conference I don't know that we always do so, but calls. certainly if requested or needed we are very willing to find the right person to participate.

And as I mentioned in the SEC section we have facilitated secure meetings and videoconference calls where members of the Board, NIOSH or DOL are able to discuss classified information in a secure setting and have unencumbered discussions.

These are two projects here that we are very proud of that took a tremendous amount of effort, and have both been recently The first is a project to revise completed. the contracting provisions in the acquisition guide within DOE to ensure that DOE has a access and maintain ownership of right to records. It sounds a little confusing, but basically changing our contracting we are

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guidance so current contracts and
subcontracts, which are really the part where
we may have been lacking in the past, current
subcontracts will have or should have
provisions in them that allow DOE to maintain
ownership and management of these records when
that subcontract may be finished. I think as
many of you know subcontractors, particularly
historically, it's been very tough for DOE to
provide records. Some times when these
subcontractors, particularly smaller
construction subcontractors, leave the site
when their project is finished, they may leave
the project records behind, but their human
resource records, their worker records, often
they take with them. So particularly for
older companies or older work, this is more
historic work, those companies may have been
bought and sold numerous times. They may not
be in existence, or they may have subsequently
destroyed those records. So this change is,
we feel very important, and will allow DOE to

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maintain control of these records for future workers and future claimants.

And then the second project is the Alamos Medical Center Records project. Los This has taken almost two years, but earlier this fall we finished it up, and we took possession of the pre-64 records. This is a the medical center project where at in the early days before 1964, behind the fence in part of the site. the site opened up it was sold to a private company and they have retained those pre-64 medical records. We went in and they were with actually contaminated the various substances, mold, and potential hantavirus, things like that. We went in, decontaminated the records, organized, indexed them, and took possession of pre-64 records. And now the Medical Center has the post-64 records, which of course being a small town with primary employment at Los Alamos there are many worker records in those as well. And the Medical

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Center is committed to preserve and provide the post-64 records to DOL and NIOSH. And they are already doing so.

We've also in the last year year and a half or so taken a greater role in outreach along with DOL, NIOSH and the various We have initiated what we are calling the Joint Outreach Task Group, which includes DOE, DOL, NIOSH, DOL ombudsman and the DOE former worker medical screening programs. These are all groups that in some form or another are trying to reach roughly the same population, and many were doing individual And because of the overlap outreach efforts. of the various programs, we work with those combine outreach efforts when groups to coordinate possible and to and improve outreach. In the last year we held 18 townhall meetings near nine DOE sites, and we continue to do so. In fact last week we were at Rocky Flats and held a meeting with the So if anyone wants to see the current group.

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schedule it is up on our website at that link.

And then the one last effort I want to talk to you about is the review of the DOL Site Exposure Matrices database. Exposure completed Site DOL the Matrices database, or what they are calling SEM. Ιt contains data from all major DOE sites. focused primarily on exposure to substances, not radiological exposures, used as а tool by the DOL examiners to adjudicate claims. When it was created in 2006 through 2008 the DOE site POCs worked closely with DOL to identify and gather the records that they needed for that We provided -- similar to the SEC database. research projects, we provided access to hundreds of boxes of records, thousands pages of information, as well as site subject matter experts were available to talk to them, and provide quidance in terms of historical work as well as where the records can be found.

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Recently DOL has requested that DOE review this database for public release, and we are working hard to do that. We've already authorized the release of the SEM information on 48 of the 116 DOE facilities in the SEM, as well as all uranium mills, mines and ore-buying stations. Those were many of those the closure sites or smaller sites that could be reviewed at DOE headquarters. Now we are currently reviewing the remaining 68 sites and most of those are being reviewed by the various DOE field sites.

Again we are hoping to have this review complete by the end of the summer and as some sites more information than others and they are more difficult to review and may have more sensitivities than others, so they may be staggered when they come out. But as various sites are finished, we will releasing them so we anticipate that the sites will be released throughout the summer. the current public SEM website can be found

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And the last thing I want to talk to you about is the Former Worker Medical The mission of the former Screening program. medical offer free worker program is to screening for former DOE workers. After they separate from DOE they can get a screening three years after separation and every three years thereafter. It's a free screening. It's tailored to the work that they may have done at the site, and the things that they may have been exposed to, although certainly if we identify or catch anything else we'll make sure to let them know and refer them to the right medical care.

Further information to the Former Worker Program can be found at the link on my presentation. And the information here is about the local screening program because there is not a current local DOE site in this area. There is no local program. However, we started the National Supplemental Screening

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Program. It provides medical screening to workers that aren't in the area of a regional project, or workers that may have moved out. So we have contracted with clinics throughout the country, and can provide screening in all 50 states, including Alaska and Hawaii. And the local outreach number is on the handout as well as the website. So questions.

CHAIRMAN MELIUS: Brad.

MEMBER CLAWSON: It is good to Ι you again. like that eight-days turnaround on the documents. I think we've still got a little bit of room there. want to ask you, what are we doing with certain sites getting documentation released a little bit faster there? Have we got anything streamlined? Because we have talked numerous times, it's too long. It has taken a lot. there anything that we can do to accelerate that a little bit?

MR. LEWIS: Well, I mean I agree.
We struggle - the eight days up there is for

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DOE headquarters. And I'm right around the
corner from DOE headquarters. We have a great
relationship with the headquarters reviewing
classification staff. We have a special
liaison over there that helps us navigate
through the various problems we run into. So
I think we are very confident with what we do
at headquarters. We agree in the field it can
take a little bit longer. Again I think for
the most part in the field, our field offices
and field sites do review the information in a
reasonable amount of time and get it back to
you. However there are certainly cases where
it slips through in certain sites where we run
into more problems than others. We did have
the DOE-wide classification officers meeting
just about a month ago. I went out to that;
Gina was there. We spoke to those groups and
met with many of them. Met with their
management and tried to emphasize the
importance of this program. We do believe
that some of the sites where we had problems.

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we have some connections now that I think will solve those problems. But again it is going to be difficult. I think communication is important, and one of the things that we have requested from SC&A and NIOSH is a list of the various -- I believe SC&A was going to put together some kind of tracking system of what's been submitted, to whom, and when received it. they've Because we necessarily at headquarters always find out about these issues until something has already been outstanding for guite some time. are kind of in catch-up mode. So that is one thing that may help, allowing us to know as things start to become an issue, not after they're an issue. But again we take it very seriously and do everything we can to work with our field sites to reduce the amount of time they spend reviewing these documents.

MEMBER CLAWSON: I appreciate that. I really would like to tell you how much we appreciate your people there at

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Germantown. They've been excellent in working with some of our documentation at Pantex.

CHAIRMAN MELIUS: Along those lines there is ___ Ι quess it's becoming infamous now -- the set of notes from Pantex that we have been waiting for for months; has that gotten released yet? We got it, okay. Good, excellent. That's good news. You should have a slide on that.

MR. LEWIS: No, and that is actually a good example, because when that was raised at the last Board meeting, my office, we weren't really aware that that was an issue. And I believe it was released a week or two after that Board meeting. So once we realized it was an issue and elevated its priority. But again we had no idea it was a problem.

CHAIRMAN MELIUS: And I also would like to thank you for your efforts on the Site Exposure Matrix, even though that is mainly a Part E issue. I'm glad that that

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1	information got out there.
2	MR. LEWIS: It's not all out
3	there yet. But we are working on it.
4	CHAIRMAN MELIUS: I know, but you
5	have done a good job in a short time period on
6	that.
7	Any other questions?
8	(No response.)
9	CHAIRMAN MELIUS: Okay. Thank
10	you.
11	We are running ahead of schedule,
12	but it won't help anybody. We will try to,
13	for the rest of the day we are on a fairly
14	tight schedule because of actually the next
15	two days with petitions. So we have got to
16	stay to the schedule, so we really can't start
17	until around 11:00 again, the time period
18	there. So we will try to catch up on a few
19	things, and then we'll take a break. But it
20	will be longer than the 15 minutes that's
21	listed.
22	MR. KATZ: So one of the

practices of the Board when a member is absent SEC vote is to collect that vote subsequent to the Board meeting, and then we register that at the next Board meeting, how at the teleconference that vote went. So March 31st the Board voted on Canoga, favor, unanimously. Fifteen members were present to add Canoga to the SEC. Dr. Poston was absent for that meeting but very shortly after, on April 2nd he contacted me with his vote in favor, so it was unanimous 16 votes in favor of that in Canoga, and I think Stu reported that Canoga has gone forward from HHS since.

CHAIRMAN MELIUS: I have two more things quickly. Just a reminder, there is a Work Group that sent out a solicitation. Got some responses, but just a reminder, if you can let me know there is a Work Group on, pending on Portsmouth, Paducah and the Oak Ridge, really a combined Work Group for Site Profile. I'm looking for volunteers for it.

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We have enough, but just in case somebody didn't see the information or whatever and would like to volunteer, just let me know.

Second thing on the case reviews, everyone should have received their assignment for the case reviews, and I believe SC&A has started to contact people, although I think they have only done -- or at least they only copied me on one group that they contacted, I think it was Josie, and so Josie and Henry, they started with A and B, the alphabetical approach. So which wasn't intended when we set up the group.

Okay, let's start with Ziemer next time. We'll work it out. And then Mark's group, I believe your dose reconstruction group has, in terms of the next round of cases

MEMBER GRIFFON: We did a preselection of cases, but the notion is that our usual process is to go through and pre-select cases, and then NIOSH provides additional

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details of those cases to allow for a refined selection of the final set to give to SC&A to work on. And they -- I mean we just had this phone call maybe a week ago, so they are not going to have that information ready for this meeting. What I proposed was to have it ready for the phone call Board meeting and then we could make the final selection at that point. That's simple.

CHAIRMAN MELIUS: We do have the letter to do?

MEMBER GRIFFON: Yes, I have a letter which I didn't circulate, I'm hoping to circulate tonight, I forgot to -- it's the summary report for the Dose Reconstruction Subcommittee. If you recall we were asked to follow up on the report that we submitted to the Secretary on the first 100 case review, and to follow up on sort of -- so what does this mean, and can we make any more specific recommendations to NIOSH in terms of the path forward and looking at their program.

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It does have some overlap with some of the items in the report that Lew Wade is actually assembling. So it might be useful from that standpoint as well. I have to email it tonight, so I'll get that around to you, but I forgot to send it over. It's only, I think, three pages or so long, and we can discuss it on Friday.

Right, CHAIRMAN MELIUS: and while we have a little time, because I will not be here on Friday; we had a meeting of the SEC evaluation Work Group in Cincinnati last week to address the 250 day issue and it was a very good meeting. made, We Ι believe, significant progress on that. We have some work assignments to do, but I expect that by Idaho Board meeting will our we have proposal for the Board to consider on that. That is at least our target, and then actually also specific recommendations on at least two of the sites that are sort of covered under I'm not sure on the third site whether that.

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1	we will be ready or not by that time. It's a
2	little bit more complicated, but we are moving
3	along. But the two sites are the Met Lab and
4	Ames Lab. The third one that we have been
5	looking at is Nevada Test Site, and that is a
6	lot more complicated. But we I do think
7	that we have got some agreement on our way to
8	move forward on that with NIOSH, and I think
9	we will be able to address that at our next
10	meeting.
11	Any other administrative matters?
12	Lew?
13	(No response.)
14	CHAIRMAN MELIUS: We will be doing
15	some tasking for SC&A while we're here, put
16	that in and so forth. Then some of that will
17	depend on sort of follow-up to a report,
18	update from our Procedures Subcommittee. So I
19	may be calling on you, Wanda, surprise you at
20	some point.
21	MEMBER MUNN: Don't.

CHAIRMAN MELIUS: That's why I'm

1	warning.
2	Why don't we take a break and
3	reconvene at 11:00.
4	(Whereupon, the above-entitled
5	matter went off the record at 10:20 a.m. and
6	resumed at 11:01 a.m.)
7	CHAIRMAN MELIUS: We will get
8	started. Our first presentation will be Brant
9	Ulsh will be presenting on the Mound SEC
10	petition. And Ted, do you want to check the
11	phones.
12	MR. KATZ: I already did check
13	the phones.
14	Dr. Richardson, are you with us
15	now?
16	(No response.)
17	CHAIRMAN MELIUS: Okay, they
18	should be able to get him. Go ahead, Brant.
19	MOUND SEC PETITION
20	DR. ULSH: All right, thank you.
21	As Dr. Melius indicated I am here to talk
22	about the Mound SEC petition. This has been a

long running process, and I'll go through some of the history of the SEC implications and the process that we have gone through over the last couple of years.

I'm going to focus pretty tightly on one topic, and that topic is radon. There are a number of other issues that are still before the Working Group under active consideration, and I won't touch too much if at all on those. I think the current plan is to bring those up for discussion between now and the August Board meeting in Idaho.

All right just to give you a very quick background on the Mound facility, the initial mission of the Mound facility was work with polonium-210 production, and that was for initiators in nuclear weapons. They also had a very active program in the early `50s, early and middle `50s, looking at alternatives to polonium-210, and that dealt with radium-226, and actinium-227.

As I mentioned those were for

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alternatives, and that program formed the basis of the SEC Class that we recommended a couple of years ago. And I'll talk a little more about that as we go on too.

Mound also had some small research programs with uranium and protactinium-231. They did some work with plutonium, and one of the bigger programs, in fact the main focus of Mound in the later years certainly was the space program. They produced radioisotope thermal electric generators -- now that is a mouthful, so you can just say RTGs - and that involved work, some work with polonium-210 but mainly with plutonium-238. Mound also did tritium research beginning in 1957. Now that was a very extensive program at Mound and long running. And that is one of the issues that is currently being deliberated upon by the Working Group.

A little more history of the Mound Site. It didn't spring into existence from nothing, it actually was the follow-on to the

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Monsanto Site, the Dayton Project as it is also known. That preceded the Mound Site, and then in about 1949 the work of the Monsanto Site was transferred over to the Mound laboratory, which is Miamisburg, Ohio, so right outside of Dayton, just south.

The next bullet here, I don't want to draw too bright a line here. Production occurred through 1994, however some D&D work occurred over the entire history of the Mound Site, and it's not like in 1994 everything stopped with production and D&D took over; it's not like that. It's just that in 1994 I think the official decision was made to shift the Mound mission from production to D&D. But there is some overlap on both ways on that with D&D and production.

D&D was the primary mission from about 1994 through 2006.

Now in terms of the SEC process that has gone on with the Mound Site, and I want to clarify here that I am talking about

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83.13 process, so this is a petitionergenerated process, we received two petitions in June of 2007, and they were qualified a couple of months later, and the petitions were subsequently merged.

And as we do with most 83.13s -in fact all of them -- we spend a few months evaluating the report --I'm sorry, then issued the NIOSH petitions, and we Evaluation Report in December of 2007, and presented that report the at Las meeting, I think that was January, 2008. right after we issued our report.

And the initial findings of that Evaluation Report we recommended a Class be added to the SEC based on the radium, actinium and thorium separations program that occurred in what was known as the Old Cave of the SW building. And that is an important facility, and I'm going to be talking about that at some length.

We concluded that we could not

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reconstruct internal doses associated with that program from 1949 through February 28th of 1959. Now there were a couple of efforts to decontaminate the facility that was used, the Old Cave, that was used for this program that occurred throughout the `50s, but the final -- I don't even want to say final, but the last of the major D&D effort was completed 1959. And basically that involved in concreting over the entire facility, that laboratory, pretty part, and abandoning that facility.

The problem is that some offices were built on top of that concreted in facility, and I'm going to be talking about that too.

Okay, now the radon petition: up to now I've been talking about an 83.13 process, so that was petitioner generated. Over the course of considering those 83.13 petitions, the Working Group and NIOSH and SC&A have discussed one of the topics that has

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come up is radon. And we have come to the conclusion that we need to recommend a Class based on radon. For administrative reasons we did that via an 83.14 petition, and that's what I'm here to present today.

So this is the standard language that you find for an 83.14. Basically we concluded that we could not adequately bound the radon dose for members of this Class. In April, so just last month, we embarked on the 83.14 process. And the petition was submitted earlier this month.

Okay, this is a very simplified schematic of what we are talking about at the Mound Site. I've drawn R and SW buildings. They are referred as RSW, they are referred to as two separate buildings but they are not really. These buildings were contiguous. They shared a wall. There were doorways between them. It's basically one structure, but two buildings were joined together.

And I've put in a red box down

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there, SW 19. That is a particular room in the SW building, and that's the one I told you where the office space was built right on top of the Old Cave. The Old Cave again is where they did the radium-actinium-thorium separations, formed the basis for the earlier Class, and during that process from 1949 to '59, contamination was pretty widespread through the R/SW building, and we determined that we couldn't really limit the Class from '49 to '59, so that covered everybody on site at that time.

I think it's worthwhile before I slide these leave this to talk about buildings. As I mentioned there are extensive tritium operations in these buildings, and work with plutonium in there was these buildings. So these aren't -- unless you have worked in a facility like that you may not really grasp the implications of that. But basically this building was operated negative pressure. Air was sucked up stacks.

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You certainly don't want to have work in one room in this building, have that air recirculated throughout the building and if something goes wrong contaminate the entire facility. So that's important to keep in mind here.

All right, now let's move forward past the earlier SEC Class from 1959 through 1980. And what you see on the first bullet of this slide is that I have a 20-year period from 1959 which was the end of the first class, up through 1979. And this is the real problem with the radon issue, and that is that we have a 20-year period where we don't really have radon monitoring. And that for me was kind of the straw that broke the camel's back on this Class.

Now I need to clarify here that when I say radon, I'm might be speaking in a - using the term in a different way than what you might be used to if you think about radon in your basement. I'm talking about three

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different isotopes of radon. Radon-222 is what you are most familiar with perhaps in your basement, but there is also radon-220, which is also known as thoron, and radon-219, which is actinon. And all three of those isotopes of radon are radioactive, and they all three generate a series of daughter products until they reach a stable species.

So what happens here. Well, 1979 a worker went in for a whole body count and it turned up high, so they were concerned he might have received а plutonium So they began to investigate where exposure. could exposed, because he have been he shouldn't have had any plutonium exposure. They checked out his office, which was in SW And what they discovered was that -- they 19. did some measurements near his desk, and they discovered high levels of radon. Now I want to clarify here that these early measurements -- and I'll get more in detail on the next slide I think -- these early measurements

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first of all used uncalibrated instruments. They had an estimated calibration factor. But basically it was sufficient to indicate that there might be an issue here. Then in July of 1979 they discovered a small hole in the floor of the office, and cracks along the baseboard, and they discovered really high concentrations of radon in the air that was streaming out of that hole in those cracks.

So here is a drawing of And you can see that there are three different see if office spaces here. Let me figure out how to work this. These squares right here represent desks. And I've got a number of radon measurements presented on this slide, now this is in June and July, so it's right after that worker showed up with this high body count. And I want to point out that all of the numbers on this particular slide are grab samples. So it's very short term; take a quick sample, and see what you've got. They are not time integrated samples on this

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here is that And what you see there high pretty numbers. are some Especially -- well, here is near the worker's desk, that had the high body count. And you can see 66 and 80 picocuries per liter. That is fairly high. Keep in mind that at the time the regulatory standard for a controlled area such as the R/SW building is 100 picocuries per liter. So it's still below the regulatory standard for that time. But you can see, I've got in red over here, this little dot, this represents the hole in the floor that they found. And they took some measurements right at the egress at that hole, and they measured over 700 picocuries per liter, and even over 800 picocuries per liter.

But what you can see here is that the concentrations first of all that they measured between where the radon was entering the room and over here in the breathing zone where the worker would have been exposed where

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he worked at his desk, there is about a factor of 10 roughly decrease, and that is an important thing to keep in mind. I'll talk more about that as we go.

After this time period, after June and July of 1979, so the latter part of 1979, they began to look for the source. Where was material coming this from? And thev discovered a tunnel, an inaccessible tunnel that ran underneath SW-19 pretty much down the hallway here, and this tunnel is about two feet and some odd inches, and they had to access it by drilling a hole through a manhole, sealed manhole cover. And they stuck a tube down in that tunnel and they measured what was in there, and let me tell you what they found.

For radon-222 they detected 88,000 picocuries per liter. For radon-220, that's thoron, they detected 28,000 picocuries per liter. And for radon-219, this is a whopper, 640,000 picocuries per liter. That's hot.

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That is a lot of radon.

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So one thing you need to, if you can do the math quickly in your head, I'll save you the trouble, 85 percent of activity, if you add up the three different radon isotopes, 85 percent of it is from actinon. That's the radon-219. Now interesting to note that actinon has about a four-second half-life. The daughter products are mostly short lived, if you look at the longest one, I think there is one in there that takes up to about 36 minutes. total half-life of actinon and the daughters is about on the order of 40 minutes.

So I think that at least is one reason that you see this factor of 10 decrease between the hole and the desk. One is just the dilution with room air, but the other thing is, we are talking about 85 percent of this activity is very very short half-life, so there is some decay before the air ever gets out of the tunnel and into the room, and

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further decay of those very short lived halflife species as it spreads out into the room.

let's move forward a Okay, now They discovered that the tunnel little bit. was the source of what they were measuring here. And I want to point out here that this was certainly a source of technically enhanced radon, but this is layered on top of naturally occurring radon that occurred throughout the Mound Site, not just in the R/SW building, but all over the Mound Site, from a couple of different processes and phenomena. There was a coal plant that was near the Mound Site, and as you might know the emissions from a coal plant are themselves radioactive, and can lead to detecting radon. And also -- in fact this was in an interview that SC&A conducted with the guy that took all these measurements -they had a problem with radon especially during the summer months when the soil was very dry, and you are operating a building here at negative pressure, so it's going to be

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sucking air in. And so during those summer months they could get some high radon levels. Now when I say high, I'm not talking about 640,000, but I'm talking high by the standards of what you might think of in your basement. And in fact since this building was monitored for plutonium, and you are measuring alpha activity in the air, sometimes those monitors would go off and subsequent investigations would reveal that radon is probably source.

So I'm not saying that the radon was zero at any point in time in this building. But what I am trying to point out here is that this is a technically enhanced source layered on top of natural sources of radon.

So in January of 1980 -- we don't have the exact date -- they decided to initiate some remediation of this tunnel, and tried to knock down some of these radon concentrations. So what they did is, they ran

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a stack from the tunnel and they installed a
turbine to suck the air from the tunnel, vent
it up a stack through a filter, and out the
stack. And they took some confirmatory
measurements afterwards, and what they found
was that they had significantly knocked down
the radon concentrations. And I have here the
measurements that they took in May of 1980.
Now the 8.2 picocuries per liter that were
measured near the worker's desk, the same
location as before, is a grab sample, but the
15.4 picocuries per liter is not a grab
sample. It's a time integrated sample using a
PERM, type of radon detector. And they
measured in the office next door with the same
instrument. These are again integrated
numbers, not grab samples. And they are on
the same order, eight to 15 picocuries per
liter. And again RCG for a controlled area is
100 picocuries per liter, was at that time.
So they were pretty successful in knocking
that down And they took further confirmatory

samples through the `80s and into the `90s and the concentrations were again confirmed to be low.

So just to sum up here is chronology of what happened with this issue: '49 to '59 you had the source-term. This is when they did the radium, actinium, thorium separation activities. That was the source for the technically enhanced radon, and we have an SEC Class that covers that time period.

From 1959 up through June 1979 you have essentially what is an undetected radon leak into SW-19, and then in January of 1979, then the worker turned up with the whole body did county. They the subsequent investigation. In January 1980 thev remediated the source of the radon. As I told you they stacked it. And in March 1980 they did some confirmatory sampling to confirm that their remediation was successful.

Now March 5th is what we propose as

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the end date of this 83.14 Class, because that is when they took the follow-on sampling. They did a grab sample that I showed you on the previous slide, and then they a confirmatory time integrated sampling. So that is the date that we are proposing that the Class for this 83.14 be ended.

They also recounted that worker that had the high whole body count, and they recounted him twice in May of 1980. The first count, since they measured such drastic reductions in the room air concentration of radon, they were a little bit surprised when they first count they did on this employee was still high, higher than they expected. is some documentation on this where they tried to consider why this might be. And eventually they did a second count five days later and it was much lower, near baseline.

So what they concluded just to quote from that, the report on that incident, the author said it is my opinion that someone

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breathing air containing a concentration of radon decay products at or near the non-occupation MPC of $1/30^{\rm th}$ of the working level would produce a lung count which is elevated and above normal or baseline.

He went on to conclude that the magnitude of this person's lung count is neither surprising nor alarming. And in fact they mentioned that especially during the summer months, in the month of August, workers would occasionally turn up with high body counts, and they sent home radon monitoring kits, and they usually came back positive, and that explained why they got the results that they did.

So we are proposing a Class, here is the two-pronged test for an SEC. You all on the Working Group have seen this before. The first test is, is it feasible to estimate the level of radiation doses for the individuals in this Class. I guess it occurs to me that I haven't really even told you what

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Class we are proposing. I'll get to that in the next slide. But the answer that we've it's probably come to there is, no feasible to reconstruct the dose for the people in this Class.

And the second test then is there a reasonable likelihood that such a radiation dose may have endangered their health? Well, when you are talking about hundreds of thousands of picocuries per liter I'd say the answer is yet to that.

Okay, sorry it took me so long to get to this, but here is the wording of the recommended Class. I won't read it verbatim to you, but this is what we have proposed. Now this might be a little confusing. Because we are basing it on tritium bioassays. Now let me walk you through the reasoning that we've got here. First of all let's start with that worker. Should that worker ever become a claimant, and come down with lung cancer, and not already be paid because he works in - for

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missed dose on plutonium, I can see that I
probably cannot put a reasonable bound on his
dose, nor can I put a reasonable bound or
anyone else that might have worked in that
office over the years. I don't know if there
was anybody, maybe. So that is the dose that
I can't reconstruct. Now we've been working
with the Department of Labor to come up with a
Class that they can administer. And I have to
tell you that we are still working with DOL to
come up with a Class Definition that they can
administer. What we proposed to DOL, since
they said that they can't really administer a
Class, that is limited to one room, just can't
do that. So we proposed, or it was decided
that we would make this all of R/SW building.
Now I don't think that everyone in the R/SW
building had an unboundable radon dose, but
it's an academic point at this stage of the
game anyway, because we are proposing all of
R/SW building. So then the question
becomes, how do you put people in R/SW

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building. Well, as I mentioned, they had an extensive program to work with tritium in this building. So people who worked in this building were on the tritium bioassay program. And that's why we are proposing that tritium bioassay be a marker for someone who had the potential to work in this building. Now that net is a little wider than just R/SW, because they did tritium work certainly in the building at the Mound facility, so we'll be capturing some people form there. And that's just fine, that's just the price you pay for making sure you get everybody. Another thing to consider is, and we heard this in a worker interview that was conducted with members of the Working Group, and SC&A and NIOSH and some former Mound workers. It was certainly the policy, and it was well known and well posted that if you went into the R/SW building, the policy was that you left a tritium bioassay sample, even if you only went in for only five minutes. That was the policy. Now the

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reality: the workers that we talked to told us
that well, yes, that is the policy and
everyone knew it, but it's not like there were
armed guards standing thee making you give
your sample. It's possible that someone could
have gone into R/SW building for a meeting,
maybe in an old lab, or just gone in for a
meeting, to deliver a letter, whatever, and
not left a sample; that's possible. So that's
fine, but I did I talked to them, followed
up with them on that, the workers, that we had
talked to, and it's just not plausible that
someone could have worked for 250 work days in
the R/SW building and not left a single
tritium bioassay sample. That is just not
plausible. In fact the worker I talked to
two of the workers that we interviewed. One
of them said that would just be highly
unusual. He said that I would go so far as to
say it couldn't have happened. The other
worker said, no, that's just couldn't have
happened. So we are pretty confident that if

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we use tritium bioassay data as a marker for potential there, work in that building, that we will capture the people we want to include in this Class.

to summarize here is So t.he feasibility and health endangerment findings. The already established Class from '49 to '59, that stays in place; there is no change to that. The change comes in from 1959 to 1980, and what we are saying is that we no longer have confidence that we can bound the dose to those three radon isotopes and their daughters from 1959 through 1980, and we are recommending the Class that you saw in the previous slide.

For the whole time period we are saying that we can reconstruct doses with sufficient accuracy for all the other radioactive isotopes that were present, and for external doses.

So I think that is my last slide, so if you had any questions I'd be happy to

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1	try to answer them.
2	CHAIRMAN MELIUS: Questions?
3	Bill.
4	MEMBER FIELD: I had a question
5	about the confirmatory tests that were
6	performed.
7	DR. ULSH: Yes.
8	MEMBER FIELD: You say they were
9	integrated?
10	DR. ULSH: Yes.
11	MEMBER FIELD: Can you give us any
12	more detail? Was it a one-hour integration?
13	One day? One week?
14	DR. ULSH: I can if you give me
15	five seconds or so. Okay, seven seconds. Ah,
16	here are the numbers I'm looking for. From
17	March 3 rd through March 11 hold on, let me
18	go back to the slide that showed that, from
19	March 3 rd to March 11 th they placed an RDT-310
20	PERM, that probably means more to you than it
21	does to me, beside this person's desk. So
22	that is a period of what, about a week. And

they got the result of 15.4. There it is
right there. And I told you the 8.2 is a grab
sample. From March 14 th , 1980, through March
$27^{\rm th}$, 1980, they measured this one, 7.7, from
March 27^{th} to April 17^{th} , 1980, 7.8, and from
April $18^{\rm th}$, 1980 , through May $8^{\rm th}$, 1980 , they
measured a 13.4. And then I'm reading from
the notes of the health physicist who did
this, and he said that another reading was
started on May 8 th , 1980. Beside this guy's
desk. So I don't have a result for that one.
So those are the confirmatory samples. There
were a couple of others up to 1980. I think
they did some more in 1982, and then they did
some more in 1990. Both of those were low.
Now I talked to a number of people, former
Mound workers, who are now on the ORAU Team,
simply because it is easy for me to do that; I
know how to reach out and touch them. They
were here around 1990, late `80s, and some
time between when this was discovered in 1980,
and 1990 when they were there, this office

space became unoccupied. In other words, they moved them. They didn't want people routinely working in there. I don't know exactly when in that 10-year time period that happened. I suspect right after they discovered this, but I can't say that for certain.

Now I don't mean that people never went in there; that's not the case. They did occasion of qo in to use some the on facilities that were still there, but it short basis. And it's on term my understanding that they did radon monitoring when they did. So that is some more detail on that.

MEMBER FIELD: I just have one follow-up. Do you know if there was any thoron monitoring done? After that point?

DR. ULSH: I know that they measured for actinon. They were looking for radon-219, because they specifically mentioned that. I would have to go through and look for the details to see if they were actually

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1	looking for radon-220, I'm not certain. But
2	it was done by the same guy who did these
3	measurements.
4	MEMBER FIELD: The reason why I
5	was asking, I don't think the E-PERMs would
6	measure those. So I just wanted to make sure
7	that we are not missing some exposures.
8	DR. ULSH: I would have to follow
9	up on that, I don't know off the top of my
10	head. But keep in mind that this was all
11	mixed in together in this tunnel, and so they
12	stacked it, and you could make the logical
13	assumption that if the radon and the actinon
14	is gone, the thoron probably went the same
15	way.
16	CHAIRMAN MELIUS: Bob.
17	MEMBER PRESLEY: I believe you
18	have a number of approximately how many people
19	that this would affect?
20	DR. ULSH: No, actually I wish I
21	did.
22	MEMBER PRESLEY: You don't? I

thought you did. I'm sorry.

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DR. ULSH: This is one of those things that in the flurry of the lead up to this meeting it occurred to me I should find out how many outstanding Mound cases there are, and then I forgot to do it. But what I can tell you is that I think there are on the order of 500 cases from Mound. The problem is I can't tell you how many we have completed, and how many are left outstanding. other thing to consider is that this is work in a plutonium building, so at least the lung cases, what we found at other sites is that three-quarters of them are paid on missed dose can't tell you about anyway. But Ι others.

CHAIRMAN MELIUS: Brad.

MEMBER CLAWSON: So how many total samples do we have? Because I was under the impression that there not that many samples. How many total samples do we have?

DR. ULSH: Do you mean after --

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1	MEMBER CLAWSON: After the initial
2	sample, after they found the crack and the
3	hole, how many times did they sample?
4	DR. ULSH: Well, okay for radon
5	it looks like one five times for radon
6	before remediation, and six working level
7	measurements before remediation. Here's after
8	remediation, we've got five samples there, and
9	you would have to add to that the samples that
10	were taken in 1982 and also again in 1990. I
11	don't know exactly how many samples that round
12	consistent of.
13	MEMBER CLAWSON: Was this done by
14	the same person that found that first crack?
15	DR. ULSH: Yes.
16	MEMBER CLAWSON: Because is there
17	some questions about his instrumentation and
18	how he did this, but that is kind of beside
19	the point here.
20	On your next slide where you kind
21	of got the Class Definition, "who were
22	monitored for tritium."

1	DR. ULSH: Yes.
2	MEMBER CLAWSON: Okay, is that
3	supposed to be should have? Were monitored or
4	should have been monitored.
5	DR. ULSH: No, it is not supposed
6	to be that. We purposely did not put that in
7	
8	MEMBER CLAWSON: That's what I
9	want you to explain to me.
10	DR. ULSH: I might need some help
11	from some other NIOSH folks, because I tend to
12	get focused on the SEC petitions that are
13	mine, and don't
14	MEMBER CLAWSON: Right.
15	DR. ULSH: It's my understanding
16	that that should have been monitored, has
17	caused no end of heartache for other
18	petitions. Am I right on that, Bomber? Yes.
19	So I think we have decided that
20	that is not something we should put in.
21	MR. RUTHERFORD: Also, and the
22	thought process is that we are identifying

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this Class from people who were monitored. So if we feel there are individuals that should have been monitored, then the Class Definition wouldn't work as defined.

CHAIRMAN MELIUS: Can I follow up on that?

may have misunderstood but thought at some point there were people that had indicated, the workers in their interviews, that they had been in building and weren't monitored for tritium. And how sure are we that I think your -- I think what Brant said was that well, we don't think they could have been in there for 250 days and not have done -- I think that's what might have been captured by the should have been monitored. And I guess my question is, how confident are we that the people wouldn't have been in the building for a significant period of time, and might not have been caught in the tritium monitoring program?

I think the original rationale for

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"should have" is we know that these monitoring programs were far from -- often far from complete in terms of capturing all the different workers that might be included in them.

DR. ULSH: Well, that's based on a couple of pieces of information. The first is the written policy, which is very clear in that anyone, visitor or worker, who went into this building for any operation whatsoever, was required to leave a tritium urine sample. Now the next logical question is, okay, now that is the policy but in terms of implementation of the policy how rigorously was that done. And what we heard from the workers that we talked to, who were there during the time period of this Class, was that it wasn't their take, and that's why I said earlier that a person could have gone in on an occasional basis. But to meet the requirements of this Class, 250 working days, possible; they said just that's not

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1 couldn't have happened. Now that is based on 2 what the workers who were there told us. 3 CHAIRMAN MELIUS: But the Class stretches -- I'll play devil's advocate here -4 - the Class schedule is for 21 years. 5 6 DR. ULSH: Yes. 7 CHAIRMAN MELIUS: That means 8 they'd go in there once a week, once every other week, might not -- that is not frequent, 9 10 it's -- if you work for that long a 11 period of time, you could accumulate 250 days, 12 and could you have escaped monitoring? 13 DR. ULSH: Without leaving single tritium urine sample. 14 15 CHAIRMAN MELIUS: Yes, that's what 16 I'm saying. Because again you keep using language that gives me a little heartburn 17 about, it's far from perfect, or it's not a 18 19 perfect -- it wasn't complete. And I'm trying to understand what the -- I have not read the 20 worker interviews. I know there were a number 21 I'm just trying to understand what 22 of them.

the workers actually said. And I don't know 1 2 if the Work Group has comments on that. 3 DR. ULSH: I don't have anything further to add. That's what I've heard. 4 5 I'll let the Working Group speak. 6 MEMBER CLAWSON: I still have a 7 question on this. Because we were involved in interview, and that person that 8 same pulled those samples for you, or started this, 9 10 and his name escaped me, we just referred to Radon, because that's what he 11 him as Mr. 12 continued to do, was not a part of the tritium 13 program. DR. ULSH: Right. 14 15 MEMBER CLAWSON: He was in and out 16 of that building, and his comment to us was, I didn't have to leave a tritium sample, because 17 18 I wasn't a part of the program. 19 DR. ULSH: Well, Ι read the 20 interview notes that are at least available in the SRDB, and I didn't see that in there. 21 22 Maybe he said and it wasn't captured in the

interview notes.

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This was when we MEMBER CLAWSON: went to Mound and interviewed. Because part of -- here is my issue with this, is, as we other sites have seen in numerous everything else like that, there is a program status, and this is how it is supposed to work; it didn't always work that way. with this statement that you got right there, it takes a lot of people out -- and I think That is for the you have got some issues. Work Group to discuss and go on from there. But it's out to us right now. This is part of the thing; how are you going to be able to do that?

DR. ULSH: Do what?

MEMBER CLAWSON: How are you going to be able to capture the people? You are saying that there is no way that anyone could have gone in there for 250 days over 21 year period and not left a tritium sample. And I beg to differ on that. I bet there are a lot

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of examples where it could have possibly happened.

DR. ULSH: Okay, well, I'm just telling you what the workers told us.

CHAIRMAN MELIUS: Go ahead, David.

MEMBER RICHARDSON: I've got two questions, and I'll start with the narrower I read the Class Definition as all one. employees, dot, dot, dot who were monitored for tritium exposure while working at Mound number of work days aggregating at least 250 work days. It doesn't say who ever least one tritium record and were had at employed at the site for 250 days. If I read it, or if you would read it again, when I read it it sounds to me like someone implementing this rule would say that they were monitored for tritium exposure for a number of work days aggregating to at least 250, implying that they would have to have 250 monitored days, not a single tritium record, and an employment

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history that spans 250 days.

DR. ULSH: No, we worked with DOL to come up with this. Now let me be careful when I say that, because DOL has not stated their official position on this Class. We are still waiting on that.

MEMBER RICHARDSON: But to me as someone who is reading this text, it doesn't say a single tritium monitoring record. And 250 days employment. It says, who were monitored for tritium exposure while working for a number of days aggregating to at least 250.

DR. ULSH: Right, certainly the intent, and we've received at least verbal confirmation from DOL that should this Class goes forward as defined, the way that they would implement it is, the person would have to have 250 days employment at the Mound Site, and they would also have to have at least one tritium uranalysis sample. Now that doesn't mean that they have to have 250 days of

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1 monitoring. Just a single tritium urine 2 sample is enough to put you in the Class if 3 you meet all the other conditions. So why did 4 MEMBER RICHARDSON: they want to have something written that is 5 6 different than how they are going to implement 7 it? I don't think they do. MEMBER 8 DR. ULSH: RICHARDSON: It sounds 9 10 like you are saying they are giving you their word that they are going to implement it in 11 12 this way, but it's not written explicitly in 13 that way. mean, I don't know, when I pick 14 Ι 15 this paragraph it's up, one very long 16 sentence, and my reading of it is that they are monitored for tritium exposure for -- you 17 have a bunch of parenthetical things set off 18 19 there -- but it's for a number of days 20 aggregating to at least 250. They aren't two separate criteria laid out here. 21 It would

have to say, who were monitored for tritium

while working at Mound and were employed for at least 250 days. It's sort of an "and" is missing. But that is a minor point.

The other question I had was, when I was listening to your presentation you seem to be laying out a case really for saying that you feel like there were high levels of radon but they were limited to measurements taken at a hole in an office in one area in one building, and that the levels measured within the office were you were characterizing as relatively moderate in that work area, they were certainly moderate. And that the radon mean here it like it is issue, Ι seems something that you could bound, and why is it not?

DR. ULSH: Well, if I heard your question correctly, at the very end there you said why do we feel that we cannot monitor it?

Or cannot bound it?

MEMBER RICHARDSON: Why can't you bound it?

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1	DR. ULSH: I know. I will tell
2	you that I struggled long and hard with this.
3	But you have got that 20 years, from 1959 to
4	1979 when there is simply no monitoring data
5	that we have discovered. Even in 1979 I've
6	got a little bit of reservation about the
7	measurements that were taken for a couple of
8	reasons. First of all they were using an
9	uncalibrated instrument. Now it's true that
10	they did have an estimate of the calibration
11	factor, so that is what it is. But the second
12	point to bring up is that when they first
13	started investigating this, the early samples
14	that they took, and for the people who were in
15	the room, these early samples so I'm
16	talking about the samples that were collected
17	in June or July, are by and large grab
18	samples; in fact completely were grab samples.
19	And also at the time they didn't recognize
20	that they had to consider these other isotopes
21	of radon. They were still approaching this as
22	a radon-222 problem. It was only as they got

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further into this investigation that the health physicist did something called the modified "Zivoglu," or something method. I'm looking right at Bill Fields, because he knows what that is, and I don't. But apparently he used that to determine if there was interference from other isotopes of radon. That was his kind of first clue that, hey, we might have a problem here with thoron and actinon. So Ι think that influenced sampling strategy in June and July. little uncomfortable saying to you that those measurements could be used to put a bound on dose when we are talking about doses this high for those reasons.

Now I am much more confident, once they figured out what they had here, that they had all three isotopes of radon and their daughters, and they used calibrated instruments, I've got a much higher confidence in those measurements.

So really, David, to answer your

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question, the thing that tipped me over was the 20-years where we just don't have anything.

CHAIRMAN MELIUS: Josie.

MEMBER BEACH: This is a comment just on the Class Definition. First of all is it possible to put language in that defines or captures special circumstances for those folks that didn't leave a tritium bioassay which we have reason to believe could have happened, or secondly, take the tritium out of the language and just go with anyone who worked in those facilities?

DR. ULSH: We have had several iterations of this Class Definition. And at least one of them had R/SW building in the definition. trying But I'm to remember exactly why that was taken out. I think it's because it was DOL's opinion that they just lacked the ability to place people in those buildings. So in other words, this definition is actually a bit broader, because you could

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1	have someone who worked in the T building and
2	left a tritium urinalysis sample, and they'll
3	be captured. Because we don't DOL feels
4	that they lack the ability to place people in
5	this building.
6	What was the first part to your
7	question? Were there two parts?
8	MEMBER BEACH: The first part was
9	capturing special circumstances. Can you put
10	language in there, and I know that's broad.
11	DR. ULSH: "Should have been
12	monitored."
13	MEMBER BEACH: There you go.
14	DR. ULSH: I would entertain any
15	suggestion you might want to make in terms of
16	a modification.
17	CHAIRMAN MELIUS: What about,
18	following up on Josie, what about a sort of
19	bifurcated definition that would be monitored
20	for tritium, the one you have now, or working
21	250 days in the building, and essentially the
22	burden would be on the records or through

affidavits for the claimant to show that they worked in the building, or missed by the other definition. Now, sort of the alternative to that, say someone can't show that they worked in the building, don't have the tritium, they come through NIOSH for dose reconstruction, might you identify them at that point in time. But if we added the definition up front maybe that makes it easier. I don't know how much certainty there is about, is the building -using building as a definition, or location as a definition, just for some employees, or is it for employee, there is just any documentation that they would have worked in the building in the personnel records that the Department of Labor was using?

DR. ULSH: No, I think that would be too strong a characterization. I mean we could certainly go through the worker files and in some cases identify that, hey, there is say for instance an incident report that occurred in a room in the R/SW building.

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Sure, we could do that. But I can't guarantee you that we could do that in every case.

CHAIRMAN MELIUS: Right, so we had a two-part definition, that one, that covered tritium which think the we encompasses everybody or should, maybe not everybody but most people, and then or the buildings, one or the other, you've got -- you only get So that would give compensated once. option а little more complicated administer, for Department of Labor, but it might -- it would cover the two groups. is a thought. Maybe you have already thought about it and said no.

DR. ULSH: Well, I understand what I understand the concerns you are saying. that are being expressed. All I can comment is the scientific aspects of on reconstructions. In terms of crafting a Class Definition there are other parties involved in this, and that is DOL and certainly So I think I can speak for those management.

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other parties and say that we would certainly entertain any suggestion that the Board would want to provide to us.

CHAIRMAN MELIUS: Henry.

MEMBER ANDERSON: Just quick question. Do you have any sense of what activities would have gone on where there would have just occasional visits? Were there conference rooms there? The issue over 20 years, you'd only have to be there 20 days, that's every other month. So to get your 250 davs if there was regular use of conference rooms, some people may have come over for meetings. Do we have any sense of kind of the occasional visits to the building that might have been regular? Because a whole group came and it would be aw gee, we aren't going to -- you won't hear for a couple of hours.

DR. ULSH: To answer your question directly, off the top of my head I don't know whether there were conference rooms in this

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building. I would assume that there were.
There were certainly laboratories in this
building, and there were certainly maintenance
people who came into this building. But again
they were covered, and they were certainly a
part of the policy. So any examples that I
could give you would be hypothetical. I'm not
aware of any functions that would take you
regularly into this building. I'm sure that
there are. But again the opinion that was
expressed to us by the workers that we talked
to were, people who fit that definition, who
even if it was occasionally but regularly, so
let's say once a month over the years, they
just didn't think it was plausible that
someone could have done that and not left a
single tritium urinalysis sample. One time,
two times, yes maybe; but 250 times.

MEMBER ANDERSON: Are you talking two workers that you talked to? Or how many?

DR. ULSH: Well, I talked to the two workers that were interviewed by NIOSH,

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SC&A and the Working Group. Then I talked to eight other workers who were there. Now granted, toward the end of the SEC period. So the two that I talked to were the ones that were there for the entire period, the entire Class period.

MEMBER ANDERSON: Ι was just trying to kind of back in to expanding this because there reasonably could be some people that were missed and there may be -- you could think in terms of management folks that would for regular meetings come over but didn't really work there. Sometimes they slip through the cracks.

DR. ULSH: This is a little dangerous, because I'm only speaking for NIOSH now and I'm explicitly not speaking for the Working Group and SC&A, because I know there is some disagreement on this issue. From NIOSH's perspective you have this technically enhanced radon concentration that occurred in SW-19. It did not occur -- we don't see any

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evidence that it occurred throughout the R/SW building. The other areas. That is strictly from our perspective an administrative construct to administer the Class.

like Ι said, the Now, other parties are involved here. The Working Group and SC&A have expressed some disagreement with just going into the that viewpoint. So building from our perspective again doesn't expose you to radon. If you go into SW-19, okay, I'll grant you that, there is some exposure there. Certainly if you went up and stuck your nose next to the crack in the floor you are going to get some radon exposure. just walking into the building I don't think That's just an administrative construct so. from our perspective.

CHAIRMAN MELIUS: But I also think it is -- I mean, you're right, it is administrative, but it's also, certainly if you were working in the building your probability of having been in a room with

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higher exposure is a higher probability. Now you are right, you may have never gone near the room, but the chances are, at least based on the information we have. And I think how we are struggling with is how do we make that definition.

Phil you had a comment, and then Josie.

MEMBER SCHOFIELD: Do you know if there is a time clock key station or any seals that the guard force or security would have had to go in that building and punch or check on a regular basis? Because I doubt they would left a sample when they went in there to just punch a clock or check a seal?

DR. ULSH: No, the answer to your question is I don't know. If you carried a time card with you and every time you went in and punched it or something like that.

MEMBER SCHOFIELD: It looks like a round clock. You go in there and you put in a key and what it does it makes a mark on a

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piece of paper, tells which key station you are at and what time. And that normally is -it's a process takes about 30 seconds. Also anything, maybe if you had a vault, maybe you had exterior doors that had to be sealed from the inside, they would go in and they would put a seal on this. And I cannot imagine those people every time they go in to check that seal or punch that clock would have left a sample.

DR. ULSH: Okay, let me tell you what I do know and what I don't know to answer your question. There may very well have been exactly what you are talking about. I don't know. I just don't. I can tell you that the written policy -- I mean I have already said this a couple of times was, even if you went in for any reason, you were supposed to leave a sample. Now there was no physical barrier there or personnel guarding the entrance to make sure that you left your sample. And hence there is this caveat that people could

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have visited occasionally and in contravention 1 2 of the policy just didn't want to be bothered 3 with it, I didn't go a hot area, whatever the reason might have been, and didn't leave a 4 I mean we are acknowledging that. 5 sample. 6 Might there have been a cohort, a 7 smaller cohort of workers, perhaps, I don't know, maintenance workers that might have been 8 subjected to the kind of thing that you are 9 10 talking about? Could have been, I don't know. I just don't know. 11 CHAIRMAN MELIUS: Josie. 12 MEMBER BEACH: Thank you. 13 CHAIRMAN MELIUS: Brant, we'll let 14 15 you sit down. 16 MEMBER BEACH: First of all, let Group 17 me say that the Work conducted discussion on 18 vigorous this issue. 19 presence of radon sources under SW-19 along 20 with tunnel underlying the foundation of parts buildings permitted radon to seep of R/SW 21

into work areas via cracks in the floor over

time. Something that was accelerated by the
use of negative pressure fume hoods for
operations. The Evaluation Report originally
indicated that there was sufficient radon data
for dose reconstruction, but it turned out to
be based on more current records from the
1990s and did not reflect on the earlier
years. The few measurements that were taken
in 1979 and 1980 and the SW-19 provided us
with an opportunity now to reach an agreement
to recommend this SEC Class for March $1^{\rm st}$,
1959, to March 5 th , 1980. But it does not
necessarily resolve the question of
reconstructability for the later years after
1980 the Work Group needs to reassure itself
that those few measurements in that one room
are bounding after 1980. For radon doses
apparently being experience elsewhere in the
R/SW complex and are in fact bounding even in
SW-19. Given some later indications that
elevated radon may have continued being
experienced there. We will be asking NIOSH

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and SC&A to continue investigating these questions over the coming weeks. I will ask both NIOSH and SC&A to come to the next Work Group meeting for Mound prepared to address the post-1980 period for radon. I will also work with Ted to schedule the next Work Group meeting, and I'm hoping that for the end of June time frame or July, and of course that depends on the action items that are on the list.

I will also be sending out the action items list to the Work Group with this additional action for NIOSH and SC&A added to it. The other reservation as we discussed was the Class Definition, and I understand we have had that discussion.

CHAIRMAN MELIUS: Thank you,
Josie, thank you for all the work that your
Work Group has done on this. My sense is that
I think from the tenor of the questions and so
forth I think we are satisfied that this
should be added to the Class. I think we are

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all still struggling, as is I think NIOSH and
DOL and the Work Group with what should be the
Class Definition for that, and maybe we can
sort of put this off, any final consideration
on this, until some time later in this meeting
to give time for NIOSH and DOL to at least
have some initial discussions on some of the
alternative approaches to dealing with this
Class that we've suggested. But I'm not sure
we are ready go vote on a conclusion, and I
would hate to have us come to a conclusion and
make a recommendation if we haven't we
really need to have some consultation on what
should be in the Class Definition. So maybe
they can work on that, and if it's okay with
the Board Members to postpone our final
consideration on that at this meeting; we'll
get back to it at this meeting to try to reach
some conclusion and decision on that; is that
fair?

MEMBER BEACH: I guess my only question about that is, we had a Work Group

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meeting last week, and NIOSH was working on that Class Definition, and came back with the same definition that was talked about during our Work Group meeting. So I wonder if NIOSH would be ready during this Board meeting to answer those questions.

CHAIRMAN MELIUS: Well, let's press them to do that, because I think we'd like to reach a conclusion, and talk, but I'm understanding that even with the Class Definition that is in the 83.14 report, DOL has not fully signed off on that yet, and I think we talked about some alternatives, and I guess I'd like to get some feedback from them. But I think to get some feedback they need some time, and let's see what we can get to at this meeting and decide. Is that fair to everybody?

MEMBER ROESSLER: It seems there are two things under consideration here, and I just want to have it clear in my mind. First of all is the definition on how do you define

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what workers would be included, and that's what you are talking about. But what Josie just brought up makes me think that there is even a question about the date, that March 15th. Am I right, Josie?

MEMBER BEACH: You are correct.

We didn't want to hold up this earlier time period by looking at the later date book.

MEMBER ROESSLER: So if you resolve, or if it's resolved, the question about what workers are included, then does this definition go through? And then your question is about the later measurements, and can they be bounded or not. That's another petition. I would think. I just wanted to be clear on that.

CHAIRMAN MELIUS: It would be most likely another petition, and I think it would be further consideration. The alternative is if we can't reach consensus on agreement at this meeting is to postpone any consideration of this until another meeting, and maybe we'd

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1 have at a last time period when we can address 2 that more fully. But let's see what we can do 3 earlier time period while we're on the 4 gathered here today. 5 MR. KATZ: Just to clarify 6 something, I don't believe it would require 7 another petition, because this is an 83.14, and they have a clause in their Evaluation 8 Report which allows for an expanded Class to 9 10 be considered following the resolution of that initial 83.14. So I don't think it would 11 require a new petition. 12 13 CHAIRMAN MELIUS: Okay, thank you for that. 14 15 Let's move on. Rochester I think 16 we can do relatively quickly, and I think there might be a petitioner before we do that. 17 MR. KATZ: Yes, I thought we'd get 18 19 to a vote, so I didn't address conflicts on 20 the front end of this. But there were three Board Members who have conflicts, for Mound: 21 Mr. Gibson, Mr. Griffon, and Dr. Lockey, and

1	they had all recused themselves from the
2	discussion which I would have noted if we got
3	into a vote. Thanks.
4	CHAIRMAN MELIUS: Okay, Rochester.
5	Laura is not here?
6	UNIVERSITY OF ROCHESTER ATOMIC ENERGY PROJECT
7	SEC PETITION
8	MR. HINNEFELD: Hello again.
9	This is Stu Hinnefeld.
10	Dr. Hughes couldn't make it this
11	trip. She is restricted from traveling until
12	she has her baby later on this summer. So I'm
13	here to present briefly an update and we
14	simply provided to the Advisory Board a week
15	or so ago an addendum to our Evaluation Report
16	for the University of Rochester that was
17	originally presented in October.
18	Just a summary of the history. In
19	October we presented the 83.13 SEC evaluation
20	and proposed this Class: All employees of the
21	Department of Energy, its predecessor agencies
22	and contractors and subcontractors who worked

at the University of Rochester Atomic Energy Project in Rochester, New York for these dates, September of '43 through October of '71 for a number of days aggregating to 250. So that was the Class we presented in October.

And this is the feasibility table we presented. We had found that we could not have sufficient information to reconstruct the internal doses for any of the period, and then some of the external doses for other periods.

After we presented to the Advisory Board, the Advisory Board recommended that we try to look additional places for data capture to see if we couldn't some internal monitoring data that was relevant to the claim. And so specifically what was mentioned was the State of New York, and maybe if you would contact the laboratory director, the laboratory director oftentimes kept these records and would know where things went.

So following on from that, we started down the trails starting with those

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two situations, and followed the trail a number of places, ended up at six different locations looking for University of Rochester.

And what's particularly relevant is the appendix to the addendum to the SEC Evaluation Report. That appendix starts on page 11 of that document, and it lists the documents that were found during these various data captures.

Going down the locations one by one we started with New York State agencies, and we were not able to find any additional information from the New York State agencies. We had previously contacted a number of New York State agencies, in our original investigation back in 2007 during our original research on this, and we had found no more since October of 2009.

We were taken to Hanford by well Dr. Lockey found information, the collection of Dr. Newell Stannard who had run the laboratory at University of Rochester for awhile, was transferred from Rochester to

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Hanford. So we contacted Hanford for search, and found quite a lot of hits from their finding database related to Rochester, key words, we asked to see some Hanford is having trouble finding all 72 boxes that had some of those hits in them, but we have found some. Part of the collection apparently was loaned to Washington State University, so that then becomes our third location that we looked for, Hanford did finally located boxes pertaining University of Rochester. Dr. Stannard had records about a lot of places, just because Dr. Stannard's name was on a record doesn't mean it pertained to the University of Rochester

So I did find some boxes at the Seattle federal records center, and we went there and captured some documents from those.

Like I said Hanford indicated some of those documents were loaned to Washington State University, so we went there. And specifically to the transuranium and uranium

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registries. And we found indication there
that some of the documents associated with Dr.
Stannard had been sent to the University of
Tennessee in Knoxville. So we did find
finally at the Washington State Library six
documents relevant to the University of
Rochester, none of which helped us out on
internal monitoring for people working at the
University of Rochester. We went to the
University of Tennessee, found 26 boxes there,
data captured this year. We found again 51
documents relating to Rochester, but not
internal dosimetry that would help us out in
providing reconstruction of this Class. Since
we were at University of Tennessee, I think
there was some sharing of information between
University of Tennessee and the Oak Ridge
operations office vault; that is the DOE
office down there. They have a vault where
they store classified information so we
searched that and found a number of boxes,
found some film badge services that the

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University of Rochester had provided to other sites, because recall back in the early days a lot of these AECs, they would issue film badges to their workers; those film badges were processed by the University of Rochester. We found a limited amount of film badge data for the University of Rochester, but in our feasibility issue with the was internal So we couldn't solve our monitoring data. issues there.

We also discovered a finding aid for NARA, that's National Archive and Records Administration, facility in College Park, Maryland. We were actually looking there in October when we presented to the Board and at that time that we were searching there. received some documents in November, and we made additional data capture this year. After finding a new finding aid and we captured a number of documents, but again, nothing that would allow do internal us to dose reconstructions for the University

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So having concluded all that, think the actual documents, a description of the documents found are listed like I said in the appendix of the evaluation, the addendum to the Evaluation Report. You can see it's quite a long list of publications, and a great deal of animal studies having to do with the radiobiology of radioactive materials when administered to animals, so the University of Rochester was heavily involved in that early determining the fate of radioactive materials that are administered to animals. Very much of what we found fit into that. did not find any information though that would make it feasible for us to reconstruct internal doses of the people working at the University of Rochester.

So we -- our addendum to the Evaluation Report did not change our proposed Class Definition; did not change our conclusion about feasibility, and so naturally

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1	if we find the dose is infeasible, we always
2	determine that if we can't bound the dose that
3	we have to conclude there is a potential for
4	harm, and so we present that to the Board as
5	essentially our recommendation again that this
6	Class be added for those purposes. We were
7	not able to find additional information,
8	despite the search over the last six months
9	roughly to allow us to complete those dose
10	reconstructions.
11	CHAIRMAN MELIUS: Okay, thank
12	you, Stu. Do Board Members have questions?
13	We have talked about it a couple of times
14	before. If there are no questions, could we
15	actually before I do that, Paul Ziemer or
16	David Richardson, do you have questions?
17	MEMBER ZIEMER: What was the
18	question again?
19	CHAIRMAN MELIUS: Do you have
20	questions.
21	MEMBER ZIEMER: No, I'm clear.
22	CHAIRMAN MELTIIS: Dr Lockey

MEMBER LOCKEY: In your definition, I forgot, but what did you say about graduate students, et cetera.

MR. HINNEFELD: Well, our definition doesn't say anything about graduate students. It says people who worked for the Department of Energy, its predecessor agencies, and contractors. So if you are employment question, asking an verified а employment question of would а graduate student be included. And I quess I don't really know; that would be the Department of Labor. We would make no distinction. The of Labor identifies someone Department as having verified employment at the University of Rochester Atomic Energy Project then it would make no distinction for us, and in fact we would not see these claimants. wouldn't even come to us.

MEMBER LOCKEY: So anybody who received funding through the Department of Energy that funneled through the University of

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Rochester would fit into this Class?

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MR. HINNEFELD: Well, the Class, the work facility is defined as the University of Rochester Atomic Energy Project, which was think that specific I was specific building or set of buildings at the University of Rochester. So whether or not people who are funded through the University of Rochester and did the work, certainly somewhere other than University of Rochester, I don't see any possible way they would be included. who worked at the University of Rochester and have a valid claim with the Department of Labor, by our definition would be included. It all comes down to whether their employment was verified by they Department of Labor as being appropriate coverage.

MEMBER LOCKEY: Is there a building definition in this Class?

MR. HINNEFELD: Well, the name of the facility, which is the name of the covered facility on the DOE database, it's the

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University of Rochester Atomic Energy Project.

That is in our Class Definition. That is the name of the covered facility certainly on the database and probably on the Federal Register that listed the covered facilities.

MEMBER LOCKEY: I guess what I'm trying to get a handle on is we have a covered facility and a covered project. And there might have other people been who employed through the Department of Energy on the University of Rochester campus. thev walked this building, into and accumulate 250 days in this building, are they included in the Class Definition?

MR. HINNEFELD: Well, as always, if they are included in the Class Definition is a determination of the Department of Labor. I hate to be evasive about your question, but I don't know that I can answer, don't know all Ι that Ι know the employment relationships or arrangements at the University of Rochester. I don't know

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that a graduate student, if a graduate student is considered an employee or not. So I suppose Jeff is prepared to opine on those questions.

CHAIRMAN MELIUS: Do you have anything to add, Jeff?

Jeff Kotsch, Labor. MR. KOTSCH: I guess the only thing I would say is unless you would -- you might lose grad students if they were members of the university who were not employed by the Department of Energy or its predecessor agencies or contractors subcontractors because they worked project. don't know if that Ι is possibility or The not. way the Class Definition is written, because there is the employment portion as well as the facility portion.

MEMBER LOCKEY: Working at a university, the reason I raise the issue is, graduate students do a lot of the work, if not most of the work in the lab. And the PI sits

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in his office and supervises. So it really is an important issue, which I think somehow has to be addressed.

MR. KOTSCH: I have to admit, I don't know like if you were on -- I know when I was in grad school, if you were on a grant or something that came through that, how that would show up as far as I was always just an the university employee of grad as а assistant. Again, I don't know -- again it comes down to the situation for those types of people having to be treated on a case by case basis, looking at the available information, trying to determine if they fit into the Class From the information that was Definition. presented in the Class Definition.

Maybe we can ask CHAIRMAN MELIUS: for an update on that in a more general sense so we understand it better. Because it comes of up, it comes up in some the university-affiliated facilities also, think understand it better. we need to

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Recognizing that there are limitations on what you can do given the law and the various contractual relationships that are out there. Thank you.

Josie.

MEMBER BEACH: And I just have an addition. I remember asking that question specifically the last time this came up at a Board meeting. There were 799 grad students, and I believe we were told that they weren't covered. That is just memory from this last discussion we had. And I am also concerned about that.

CHAIRMAN MELIUS: I don't remember 799, but Wanda.

MEMBER MUNN: My memory of the language of the law itself is specific about employees of the Department of Energy. That being the case there is nothing we can argue here if there is an issue, then the folks who were instrumental in writing the original law need to address the Congress of the United

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States on this issue. I can't see how it can be resolved if my memory is correct. My memory may not be correct; I haven't read the law in several months.

Other ways of designating facilities if they meet the definition, so that can be changed. So it's not just necessarily Congress. But you are right, there are limitations based on employment, and what they recognize in the contracts and those are usually not straightforward, and they change over time. Henry.

MEMBER ANDERSON: I would have thought that this project was contracted to the University of Rochester so that grad students were employees and could show that they had worked on a project for the 250 days, you would think they would be covered because they are subcontractors.

CHAIRMAN MELIUS: Yes. I would think so. Other questions or comments.

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MEMBER ZIEMER: This is Ziemer. I
do have a question, and this may have come up
before, but has it been established that the
quote project, the Atomic Energy Project, was
in fact building specific? And the reason I
ask that is those who work on campuses
recognize that grants that are made to either
a project or a center or whatever the name is
are frequently scattered throughout multiple
buildings. The project is a paper entity, and
may not be building specific. It may include
labs in a number of different places, and then
that opens the door to accessibility by
others, and I suspect on this particular one
it could have certainly been building specific
or location specific. But has that ever been
confirmed.

CHAIRMAN MELIUS: Go ahead, LaVon.

MR. RUTHERFORD: Dr. Ziemer, yes, we actually did correspond with the Department of Labor, and it is building specific.

MEMBER ZIEMER: I thought that had

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1	been asked before, but I couldn't recall if it
2	had been resolved. Thank you.
3	CHAIRMAN MELIUS: We also may have
4	the petitioner on the line, if she is
5	available.
6	MS. KESTON: Yes.
7	CHAIRMAN MELIUS: Do you have any
8	comments to make?
9	MS. KESTON: That it was a
10	specific building when I worked there from
11	September of '43 to June of '45. It was only
12	in a specific building.
13	CHAIRMAN MELIUS: Okay, thank you.
14	If there are no further comments, do I hear a
15	motion?
16	MEMBER PRESLEY: So moved.
17	CHAIRMAN MELIUS: So moved what?
18	MEMBER PRESLEY: That we accept
19	it.
20	CHAIRMAN MELIUS: Thank you, Bob. MEMBER
21	ANDERSON: Second.
22	CHAIRMAN MELIUS: Second from

Henry. And Bob if you will accept a friendly amendment. Let me read the statement.

Maybe I should have said, what I hope to be a friendly amendment.

This is our letter and forbear a little bit.

The Advisory Board on Radiation Worker Health, the Board has evaluated SEC Petition 00140 concerning workers University of Rochester Atomic Energy Project on the statutory requirements established by EEOICPA incorporated 42 CFR Section 83.13. The Board respectfully recommends a Special Exposure Cohort be accorded to all employees of the Department of Energy, its predecessor agencies, and their contractors or subcontractors who worked at the University of Rochester Atomic Energy Project in Rochester, New York, from September 1st, 1943 through October 30th, 1971, for a number of work days aggregating at least 250 work days occurring either solely under this employment or

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1 combination with workdays within the 2 parameters established for one or more other 3 Classes of employees in the SEC. recommendation is based 4 This following factors: the University 5 the 6 Rochester Atomic Energy Project conducted research in technical projects related to the 7 development and production of nuclear weapons. 8 Number two, NIOSH found that there 9 10 was insufficient monitoring data 11 information on radiological operations at this 12 facility in order to be able to complete accurate individual dose reconstructions. 13 Four, the University of Rochester 14 15 Atomic Energy Project employees during the 16 time period in question. Board with this 17 The concurs conclusion. 18 19 Three the Board has reviewed 20 information which confirms radiation the University of Rochester 21 exposures at the Atomic Energy Project during the time period 22

in question could have been dangerous to the health of members of this Class.

The Board also concurs with this conclusion. Based on these considerations and the discussions held at our May 19-21, 2010 Advisory Board Meeting in Niagara Falls, New York, and our two previous Board meetings, the Board recommends that this Special Exposure petition be granted; enclosed Cohort documentation from the Board meetings where the Special Exposure Cohort Class was discussed. The documentation includes transmits of deliberations, copies of the NIOSH review thereof, petition, and related materials. If any of these materials are unavailable at this time they will follow shortly.

Do you accept that?

MEMBER PRESLEY: I accept it.

CHAIRMAN MELIUS: Okay. And I will note for the record, I don't think counsel has had a chance to review this yet.

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1	So everyone will get another look at it at
2	some point, but I wanted to at least get it
3	into the record now.
4	So any further comments or
5	questions? If not I think I'll ask Ted to
6	call the roll.
7	MR. KATZ: Okay, we will just do
8	this alphabetically this time. Dr. Anderson.
9	MEMBER ANDERSON: Yes.
10	MR. KATZ: Ms. Beach.
11	MEMBER BEACH: Yes.
12	MR. KATZ: Mr. Clawson.
13	MEMBER CLAWSON: Yes.
14	MR. KATZ: Dr. Field.
15	MEMBER FIELD: Yes.
16	MR. KATZ: Mr. Gibson.
17	MEMBER GIBSON: Yes.
18	MR. KATZ: Mr. Griffon.
19	MEMBER GRIFFON: Yes.
20	MR. KATZ: Dr. Lemen.
21	MUMDED I EMENA MAR
j	MEMBER LEMEN: Yes.

1	MEMBER LOCKEY: Yes.
2	MR. KATZ: Dr. Melius.
3	CHAIRMAN MELIUS: Yes.
4	MR. KATZ: Ms. Munn.
5	MEMBER MUNN: Yes.
6	MR. KATZ: Dr. Poston.
7	MEMBER POSTON: Yes.
8	MR. KATZ: Mr. Presley.
9	MEMBER PRESLEY: Yes.
10	MR. KATZ: Dr. Richardson.
11	MEMBER RICHARDSON: Yes.
12	MR. KATZ: Dr. Roessler.
13	MEMBER ROESSLER: Abstain.
14	MR. KATZ: Mr. Schofield.
15	MEMBER SCHOFIELD: Yes.
16	MR. KATZ: Dr. Ziemer.
17	MEMBER ZIEMER: Yes.
18	MR. KATZ: So we have 15 in favor
19	and one abstention, and the motion passes.
20	CHAIRMAN MELIUS: Okay. Lunch
21	time. Take a break. We are a little bit
22	late. We do have I believe a petitioner that

1	will be here, so we'll try to start maybe
2	1:35. That gives an hour and 15 minutes.
3	Hopefully everyone can make it back by then.
4	Thank you.
5	(Whereupon, the above-entitled
6	matter went off the record at 12:24 p.m. and
7	resumed at 1:46 p.m.)
8	CHAIRMAN MELIUS: We can
9	reconvene. Ted, you want to check the phones?
10	MR. KATZ: Yes, let me check the
11	lines and see that we have Dr. Ziemer and Dr.
12	Richardson?
13	MEMBER ZIEMER: Ziemer is here.
14	MEMBER RICHARDSON: Hi, David
15	Richardson.
16	MR. KATZ: Great. And for
17	everybody else on the line, maybe some new
18	people will have joined us post lunch, we are
19	starting a little bit late. We had a late
20	ending of the morning session, and we are
21	about to get going. So let me note for any
22	people who may have joined us just freshly now

1	this afternoon who are on the call, please
2	mute your phone. Use the *6 button if you
3	don't have a mute button, and that will mute
4	your phone. Use *6 again if you want to speak
5	to the group. And please don't hang up I
6	mean please don't put your call on hold. Hang
7	up and call back in if you need to leave at
8	some point.
9	CHAIRMAN MELIUS: And we will
10	start this afternoon, first order of business
11	is BWX Technologies. And LaVon, you're up.
12	
13	BWX TECHNOLOGIES (LYNCHBURG, VA)
14	SEC PETITION (83.14)
15	MR. RUTHERFORD: Thank you, Dr. Melius.
16	I'm going to talk about BWXT, and our Special
17	Cohort petition for that facility.
18	We in April 6, 2010, after many
19	data capture efforts and work we decided that
20	we could not reconstruct dose for a period of
21	time at BWXT so we at that time sent a letter

to a claimant indicating to that claimant that

1	dose reconstruction was not feasible, and we
2	provided that claimant a form A petition to
3	submit an SEC petition for that site.
4	On April 14 yes.
5	MEMBER PRESLEY: Can we do one
6	thing, can we designate that as BWXT Virginia?
7	Because there are BWXT sites now all over the
8	United States in the weapons complex.
9	MR. RUTHERFORD: I actually had
10	that on my next slide.
11	CHAIRMAN MELIUS: Okay. We will
12	make sure.
13	MR. RUTHERFORD: Okay. All right,
14	so on April $14^{ m th}$, 2010, we received an 83.14
15	SEC petition. That petition qualified on that
16	day. And we after a month or so we issued our
17	Evaluation Report on May 4 th , 2010.
18	A little background: as Mr.
19	Presley indicated, BWXT is located in
20	Lynchburg, Virginia. It was an Atomic Weapons
21	Employer for three separate time periods,
22	which is kind of a little unique: January 1,

1959 through December 31st, 1959; January 1, 1968 through December 31st, 1972. And January 1, 1985 through December 31, 2001.

So during those three separate time periods there were different AEC activities. There were commercial activities that went on all the way from 1956 all the way to present.

From our interview indication we have that the workforce ranged from roughly 1,000 employees up to 3,000 employee at that facility.

The facility is actually BWXT is designed as a single site. However there are two licensed locations: Naval Nuclear Fuels Divisions; and the Lynchburg Technology Center. The Naval Nuclear Fuels Division, the involved primary mission is in fabrication using enriched uranium. The LTC which is -- I'm going to use NNFD and LTC as acronyms instead of repeating their names --LTC's work primarily involved reactor

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research, fuel testing and hot cell work.

Our sources of information that we looked to retrieve data, we looked at our existing Site Profiles. We have no Site Profile for BWXT. Technical Information Bulletins, NIOSH Site Research Database, data captures, and worker interviews. We did 36 worker interviews. Those were interviews, and I'll talk a little bit about them shortly.

We did data capture efforts with BWXT, DOE Legacy Management, DOE Germantown, and NRC, NNSA, the Virginia Department of Health, Westinghouse Site, the Hematite actually had some documentation that pulled; R.S. Landauer. Landauer did the film badge for BWXT. And U.S. Transuranium and Uranium Registries, as well as Washington State University's DOE OpenNet, Internet searches, CEDR database, Hanford's DDRS, and National Academies Press.

Existing claims, this is

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information as of the $4^{\circ\circ}$ of the May. We had
78 claims submitted to NIOSH. Of those 78, 62
meet the Class Definition that we are
recommending for the SEC. We have completed
two dose reconstructions within that Class,
and I broke down the claims, internal
dosimetry and external dosimetry, based on the
internal and external monitoring data for
those periods. So we had, you can look at it,
you can see that we had three claims within
internal dosimetry in 1959. We had 39 and 43
for the other two periods. And then the
external dosimetry. I could have actually, it
would have probably been a little more helpful
if I had put in the actual total numbers as
well for that period, but it does give you
some indication. And I'll talk more about the
internal dosimetry shortly and external.

Site operations: LTC was in mу mind very much almost like a national lab that dealt with. did They research we and radioactive development with materials and

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reactors from 1956 to 1984. Unencapsulated fuel work with uranium and thorium from 1957 to 1984. Reactor operations, there were a number of reactors, there is a table in the actual report that lists some of the reactors that operated and their time periods. Reactor operations from 1957 to 1983. They did thorium U-233 fuel 1964. research in Plutonium fuel research in 1966 through '71. And then laboratory analysis work for all the facilities, both the operations going on at LTC and NNFD occurred at the Lynchburg And they also had cask Technology Center. handling, liquid waste disposal, hot cell storage of highly activated work, and contaminated materials, and fuel cell inspections. And that all occurred from 1960 to the present.

NNFD, which was -- a lot of people considered the main plant, did uranium fuel fabrication from 1956 to present; thorium fuel fabrication from 1956 to 1963; and

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downblending of highly enriched uranium to fuel grade enrichments in the later years of 1995 to 2000.

Potential radiation source of exposure, primary radionuclides that were the source of external and internal exposure was uranium, typically enriched from 4% to over 90% and thorium-232.

The LTC had primary radionuclides that were of concern were fissile materials which included enriched uranium, thorium, plutonium, and U-233; transuranics; irradiated fuels and materials; and fission and activation products.

I'm going to talk a little bit about internal monitoring data, first with the processing facility, NNFD. fuel Uranium bioassay exists for all time periods. However, fluorometric analysis was used for the first operational period. Fluorometric analysis was used from 1956 I believe, '56 or '59, up through 1965. So it does cover that

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first operational period in 1959. And fluorometric analysis measures uranium by mass. Unless we have well-defined enrichment values this analysis is not really capable of measuring enriched uranium.

There is no bioassay data for thorium.

General area monitoring exists for uranium and some for thorium, but no breathing zone data are available. I should say, it says no breathing zone data are available for first operational periods, the two exception to the 1959 HASL report, the 1959 HASL study actually looked at two different operations in 1959, some pellet oxide fuel fabrication for Savannah River, and also some fuel fabrication for some Navy work. That 1959 study does contain breathing zone data. However, other than that there is no other breathing data for the first zone two operational periods.

Internal monitoring data at the

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LTC, we have uranium bioassay exists for all periods. We have one worker that indicated, or that we have routine bioassay analysis for uranium and fission products, that also includes some bioassay for plutonium and americium related to a 1969 incident. That is the only worker that we have that actually has any plutonium or americium bioassay data.

Okay, other than uranium and mixed fission products, again, bioassay sampling appeared to be incident specific. The one individual who we did have uranium -- or americium and plutonium bioassay was based on a 1969 incident, and they did continue that monitoring of that individual over time past that 1969 incident apparently to watch it clear.

No air sampling data for the first two operational periods could be directly attributed to the LTC. All the air data that we have right now that we have received is only for the fuel processing facility.

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External monitoring data, we have film badge data for both NNFD and LTC, again I've mentioned Landauer did the work for them for all operational periods. Neutrons from the records that we have, neutrons were not assessed at NNFD, because it was felt neutrons were not a significant exposure source. Neutron exposures were evaluated at the LTC during the period when commercial reactors were operating, which they operated up until 1986.

Source-term data, have we radioactive material inventory data that would NIOSH to place enable an upper bound on potential exposures to the wide array commercial and AEC radiological sources that could have been encountered at **BWXT** facilities.

Our feasibility of dose reconstruction, there are insufficient monitoring and source-term data from which to draw conclusions regarding potential magnitude

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of internal dose. Again I mentioned fluorometric analysis is inadequate for enriched uranium without specific knowledge of enrichment. We do have a range, we do have one campaign that we know that ran 5.9 % that we could tie workers to. However, all the other ones, we know there was a range of 4-90% and that would be a significant factor placed -- correction factor placed on on that bioassay.

Thorium exposures cannot be estimated for the 1959 period at NNFD. Only incident specific personal monitoring is available for plutonium and americium at LTC during the first two operations periods, and we have no breathing zone data for the first two operational periods.

External exposures, NIOSH believes there are sufficient monitoring and source-term data from which to draw conclusions regarding potential magnitude of external exposures. There are some questions about the

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1959 period and the early period of neutron exposures since they were operating with highly enriched uranium. But we believe that can be reconstructed.

Reconstruction of medical dose is likely feasible using claimant-favorable assumptions. And we will also use monitoring data personal that available for completing partial dose reconstructions.

We did, as I mentioned earlier, 36 interviews. Our interviews focused were because we actually as I mentioned earlier, there are two main plants, at least they would be defined as two main plants now, which would be the LTCand the NNFD. During our recognized that operations we there were feasibility limitations in for dose reconstructions, and we looked at different ways that we could possibly limit the Class. limit this Class to just the facilities which are actually -- there are

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buildings, A, B, C and D, that listed. Can we limit it to that? And separate out and do dose reconstructions for the NNFD? And vice versa. However, ultimately we came up with issues with both facilities. But the worker interviews did show us that some of the workers, we had received indications through one of the health physicists that workers were assigned specific buildings and they didn't move back and forth, and that may be true during the However we have indications later period. during some worker interviews that there were workers that moved from the LTC to NNFD and it is not apparent in their exposure monitoring records. And because of that we didn't even pursue limiting the Class because of that, even though there were infeasibilities tied to both facilities.

Okay, so our feasibility determination is that internal dose is not feasible, and external dose is feasible, for

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the period, we have two periods, January 1, 1959, through December 31, 1961, and January 1, 1968 through December 31, 1972. The third period of 1985 through 2001 we have not weighed in on that. We are still looking at that information, so this 83.14 only covers the first two operational periods.

The evidence we reviewed in this evaluation indicates that some workers in the Class may have accumulated chronic exposures through intakes of radionuclides and direct exposure to reactor materials. And NIOSH is specifying that health may have been endangered.

Our proposed Class, and I'm not going to read all of it, but it does cover the January 1, 1959 through the December 31, 1959, or from January 1, 1968 through December 31, 1972.

And again our recommendation: dose reconstruction is not feasible for those periods, and health was endangered.

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Questions.

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CHAIRMAN MELIUS: Questions for LaVon? Josie?

MEMBER BEACH: Hi LaVon, and you may have mentioned this and I didn't catch it, but the years 1960 to 1967, I know there was dose, I know there was still source there; why aren't those years covered?

MR. RUTHERFORD: Those are covered under the residual -- those are considered residual contamination years. We have not the residual contamination weighed in on period. What we would be doing, ultimately if we get to a point where we determine the residual periods we couldn't reconstruct, we would come back and do another 83.14. So we haven't weighed in on that in this evaluation. This evaluation only addresses those operational periods from 1959, which are considered -- if you remember from 1959 and 1968 through '72 we have to reconstruct all exposures, okay? For those residual periods

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1	of 1960 through 1967 we would only reconstruct
2	the residual exposure from the AEC activities.
3	The AEC covered activities. And that would
4	have been the production of the fuel for the
5	Savannah River reactors in 1959. So we
6	haven't weighed in on that. There is a
7	possibility down the line that that could
8	happen that we could recommend a Class for
9	that period.
10	MEMBER BEACH: Okay, thank you.
11	CHAIRMAN MELIUS: Just to follow
12	up on that, what is your process now going
13	forward?
14	MR. RUTHERFORD: Well, right now
15	we are still pursuing records through BWXT.
16	CHAIRMAN MELIUS: Okay.
17	MR. RUTHERFORD: And we are
18	working through them mainly because we feel
19	the 1985-2001 period we have much more data.
20	We have a lot, we have a lot more external
21	monitoring data. Also the analysis techniques
22	that took place with bioassay had changed. We

1	have alpha spec bioassay data. We also have
2	the enrichments are identified on the bioassay
3	card. So we have a lot more information
4	during that period. So but we are pursuing
5	the thing, the one thing that we are missing
6	is the understanding of their decision process
7	when to monitor and when not to monitor, so we
8	are trying to get that information from them
9	right now.
10	CHAIRMAN MELIUS: But would you
11	come back with a Site Profile type of document
12	or technical document?
13	MR. RUTHERFORD: I think that I
14	don't know that we have made a decision on
15	that.
16	CHAIRMAN MELIUS: I'm just trying
17	to follow up on what Josie was asking is, when
18	would we be considering, we as the Board, or
19	when
20	MR. RUTHERFORD: From the 83.14
21	there would be nothing that would happen, but
22	from us we have dose reconstructions that have

1	to be completed. So as we get this
2	information and we feel that we have enough
3	information to complete the dose
4	reconstruction, we will either complete the
5	dose reconstruction with all the information
6	inside it, or we will move forward with a Site
7	Profile.
8	CHAIRMAN MELIUS: Okay. Phil.
9	MEMBER SCHOFIELD: On your data,
10	your '85 through 2001, do you have any in vivo
11	counts?
12	MR. RUTHERFORD: Whole body
13	counts? Yes, for the later years we have
14	whole body counts for the later years. But
15	that is why we we have not weighed in on
16	the 1985 to 2001 period in this report.
17	MEMBER PRESLEY: LaVon have you
18	come up on any data on munitions fabrication
19	up there?
20	MR. RUTHERFORD: No, not any
21	no. Not that I can recall. Now I would have
22	to ask my ORAU counterpart if he noticed

1	anything in that. But no.
2	CHAIRMAN MELIUS: Brad.
3	MEMBER CLAWSON: LaVon, I'm
4	looking at this January 1 st , 1959 to December,
5	what if right halfway in the middle we have a
6	person that starts working there.
7	MR. RUTHERFORD: You are correct.
8	MEMBER CLAWSON: The 250 days is
9	what what is kind of bothering me, because
10	it goes right into another area that are going
11	to kind of be held in limbo.
12	MR. RUTHERFORD: I understand, and
13	that is a problem. I mean when you have a
14	short operational period of only one year, and
15	I did not review each claim to see if there
16	were claims affected by that. I probably
17	should have; I didn't think about it,
18	especially since I have had this happen to me
19	before. But there is definitely a possibility
20	that that could happen.
21	CHAIRMAN MELIUS: Again, I have a
22	related question which is, you mentioned

1	incidents. Are there potentially at this site
2	incidents that were maybe documented that
3	occurred but there may not be adequate
4	information on?
5	MR. RUTHERFORD: We are pursuing
6	that information as well.
7	CHAIRMAN MELIUS: Okay, good.
8	MR. RUTHERFORD: And I know this
9	isn't a great answer for you, Brad, but one
10	thing if we did ultimately determine dose
11	reconstruction wasn't feasible for the
12	residual periods, that would add on to that.
13	But we can't the problem, the Department of
14	Labor has defined it as 1959. We do use 1959
15	as January 1 to December 31 st . That is about
16	all we can do.
17	CHAIRMAN MELIUS: Not much else
18	in 1959. If you find anything let us know.
19	David Richardson or Paul Ziemer.
20	MEMBER ZIEMER: This is Ziemer.
21	I have a question. Am I on?
22	I'm never sure whether I put it on

mute or not. Okay, LaVon I have a question the air sampling. that relates to mentioned the general air sampling for uranium and some thorium, and the lack of breathing Just for clarity on the uranium zone samples. air sampling is there information on the enrichments involved?

MR. RUTHERFORD: Only for the 1959 study that was done by HASL.

Okay, and then on MEMBER ZIEMER: the -- okay that sort of takes care of it I guess I'll go ahead and ask the overall. rest of the question. Were the operations there such that the breathing zone samples would be markedly different than monitoring? There are many places where the area monitors and the breathing zone samples are quite similar, simply because they are not involved things that provide heavy -- things like grinding and so on that provide heavy localized air concentrations.

MR. RUTHERFORD: And actually if

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you look at the 1959 sample as an example, if you use that as indicative of all operations, then yes there is significant difference between the breathing zone data and the general area monitoring.

MEMBER ZIEMER: Okay, thank you.

CHAIRMAN MELIUS: David?

MEMBER RICHARDSON: Yes, I had one Something I'm struggling with. question. There are some sites where it's proposed that dose reconstruction can be carried out despite the lack of individual monitoring data, or even general area monitoring data. And yet here we are in a situation where there is individual quantitative estimates of intakes as you are saying, fluorometric analysis of mass, which takes us part of the way towards kind of individual dose reconstructions, and saying there is a key piece of you are information missing on enrichment of material being worked with. But it -- I mean it seems to me it's a difficult thing for me

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to reconcile. That here we don't feel comfortable either exploring information about the source-term, you're saying it's difficult to locate. But that we can't -- that there is not enough to work with when there is bioassay data actually available for workers, and yet another situation we are comfortable using surrogate data from other facilities. How do you reconcile that?

MR. RUTHERFORD: I think it's a If you look at different great question. operations, different operations we have data, lot more data on, Ι think. When we typically use surrogate data, especially if you look TBD-6000 and such, we've taken data from a number of facilities to derive, or to come up with what we feel are good surrogate data numbers. You have a -- I realize that there other enrichment facilities out are but I think in this process when we there, looked at, from the data that we had, we knew of one enrichment from the 1959 study, but we

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also had indications that enrichments ranged from 40% to 90% so that was one of our issues.

Our other issue was obviously that in 1959 was the thorium work.

MEMBER RICHARDSON: But when you couple this with a few plausible or even claimant friendly assumptions, and you couple those with bioassay data, is it not possible to bound something in a way that maybe more useful than kind of just stepping entirely and saying it's not possible to do reasonable dose reconstruction, is question. Or to take a year, for example, '59, where you said the work, the AEC-related work, was making fuel rods for Savannah River, knowing that, isn't that enough to kind of bounds level give you some on the of enrichment that would be used? Not completely unknown anymore.

MR. RUTHERFORD: We know that one. That one is part of the '59 study, so I know that one. The problem we have is because

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1	it is an AWE it's an Atomic Weapons
2	Employer, we have to be able to reconstruct
3	all exposures from that site. So I can
4	reconstruct in 1959 the work that was done for
5	the Savannah River reactor, but it's the
6	other enrichments that I don't know and that
7	occurred during that period.
8	MEMBER RICHARDSON: During that
9	same calendar year?
10	MR. RUTHERFORD: Exactly. And we
11	have went to BWXT and we've attempted data
12	captures, over the last three years we have
13	attempted to get additional information from
14	BWXT. We are back there again now trying to
15	get additional information.
16	MEMBER RICHARDSON: But for the
17	1,000-3,000 workers who were there, some small
18	fraction of them would have or am I right
19	or wrong, some small fraction of them would
20	have confirmed internal depositions based on
21	just the kind of bioassay data?

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MR. RUTHERFORD:

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It's not a small

1 fraction, no, I won't say that. We have from 2 there are actually a number the NNFD 3 positive urine samples. I didn't actually do 4 a percentage, but it's not a small sample. MEMBER RICHARDSON: I don't mean a 5 6 small number of people, I mean in percentage. 7 But I mean that is besides the point. Okay, I mean it's just still something that --8 This is Jim 9 CHAIRMAN MELIUS: 10 Melius, I think those are good questions, David, and I think the other way of looking at 11 12 that is, NIOSH also needs to be able to place 13 people within certain operations, so it's not only what operations you have information on 14 15 but how well can you place some people there 16 and how long they spent and what they did and so forth. 17 18 MEMBER RICHARDSON: Tn the 19 presence of bioassay data? 20 CHAIRMAN MELIUS: Not in the bioassay, I'm talking about the operations 21 22 side. Yes, you are right.

MR. RUTHERFORD: One of the persons we interviewed actually worked at the fuel processing facility for NNFD, the number of years, and actually they worked at the LTC, the actual technology center for a number of years, switched over to the NNFD for a year, I can't remember exactly how much time it was. And then switched back. The exposure monitoring records don't show any of that, they just show exposure monitoring The difficulty is while they were at uranium. LTC they were potentially exposed to a number of other radionuclides, and if we don't have information that makes it very difficult for us to reconstruct the dose. So even if we come up with, if we take generous methods on believe asked because me, Ι the question, I asked our internal dosimetrist why can't we just throw a factor on top of it and assume it's all 90 percent enriched uranium. And the problem you get is the exposures from that because of the factor you are throwing on

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1	top of it are not really plausible, they are
2	so high.
3	CHAIRMAN MELIUS: Any further
4	questions?
5	MEMBER ZIEMER: This is Ziemer,
6	let me just follow up on that. Because I was
7	having the same thoughts as I listened to
8	David's question. I assume in a sense you
9	might bound the uranium based on an assumption
10	of the high enrichment. Is it the thorium and
11	the other stuff that really causes the real
12	problem?
13	MR. RUTHERFORD: Yes, and we
14	you know we we just have no monitoring data
15	at all for the thorium. We have a few general
16	area samples, but we have no breathing zone
17	data.
18	MEMBER ZIEMER: So in principle
19	you could probably bound the uranium by making
20	a sort of worst case assumption.
21	MR. RUTHERFORD: Yes.
22	MEMBER ZIEMER: Because you have

the uranium bioassay but you don't know the enrichment, is that correct?

MR. RUTHERFORD: That's true.

MEMBER ZIEMER: Okay, thanks.

CHAIRMAN MELIUS: Now Mark

Griffon.

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MEMBER GRIFFON: I quess I was going to say the same thing as David and Paul. I mean the more convincing argument to me was also the thorium. And we have been sort of down this path with the uranium stuff before. But the question I had is if you determine '59 was an SEC, and then the next operational period was an SEC, what do you expect to learn in the next couple of months about that residual period in between that is going to allow you to reconstruct doses if you don't -if you can't reconstruct before, you can't reconstruct after, why don't you just roll the residual period in between now. I can see an argument made for the later residual period, but I can't understand how you would have any -- because we have had models before that back extrapolated or forward extrapolated.

MR. RUTHERFORD: Well, see, the difference is remembering that the residual period, the only thing we are required to reconstruct is the AEC activity. So the AEC activity was only the uranium portion of it. Now can we reconstruct the residual period assuming we take some data that we have the uranium data that we have and assume a TIB-0070 approach up to the next period. I don't know yet, so we couldn't -- if it was the same -- if we had to cover everything we would probably be in the same boat we are in now.

CHAIRMAN MELIUS: For the petitioner is not going to be speaking. She has opted not to. Do we have any other further questions? Do we have a motion? Yes, Phil.

MEMBER SCHOFIELD: Just one quick question. I'm kind of going off Mark's question there. Do you really know what

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1	occurred between those two time periods, '59
2	to '68. Was there any demolition that went
3	on? Was there any processing that you are
4	aware of that went on?
5	MR. RUTHERFORD: We haven't
6	completed that review, so I can't really say.
7	Again, we are looking at that period, but
8	this is only addressing those two operational
9	periods right now.
LO	CHAIRMAN MELIUS: Phil, that is
11	why I was asking for the, sort of what is the
L2	next document or next step that we how do
L3	we get involved, and what would we be
L4	reviewing at that point in time. And I think
L5	they will be pursuing this, and they will be
L6	coming back to us in some way in the future to
L7	address both that interim and the follow-up
L8	time periods. Yes.
L9	MEMBER LEMEN: How long will we
20	wait to hear back from them?
21	MR. RUTHERFORD: How long do we
22	wait to hear back? Oh. I'm sorrv. I didn't

1	know the question was to me, I didn't hear it.
2	I will get back with the Board on that. I
3	can give you a time period. I'm not sure when
4	our data capture efforts are going to be
5	complete at BWXT, and I'm not sure how long
6	it's going to take us to evaluate.
7	MEMBER LEMEN: Are we talking
8	months, years?
9	MR. RUTHERFORD: I should be able
10	to give you an update at the August meeting.
11	CHAIRMAN MELIUS: Wanda.
12	MEMBER MUNN: I don't have a
13	question. I'm prepared to make a motion.
14	CHAIRMAN MELIUS: Well, go ahead.
15	MEMBER MUNN: I move that we
16	accept the NIOSH proposed SEC Class for all
17	Atomic Weapons Employees who worked at BWXT,
18	Inc., in Lynchburg, Virginia, during the
19	periods from January, 1, '59 through December
20	31, '59, or from January 1, 1968 to December
21	31, 1972.
22	CHAIRMAN MELIUS: Do we have a

second for that?

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MEMBER CLAWSON: Second.

CHAIRMAN MELIUS: Second from Brad. Again, Wanda, I would ask if you would listen and maybe accept a friendly amendment.

MEMBER MUNN: I might.

CHAIRMAN MELIUS: I know, that's why I'm asking.

Dear Madam Secretary, the Advisory Board on Radiation Worker Health, the Board has evaluated SEC Petition 00169 concerning BWX Technologies, workers at Inc., Lynchburg, Virginia, under the statutory requirements established by EEOICPA incorporated in 42 CFR Section 83.14. Board respectfully recommends Special Exposure Cohort SEC status before all Atomic Weapons Employer employees who worked at Technologies, Inc., in Lynchburg, Virginia, from January 1, 1959 through December 31st, 1959, or from January 1st, 1968 through December 31st, 1972, for a number of work days

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aggregating at least 250 workdays for either solely under this employment or in combination with workdays within the parameters established for one or more other Classes of employees included in the Special Exposure Cohort. The recommendation is based on the following factors: people working at facility during the time periods in question worked on fuel fabrication, uranium recovery and commercial reactor and laboratory operations related to nuclear weapons The NIOSH review of available production. data found that they lack adequate source-term process and monitoring data in order to be able to complete accurate individual dose reconstructions for internal radiation doses for employees at this facility during the two time periods in question. The Board concurs with this determination, agree that NIOSH determined that health may have been facility workers. endangered for these BWX The Board concurs with this determination.

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1	Based on these considerations and
2	the discussions held at our May 19 th through
3	21 st , 2010, Advisory Board meeting held in
4	Niagara Falls, New York, the Board recommends
5	that this Special Exposure Cohort petition be
6	granted. Enclosed is documentation from the
7	Board meeting where this Special Exposure
8	Cohort Class was discussed. Documentation
9	includes transcripts of the deliberations,
10	copies of the petition, the NIOSH review
11	thereof, and related materials. If any of
12	these items are unavailable at this time they
13	will follow shortly.
14	MEMBER MUNN: That is a
15	significant amendment, and as a matter of fact
16	is an order of magnitude greater than the
17	original motion, but it can be accepted.
18	CHAIRMAN MELIUS: Orders of
19	magnitude.
20	And I think Emily has some
21	clarification. There are a couple of typos in
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there.

1	MS. HOWELL: I wasn't even going
2	to mention the typos. Dr. Melius had
3	previously asked me if we were creating any
4	sort of a loophole for people who had worked
5	during both of these covered periods that are
6	in this Class Definition and needed to combine
7	work during those periods to get to the 250
8	day threshold. Normally we have a single
9	period, and so the aggregating language allows
10	that. In this instance, since it's a single
11	Class with two different periods we were just
12	going to suggest we don't know that it
13	would be a problem. It would be in DOL's
14	interpretation, but if you put an "and slash
15	or" that should take care of it.
16	CHAIRMAN MELIUS: Further friendly
17	amendments? Thank you, Emily, for that.
18	Any further discussion or
19	questions on that? If not, go ahead Ted.
20	MR. KATZ: Thank you. I will do
21	the roll call in reverse order. And let me

note at the head of this that Dr. Poston had a

1	conflict for this site, he did not participate
2	in the discussion nor will he participate in
3	the vote. So beginning with Dr. Ziemer?
4	MEMBER ZIEMER: Yes.
5	MR. KATZ: Mr. Schofield.
6	MEMBER SCHOFIELD: Yes.
7	MR. KATZ: Dr. Roessler.
8	MEMBER ROESSLER: Yes.
9	MR. KATZ: Dr. Richardson.
10	MEMBER RICHARDSON: Yes.
11	MR. KATZ: Mr. Presley.
12	MEMBER PRESLEY: Yes.
13	MR. KATZ: Ms. Munn.
14	MEMBER MUNN: Yes.
15	MR. KATZ: Dr. Melius.
16	CHAIRMAN MELIUS: Yes.
17	MR. KATZ: Dr. Lockey.
18	MEMBER LOCKEY: Yes.
19	MR. KATZ: Dr. Lemen.
20	MEMBER LEMEN: Yes.
21	MR. KATZ: Mr. Griffon.
22	MEMBER GRIFFON: Yes.
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1	MR. KATZ: Mr. Gibson.
2	MEMBER GIBSON: Yes.
3	MR. KATZ: Dr. Field.
4	MEMBER FIELD: Yes.
5	MR. KATZ: Mr. Clawson.
6	MEMBER CLAWSON: Yes.
7	MR. KATZ: Ms. Beach.
8	MEMBER BEACH: Yes.
9	MR. KATZ: Dr. Anderson.
10	MEMBER ANDERSON: Yes.
11	MR. KATZ: So all vote in favor
12	with one recused, that's 15 members in favor.
13	The motion passes.
13 14	The motion passes. CHAIRMAN MELIUS: Very good. Dr.
14	CHAIRMAN MELIUS: Very good. Dr.
14 15	CHAIRMAN MELIUS: Very good. Dr. Poston, you are allowed to join us again.
14 15 16	CHAIRMAN MELIUS: Very good. Dr. Poston, you are allowed to join us again. The next item of business on our
14 15 16 17	CHAIRMAN MELIUS: Very good. Dr. Poston, you are allowed to join us again. The next item of business on our agenda is the surrogate data criteria. And
14 15 16 17	CHAIRMAN MELIUS: Very good. Dr. Poston, you are allowed to join us again. The next item of business on our agenda is the surrogate data criteria. And you all should have received a document
14 15 16 17 18	CHAIRMAN MELIUS: Very good. Dr. Poston, you are allowed to join us again. The next item of business on our agenda is the surrogate data criteria. And you all should have received a document labeled final draft criteria for the use of

the Surrogate Data Work Group met on May 13th, of -- by conference call, and we discussed a draft. I will indicate that Dr. Jim Lockey was unable, he was traveling at the time so he unable to participate, but the members of the Work Group did. And we have I had made several some further discussions. changes based on the discussions that we held at the last Board meeting. And it included changes, then we made а couple clarifications I would say since that time based on the Work Group meeting, there have just been some minor changes since then. The Work Group Members that were present I think were in general agreement with this document, and felt that we should bring it back to the Board for additional comment or adoption by the Board.

I think the -- actually most of the changes that have been made since our last meeting that were designed to clear up the confusion with some of the wording, and I

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think that is that. I think Dr. Richardson,
David, had made some comments on the hierarchy
data, and I think we tried to clarify that. I
think we all had some differences on how we
interpreted what was higher or lower in the
hierarchy, and his comments were helpful as
were others. There were again some
clarifications.

I think that the last paragraph I
had also changed around. I think we had used

I think that the last paragraph I had also changed around. I think we had used the words, rarely used, or something like that, and I think we have clarified that. So I'll open it up to comments from other Board members. Yes, Wanda?

MEMBER MUNN: There was one clerical nit in the paragraph one, hierarchy of data. We did not have a closing paren.

CHAIRMAN MELIUS: Jim.

MEMBER LOCKEY: Unfortunately I was on a plane when this was happening, but under scientific plausibility, when I was reading that, this last few days, and I looked

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at it again today, when I look at that, Jim, I guess I thought I would ask you, are the assumed models scientifically appropriate? And then are you saying in the next sentence that these models have to be validated through actual monitoring data? Or is that more appropriate to say validated where feasible? Some of the things you are just not going to be able to recreate in this day and age.

CHAIRMAN MELIUS: No, I think the -- what we were trying to do with that is identify issues. So one of the issues that one would discuss is not -- there is no absolute criteria that one has to have a validated model. It's one of the questions that would look at in making the you evaluations.

MEMBER LOCKEY: I would say that if you can validate a model by giving actual data you should do it. But I would propose putting the word in, validate where feasible.

Which means that I can't go back and recreate

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exposures that happened in a laboratory setting in 1945 or 1950; it's just not feasible under the current circumstances. But where it is feasible I think that is a good criteria.

CHAIRMAN MELIUS: Actually your other comments, that Wanda had relayed -- I think you had left the meeting, the previous meeting, at some point. I understand now.

Any other comments or questions?
Dr. Lemen.

MEMBER LEMEN: I know that you have been considering this for many many years, and I've just come on the Board. But I still question, and I understand that this document has restricted the use of surrogate data, fairly strongly, but I still question the use of surrogate data in a compensation program. And I really have a problem with the use of it. I think in doing epidemiology, yes, you might want to use surrogate data; you might want to make some assumptions about it.

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But when you are dealing with a compensation program I don't think it's appropriate. Now I may be the minority on this Board that feels that way, but that is my opinion. I feel I have to express it.

CHAIRMAN MELIUS: Ι think my response to that is, I think as we have found on the Board before. Before you joined us, and with other members in the past, we do have differences of opinion, and we are not going to always reach a consensus, particularly on a general area, because there are lots different uses, potential uses of surrogate data, and lots of different circumstances. As Jim just mentioned in some cases it may be feasible to do a test on the plausibility of a In other cases it may not, and I think model. we are all going to judge those individually, and then reach whatever consensus we can on And what I think we've found in trying forward criteria for making to put evaluations, the first thing that is

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important is identifying the issues that we think need to be addressed, that NIOSH should address in presenting to the Board the information we are going to need to make an assessment. And then I think we are going to make that assessment.

So I think we've found it hard and will continue to find it hard to come up with absolute criteria. There must be a model. There must be validation of a model, something It's going to depend like that. circumstances, how the data is used. We have also found there is, I think, a criteria, a continuum, on the use of surrogate data in this program and the actual use or application of something we are going to look at. frankly I don't think we could agree absolute criteria as a Board, and I'm not sure it's that helpful to have us do it. As long as we have a framework that hopefully we can agree on for how it should be considered, what information we want available to us. That

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would then guide NIOSH and our contractor in producing documents and information for And I think that's where we going forward. And I think we will have to see as we Maybe we can refine this as apply it. apply it. We'll just have to see. I think we taken it fairly far in terms identifying the issues. But that would be my Jim, do you have a follow-up? comment.

MEMBER LOCKEY: I think that this document really sets a format to be followed for surrogate data, and then also the system does allow, when we are looking at dosimetry, we are looking at the 99 percent confidence intervals, which gives you a broad margin of safety to encompass potential exposure. think when you look at them together, I think friendly, certainly, it is towards claimants. And I think having it outlined --I think this is a great document, having it outlined like this is a good roadmap to follow.

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CHAIRMAN MELIUS: Henry.

MEMBER ANDERSON: I just wanted to probably underscore what you already said, that I really think this is a useful guidance document that hopefully answering all these questions, NIOSH goes through that in advance, so rather than bring proposals to the Board and then having us disagree with NIOSH this should, I would think, bring it closer to an agreement. But I just want to be sure. you said is that the quidance, or this is strictly to be used to guide, criteria, kind of, the discussion, but the decisions will be one at a time on the specific site, and you can never have something that is enough for all the gray zone areas. So this isn't going to eliminate all disagreements, but I think it will help people begin to think about when to bring surrogate data to the Board for use.

CHAIRMAN MELIUS: And how to bring it, I think that as much as anything --

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And I think it's going to take us some time to do that. Yes, Dick.

MEMBER LEMEN: I just had one follow-up, and that was, how are we assured that NIOSH will take this under advice, and follow these? And the second point is, I'm not sure I understand, the Type II, exactly what you are talking about here.

CHAIRMAN MELIUS: Let's do number one first, and I think that NIOSH -- first of all, why do we think these are helpful, and address did NIOSH it. how These relatively close to the criteria that NIOSH has developed and published. I can't remember the number of the document, but something --IG-004. And that document, like other documents they have is undergoing change, as they better understand and apply -- actually next discussion of the Hooker our Electrochemical facility, I think they will have some comments on how they are approaching surrogate data in that, and that's why we sort

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of paired up the agenda in this way.

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Secondly, what this document meant to do is to guide the Board on what issues we are going to be looking for, and looking at, when we are reviewing the use of surrogate data, mainly in the Type Ι situation, the use of it for dose reconstruction or potential use for reconstruction.

And so I think as we found in the past that is helpful for NIOSH; it speeds the They understand what process up. looking for. Our previous example, this is with the SEC evaluation which I think is, we were struggling, were having problems as a Board, and working with NIOSH on evaluation of and that document I think provided a SECs, focus to what is now in the Evaluation Reports, what issues are addressed. And it facilitates decision-making, our facilitates how NIOSH approaches -- we are not just saying, well, you forgot this, or you

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didn't do that. Send it back. They know ahead of time. So we are here to provide independent advice to the Secretary and to NIOSH on this program and certain parts of this program, and I think it helps to have a document that guides our - how we are going to approach this particular in this case surrogate data.

Second question, Type I and Type II, they do go together. I think Type I data in a simple sense is where the surrogate data directly basis for used the as reconstruction of individual doses or parts of individual doses. There is Type II in general where it's used is sort of in supporting information for part of that dose reconstruction. The two blend together, and it's hard, I think the more we've looked at it -- it may be hard to make that separation, but primarily we are trying to focus on the Type I.

John Mauro at SC&A has done a

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1 document, which we all should have received, 2 where they have gone through, at least Site 3 Profiles I believe, and documented where it's 4 used and how it's used, and with 5 description, not only Type I, Type II, but 6 also a description of where it's used in 7 different Site Profiles, and I think you will see how it's -- sort of what the range of that 8 So if that helps to address --9 Any 10 other questions or comments? MEMBER RICHARDSON: This is David 11 12 Richardson. 13 CHAIRMAN MELIUS: Sorry, David, go ahead. 14 15 MEMBER RICHARDSON: I think this 16 is a very useful document, and I think it lays out some good principles. One issue I have 17 18 been thinking about relates to something which 19 isn't quite made explicit here, which is 20 something we have talked about several times today, which relates to not so much to the 21

kind of similarity of exposures or exposure,

possible exposure conditions, but the variance
in the exposures within a workplace, and the
difference between we talked about, for
example, what is the relationship between area
monitoring and breathing zone monitoring?
There are some situations in which exposure
conditions are relatively homogenous in a
workplace, and you might think that is kind of
a representative value, but if you take the
mean, the median, or a percentile, it is going
to give you a good sense of the exposure
conditions for any individual you draw from
that workplace. And then there are so that
the variance is relatively small. And it's
well characterized. And then there are places
where you have got real tails on the
exposures. Because the exposure conditions
are can be quite variable. And we had this
example at Mound, right, where you wouldn't be
for this worker in SW-19 room C, you really
don't feel comfortable saying that you can
reconstruct that person's dose if they come

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down with lung cancer, because their exposure
is really different. And I started thinking
about that, we had surrogate data for Building
SW, or S/WR at Mound, you might say kind of
the average exposure, the 95 th percentile,
that doesn't really tell you very much in a
situation where you've got a long tail on the
exposure distribution. So to be claimant
friendly, if that means for any worker you are
going to develop an estimate which is either a
good estimate of their exposure or an
overestimate, but isn't a substantial
underestimate of their true exposure, then you
have to get some idea about is this surrogate
data capturing those tails in a fair way? And
I think that is really hard. And it sort of
made me think is there I guess item six
here, aside from plausibility and site
processes, which is something about surrogate
data may be more useful in situations where
there is relatively low variance exposure, as
opposed to situations where the judgment is

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that there is a lot of heterogeneity in Where somebody in some exposure. locations really has a potential for peak exposures, which aren't all captured by the median or the 95th percentile. So there may be better and worse situations. If exposure intensities are kind of -- by some process which is repeatable and fairly consistent with relatively little variance because of kind of the process generating the exposure, that may be easier for us to kind of intuit that where you could take information from somebody else who is working with a similar process in a different location. But when it gets really complicated, we may want to be a little more cautious in using surrogate data.

CHAIRMAN MELIUS: I think that is a good point. I think it is more -- my reaction is, it's also a more general point about the entire program. I think that is what we spend a lot of time struggling with in individual Site Profile reviews is, which is

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the best way of characterizing the distribution exposures, and of therefore, making the proper assignment of that. The coworker model comes up all the time, and it is something important. Your point earlier on the hierarchy, maybe to think about that, and these criteria aren't usually absolute by themselves, but it's sort of the context that you are dealing with, and how the general quality of the data, and maybe distribution is one part of the quality of the data, how good the data is, how tightly does it describe the range of exposures.

MEMBER RICHARDSON: And I think the more, as you can narrow things down to -- an epidemiologist would not like to have surrogate data for a plant, but maybe for a job title and a location and a period, you may feel well, there you have started to bound how extreme those tails are going to be or what the variance is going to be, possibly, maybe not. But to the extent that surrogate

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data can be coupled with other information that lets you reduce the variance of the two distributions that you are trying to think about as being comparable.

CHAIRMAN MELIUS: The other criteria, I thought about at one point, I just think it's very hard, is well, how much of the dose for the worker is going to be made up from the surrogate data, and that is really a question of the distribution also. The two are related and so forth, and how do you judge that if it's a difference between a 49 percent and putting them over 50 percent it's still an important component of the dose situation. But again I think that is also general to the whole program; it's not just surrogate data. And for that reason I sort of left it out of the surrogate data criteria. We have to think about what you are proposing. Certainly I think we all agree that it is important, do we put in surrogate data, how do we put it into our other ways of evaluating the methods used

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1 for this program? 2 Anybody else have comments 3 that? Dr. Ziemer? This is Ziemer, 4 MEMBER ZIEMER: 5 I have a comment. 6 CHAIRMAN MELIUS: Go ahead. 7 MEMBER ZIEMER: I think you are exactly right, Jim, I think we have exactly 8 the same issue whether it's surrogate data or 9 10 real data, in terms of those distributions. don't think there is any implication in this 11 12 document that the surrogate data implies that 13 taking midpoints or averages we are anything like that. We always have to deal 14 15 with that distribution and the tails, whatever 16 the data distribution is, surrogate or real. I think the point is well made, but I think 17 it's broader than just surrogate data. 18 19 MEMBER RICHARDSON: Yes, Ι wouldn't think we would have to deal with the 20 tails. But my understanding of how you 21

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the

way

this,

implement

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it's

been

implemented, is to propose that there is a value or a distribution of values, that you are going to assign. And I believe some of proposals have been, well, will claimant friendly and rather than say that the exposures for people where we don't have information on their actual exposures is going to be comparable to the facility, and we'll 95th take the percentile of the dose distribution at that surrogate facility and we will treat that as the annual dose rate for workers at the target facility.

MEMBER ZIEMER: Yes, and I think you would have to make the case in each facility as to why you did it a certain way.

MEMBER RICHARDSON: Right, so that is what I was getting at, the issue of, this is where the tails become important. Because you don't have the true data. When there are measurements made, you also have tails, but you for an individual you can place them in the tail or not. But here we are

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going to have a play a game which is not necessarily claimant friendly, if they happen to be one of the true people in the tail and yet we impute them at some point farther down the percentiles in the distribution.

CHAIRMAN MELIUS: Any other comments?

I would propose, if people are agreeable is, with the one change from Dr. Lockey that we adopt these -- this document for now. We will continue to work on it, and David, you and I can -- maybe Paul or others can think about how incorporate point we number six. There is no reason we couldn't add to it, and we may well want to hopefully change and improve it as we go along. also would like to get some closure at this point simply so there is a document that NIOSH and others that are involved in the program and our contractor can maybe start to utilize with a little bit more certainty than in the past in moving forward. So I have a proposal.

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1	MEMBER ZIEMER: Doesn't this come
2	as a recommendation from the Work Group?
3	CHAIRMAN MELIUS: It is.
4	MEMBER ZIEMER: I think that
5	makes it a motion then.
6	CHAIRMAN MELIUS: It's a motion,
7	thank you, our parliamentarian. That was your
8	appointment, Paul, remember?
9	MEMBER ZIEMER: Thank you.
10	CHAIRMAN MELIUS: And grammarian
11	is it? Do I have a second to that? I think
12	we need a second?
13	MEMBER CLAWSON: I second.
14	CHAIRMAN MELIUS: Further
15	discussion? All in favor?
16	(Chorus of ayes.)
17	CHAIRMAN MELIUS: Opposed.
18	Why don't we call the vote?
19	MR. KATZ: Okay, so just a roll
20	call vote. Dr. Anderson?
21	MEMBER ANDERSON: Yes.
22	MR. KATZ: Ms. Beach?

1	MEMBER BEACH: Yes.
2	MR. KATZ: Mr. Clawson?
3	MEMBER CLAWSON: Yes.
4	MR. KATZ: Dr. Field?
5	MEMBER FIELD: Yes.
6	MR. KATZ: Mr. Gibson?
7	MEMBER GIBSON: No.
8	MR. KATZ: Mr. Griffon?
9	MEMBER GRIFFON: Yes.
10	MR. KATZ: Dr. Lemen?
11	MEMBER LEMEN: No.
12	MR. KATZ: Dr. Lockey?
13	MEMBER LOCKEY: Yes.
14	MR. KATZ: Dr. Melius?
15	CHAIRMAN MELIUS: Yes.
16	MR. KATZ: Ms. Munn?
17	MEMBER MUNN: Yes.
18	MR. KATZ: Dr. Poston?
19	MEMBER POSTON: Abstain.
20	MR. KATZ: Mr. Presley?
21	MEMBER PRESLEY: Yes.
22	MR. KATZ: Dr. Richardson?

1	MEMBER RICHARDSON: Yes.
2	MR. KATZ: Dr. Roessler?
3	MEMBER ROESSLER: Yes.
4	MR. KATZ: Mr. Schofield?
5	MEMBER SCHOFIELD: Yes.
6	MR. KATZ: Dr. Ziemer?
7	MEMBER ZIEMER: Yes.
8	MR. KATZ: So the total is 13 in
9	favor, two noes, and one abstain. The motion
10	passes.
11	CHAIRMAN MELIUS: We have until
12	3:15, but I think some of you may want or
13	deserve a break since it was put off late
14	afternoon. What I would like to do, we have
15	to schedule a February meeting, 2011 meeting.
16	So Ted, do you want to talk?
17	MR. KATZ: Sure. Okay, very
18	good. So we are scheduled through November
19	for meetings. And it would be good at this
20	point, because some locations are harder than
21	others to book, well, we might as well while

we are at it schedule a teleconference too,

which will be tougher. The right sort of date
range for the next face to face after November
is February. We've got some constraints,
which makes stretches it a little bit
because we have some unavailability already of
staff. So the first good week for the Board
to possibly meet in February would be February
the week of President's Day, which is the
week starting the 22^{nd} of February, that is
the Tuesday I guess afterwards, 22^{nd} , 3^{rd} , 4^{th} ,
5 th , I believe is right, so that is one
possibility. And that would be keeping it
relatively tight in terms of stretching out,
because this is a slightly longer period than
other quarters, already. Otherwise is the
week of February 28 th , so look on your
calendars for that.

CHAIRMAN MELIUS: Anybody have major conflicts the week of the $21^{\rm st}$? I guess the holiday is Monday.

MR. KATZ: Right, the 21^{st} is

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1	President's Day.
2	MEMBER MUNN: So you are looking
3	that far ahead in February because?
4	MR. KATZ: Because some places
5	are harder to book. And so we get started
6	actually quite early.
7	MEMBER MUNN: But I am just
8	wondering why not earlier in February?
9	MR. KATZ: Oh, okay, there are
10	constraints, people can't do it the week of
11	February 7^{th} or 14^{th} . So we really have to
12	start the end of January, and that leaves no
13	time between the November meeting and that.
14	MEMBER RICHARDSON: And how many
15	days is this meeting?
16	MR. KATZ: Three days is what we
17	planned for. So it could be 22^{nd} , 3^{rd} , 4^{th} ,
18	5 th .
19	MEMBER RICHARDSON: Third,
20	fourth, fifth is best for me, or next week.
21	MR. KATZ: I'm sorry, David, I
22	couldn't hear what you said.

MEMBER RICHARDSON: The 23rd, 24th, 1 and 25th would be better for me, or else the 2 3 next week. 4 MR. KATZ: Thank you. CHAIRMAN MELIUS: 5 Let's talk, Jim 6 Lockey is waiting to hear back. Let's talk 7 locations. MR. KATZ: So I have some ideas, 8 but others could certainly have others. 9 10 Savannah River site, so Augusta, we have met there for Savannah River site. That gives a 11 12 good bit of time to get a lot of work done for Savannah River site. So I don't know, it 13 might not be bad timing in that respect. And 14 generally we try for February to aim for a 15 16 place where we know weather is not going to keep us from there. And Augusta, we'd be 17 pretty safe there, it won't be balmy but it 18 19 won't be terrible probably. 20 other possible locations, The Tennessee, we haven't been there in quite some 21

time, in Oak Ridge, but we have quite a bit of

1	activity related to Tennessee still to do with
2	the Board. So that is a possibility, and I
3	think it's not too hard to get into Oak Ridge
4	even in the winter. Those are the two best
5	options actually. And I am open to other
6	suggestions.
7	MEMBER MUNN: I would prefer Oak
8	Ridge.
9	CHAIRMAN MELIUS: How about other
10	locations? Anybody think of something we're
11	missing? They want some place like Idaho in
12	August.
13	MEMBER BEACH: It seemed like
14	there was some talk about Florida?
15	MR. KATZ: There is, Pinellas,
16	but that's pretty narrow opportunity.
17	CHAIRMAN MELIUS: It's narrow,
18	and frankly President's Day week in Florida is
19	tough. There's a lot of
20	MR. KATZ: Holiday spot.
21	MEMBER CLAWSON: I'd like Oak
22	Ridge.

1	MEMBER GRIFFON: I would expect
2	we'd be pretty far along on the Savannah River
3	SEC petition by then. It might make sense to
4	be there again.
5	CHAIRMAN MELIUS: I think in
6	terms of I don't think we have any active
7	petitions at the Oak Ridge right now.
8	MEMBER CLAWSON: It could be
9	Nashville.
10	MEMBER ANDERSON: I'd hold it in
11	Knoxville. I wouldn't hold it in Oak Ridge,
12	just in case the weather does turn bad.
13	CHAIRMAN MELIUS: Right. But I
14	think Savannah River
15	MR. KATZ: So is that our first
16	choice?
17	CHAIRMAN MELIUS: First choice,
18	Savannah River, second choice, Oak Ridge, and
19	then why don't we take a break and just before
20	the start we will try to settle the date.
21	MEMBER LEMEN: Is Savannah River
22	Augusta then?

1	MR. KATZ: Yes, it would be
2	Augusta, I think that is the best location.
3	MEMBER LEMEN: That's fine with
4	me.
5	CHAIRMAN MELIUS: Good, let's
6	take a break.
7	MR. KATZ: Dr. Lockey says it's
8	good for him.
9	CHAIRMAN MELIUS: Then plan the
LO	23 rd , 4 th , 5 th and fifth in Augusta. And
L1	I will say up front we will try to make it a
L2	2-1/2 day meeting so people will be able to
L3	get out on Friday for everybody. And the
L4	meeting may start at 5:00 in the morning or
L5	something. We will take a break, come back at
L6	20 after. We will get started. We have a
L7	petitioner.
L8	(Whereupon, the above-entitled
L9	matter went off the record at 3:00 p.m. and
20	resumed at 3:22 p.m.)
21	CHAIRMAN MELIUS: If everybody
22	else could get seated, and quiet down we'll

1	get started. We will discuss Hooker. We have
2	one more administrative item to do, that is
3	our Board call, which would be ideally
4	sometime in January. And we have some people
5	that aren't available in the latter two weeks
6	in January, so we are talking about ideally
7	the week of the 10 th of January. I suspect
8	that will be a relatively short call, just
9	given the post-holidays and then given, I
10	don't think there will be a lot to update on,
11	but we should try to at least have it on the
12	books; we do need it. So I'm just going to
13	throw out Wednesday, January 12 ^{th.}
14	MEMBER MUNN: Very good.
15	CHAIRMAN MELIUS: Good, okay.
16	Board call, January 12 th , 11:00 a.m.
17	MEMBER LEMEN: 11:00 a.m.
18	CHAIRMAN MELIUS: Eastern.
19	Eastern time.
20	And first presentation now will be
21	on Hooker Electrochemical. And we have Dave
22	Allen from NIOSH, and I believe we have

1	petitioners here.
2	HOOKER ELECTROCHEMICAL (NIAGARA FALLS, NY)
3	SEC PETITION
4	MR. ALLEN: Most of you have
5	already met me before, but for those who
6	haven't, my name is David Allen. I'm a health
7	physicist with NIOSH, and as Dr. Melius
8	mentioned I am here to present the Evaluation
9	Report for the Hooker Electrochemical Special
LO	Exposure Cohort.
L1	CHAIRMAN MELIUS: Dave, can you
L2	speak into the mike?
L3	MR. ALLEN: A little closer. Is
L4	that better:
L5	CHAIRMAN MELIUS: That's better,
L6	yes.
L7	MR. ALLEN: Okay, a little
L8	background to start with. Hooker
L9	Electrochemical was classified as an Atomic
20	Weapons Employer from 1943 to 1948. There is
21	a residual contamination period that goes from
22	the end of the contract period until the end

of 1976. The Hooker primarily produced chemicals for Manhattan Engineering the District during World War ΙI and shortly One in particular, P-45, produced a waste product of concentrated hydrocholoric acid. Αt some point somebody from the Manhattan Engineering District realized they could take that waste product, put it together with product а waste from an electrometallurgical, and concentrate а uranium content in that waste product produce a material that could be -- that they could recover uranium from.

So that process was added as supplement to the P-45 contract. In order to perform this operation the equipment housed building in а that was built specifically for this, a small cinder block building. It was built by a subcontractor and turned over to Hooker on July 11th, 1944. P-45 operation including the supplement that covered the concentrating of chloride maq

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ended by January 15th of 1946.

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The incoming material, the magnesium fluoride contaminated with uranium, had a uranium content of approximately .2 percent uranium by mass. The concentration effort brought it up to between one and two percent uranium by mass.

The petition for Hooker Electrochemical, the SEC petition, submitted March 6th of 2009. The proposed Class at the time was for all the operators the laborers in furnace The room. building that I said housed this equipment did not have a furnace room. There were other furnace rooms within Hooker Electrochemical and other buildings, but this particular process did not have one. Because of that we did not qualify the petition for evaluation. Later the petitioner did revise the Class to all employees at Hooker Electrochemical, and radiation based monitoring the on no petitioner revised the Class on September

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 $26^{\rm th}$, re-qualified it then on October $16^{\rm th}$, and the Evaluation Report was issued on May $3^{\rm rd}$.

The proposed Class as I said now is all employees who worked in any location at Hooker Electrochemical, during the operational period and during the residual period, the Class that NIOSH evaluated is the same.

Sources of available information include Appendix AA to Battelle-TBD-6001 and that is essentially an appendix that describes our dose reconstruction methodology for Hooker Electrochemical. That also points to, and some of the other information we have, is some ORAU Technical Information Bulletins; various documents on the Site Research Database; we did interview some former employees; and we have some information in case files within our Claim Tracking System.

As I said the basis for the petition was an affidavit indicating to the best of the petitioner's knowledge there was never any internal or external monitoring at

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Hooker Electrochemical. We found nothing to contradict this. It's pretty consistent with what we found in the documentation.

mentioned, the As Ι dose reconstruction methodology, we put together an appendix for TBD-6001, it's Appendix AA, TBD-6001 does use surrogate data for internal Surrogate data for TBD-6001 is dosimetry. broken up into various tasks or operations. The operation we chose for Hooker Electrochemical was scrap recovery.

A little more detailed process of what occurred at Hooker involving radioactive material was contaminated magnesium fluoride was received from Electro Metallurgical in 500 pound barrels. The barrels were dumped onto a conveyor, and the conveyor brought them to a digestion tank. The waste hydrochloric acid from the P-45 process was then added to the digestion tank and delivered with water up to a pH of 4.0. The tanks were agitated for 20 hours, and then about once every two days the

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liquid was decanted from the tank, and additional hydrochloric acid and water were added, and the process started over.

What this did is essentially dissolve some of the mag fluoride leaving uranium behind and therefore concentrating the mag fluoride.

At the end of the digestion this slurry was neutralized and then pumped to a filter press where it was filtered and the filter was rebarreled and sent back offsite.

The recovery scrap process described in TBD-6001 involves calcining uranium scrap in a furnace; digesting that scrap in acid; precipitating the uranium; and then filtering that precipitate. The digesting and the filtering are very similar what occurring at Hooker to was Electrochemical. The primary airborne causing though, operation, in the scrap recovery the calcining. process was And by calcining operation I mean also loading and

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unloading the furnace, as well as simply heating the material.

The other steps were not very - did not cause a great deal of airborne, and it makes sense that it was either a solution or a material with a high moisture content. You don't get a great deal of dust from the rest of the operation.

Comparing the materials that were at Hooker versus what was in scrap recovery, the incoming material at Hooker was approximately .2 percent uranium by weight. The incoming material for scrap varied quite a bit. It could be uranium metal such as metal turnings, et cetera. But it was also low grade ores and slag, dross various other scrap materials that would occur during the processing of uranium. It was very precious material, so they always try to recover uranium from any other waste product.

The outgoing material at Hooker was concentrated to one to two percent uranium

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by weight, whereas for scrap recovery the intent of recovering that scrap was to produce a uranium compound that could then go further down the stream and be used in the weapons program. So that was high grade uranium compounds that was the output of the scrap recovery process.

Because of those reasons it seemed that the scrap recovery process in TBD-6001 appears to be a comparable match, probably a favorable or bounding match, to the materials and the process that occurred at Hooker Electrochemical.

This was labeled alternative to surrogate data, because if we chose not to use surrogate data, it does not mean that we could estimate Hooker not. t.he dose at Electrochemical. There is a well source-term and a well defined process, and it is open to the idea of modeling. Any time you model something you obviously have to rely on some assumptions, and there are some inherent

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uncertainty associated any time you are relying on assumptions.

When we looked at this at Hooker Electrochemical we decided that the scrap recovery was more specific than the model that might be more generic. It was a reasonable fit at Hooker, to the operations at Hooker, therefore we decided that surrogate data was a more robust analysis than any kind of modeling we could come up with.

And as the last agenda item was surrogate data, we are aware that the Board is reviewing the use of surrogate data. We did evaluate our use of surrogate data in Hooker based on IG-004, and I believe that comes reasonably close to your draft items; I'm not sure about that, how it might have been changed here recently. But we did evaluate the use; we did decide that this was a more scientifically sound model than we were going to come up with.

I have a couple of dose

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reconstruction examples. One being an employee from 1944-45, the demographics for this case, a male born in 1927, diagnosed with lymphoma in 2004. Using Appendix AA, the external dose for this case would be a little over 100 millirem. The internal dose a little over 76 rem. The medical dose from x-rays, approximately 84 millirem, for a total of over 76 rem.

Probability of Causation based on that dose estimate and the demographics on the previous slide will result in a Probability of Causation of 54 percent.

The second example is someone that worked at Hooker Electrochemical longer, over 30 years. Unknown job title, which Appendix AA does not try to distinguish different job titles. It puts everybody in the The demographics for this case operation. 1917, diagnosed male, born in prostate cancer in 1993. And the dose in this external, approximately 5-1/2 case,

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internal, 32 millirem, and medical dose, 25 millirem, for a total of a little over 5-1/2 rem. Probably of Causation with those demographics and that dose result in a Probability of Causation of 9.24 percent.

after the petition was qualified, prepared in accordance with 42 CFR 83. It was issued and sent to the Board on May 3rd of this year. And as you all know very well the SEC process is a two prong test. The first test is whether or not it's feasible to reconstruct the radiation dose for members of the class, and the second test is whether the likelihood of suffering radiation dose endangering the health of members of the class.

What we found in our Evaluation Report was, we are able to use source-term data to reconstruct with sufficient accuracy, they issue those for members of the Class. Because of that determination, any determination of a health detriment wasn't

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necessary for the SEC process.

And the last slide just reiterates that Evaluation Report concluded that we could reconstruct all the radiation dose at Hooker Electrochemical.

Any questions.

CHAIRMAN MELIUS: Thank you,
Dave. Board Members? Yes, Phil.

MEMBER SCHOFIELD: Okay, first off, I don't like that word likely. That leaves too much to interpretation. The other thing is, I want to know how much you know about the ventilation system there, how much you know about the size of the batches were, what kind of equipment they had, the number of hours per week these people worked, and I mean these are just basic questions you would need to be able to have answers to to do any kind of reconstruction.

MR. ALLEN: Yes, and we do have more detail than what I presented in this presentation. The drum dumping, the dustiest

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1	part of the operation, was done outdoors next
2	to the building on a concrete pad next to the
3	railroad spur. The surrogate data we've done
4	indoors without the aid of ventilation. The
5	digestion tank, the batch was 10 tons, 40 500-
6	pound drums. The tanks were wooden vats, 11-
7	foot diameter, about 11 feet high. I'm not
8	sure if that answered all your questions or
9	not. There is more detail that is included in
10	documentation and in the Appendix AA.
11	CHAIRMAN MELIUS: Other questions
12	for Dave? Yes, Jim.
13	MEMBER LOCKEY: Hooker
14	Electrochemical was not a scrap recycling,
15	right?
16	MR. ALLEN: Excuse me?
17	MEMBER LOCKEY: They didn't do
18	scrap recycling? Am I reading the slides
19	right?
20	MR. ALLEN: They did not
21	specifically do scrap recovery for uranium,
22	no. This operation you could, some would say

it's scrap recovery, it's more of a contaminated material concentrating. But the data we have for scrap recovery did not come from Hooker if that is the question.

CHAIRMAN MELIUS: As a follow-up maybe you could describe where that data did come from.

MR. ALLEN: It came from a publication that I cannot recall the authors of. It is all listed in TBD-6001, I believe it came from more than one site, but I'm afraid I don't recall the sites that it actually came from right now.

CHAIRMAN MELIUS: I'm just trying to understand the comparability of the sites. You mentioned for example that the TBD-6001 Appendix, the scrap portion of it, it deals with, also includes furnace operations. And so I guess I'm trying to understand, I think this is what Jim was getting at also, is what kind of operations are involved in - in developing that surrogate dose. I think that

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is what we are trying to judge and compare.

MR. ALLEN: The scrap recovery, that described in TBD-6001 in the was reference that it came from was what I listed the slide there, and it started with calcining to burn off any organics or oxidize any uranium that they could. And then it was digested in a tank using acid. The biggest difference there after that point was probably that the - in that case they were dissolving the uranium, then it went to another step where the uranium was precipitated, whereas in Electrochemical uranium Hooker was never dissolved. Ιt left in was a magnesium chloride matrix, and they tried to dissolve the matrix, both done in a liquid vat, it doesn't create a great deal of internal dose generally.

CHAIRMAN MELIUS: And my other -
I hate to go down -- be careful going down
this road, but can you speak a little bit more
about why you did not think that a source-term

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1	model was useful? And was there any attempt
2	to do a source-term model that would help us,
3	might help us - I should say that very
4	carefully - might help us to be more
5	comfortable with the surrogate model that you
6	used?
7	MR. ALLEN: Well, I certainly
8	never said that a model is not useful. I
9	think we could do one, and I think it would be
10	reasonably accurate. Any model does depend on
11	some assumptions. We explored a few different
12	models, one being NUREG 14000, and there are
13	various parameters that you have to
14	essentially pick a value for, depending on
15	what you have. And I think we could justify
16	those parameters. But in this case it seemed
17	like a closer fit than picking some
18	assumptions for a model.
19	CHAIRMAN MELIUS: Okay, thanks.
20	Phil then Henry.
21	MEMBER SCHOFIELD: Okay, on the

internal uptake and internal exposures, how

1	are you going to be able to limit these or
2	know what they were?
3	MR. ALLEN: I'm not sure I
4	understood the question. We are using
5	surrogate data from TBD-6001, which is
6	essentially airborne data, and we are assuming
7	they are inhaling that concentration of
8	uranium. Is that the question?
9	MEMBER SCHOFIELD: Yes, I wanted
10	to know on what basis you were coming up with
11	internal exposures.
12	CHAIRMAN MELIUS: Henry, then
13	David Lemen.
14	MEMBER ANDERSON: Is the
15	surrogate data from the same period of time,
16	from the early 40s, 43 to 48?
17	MR. ALLEN: It's from the 40s and
18	early 50s. The reference actually had data
19	for doing this operation without ventilation
20	and data for doing it with ventilation which
21	came after 52, give or take, I don't remember
22	the exact years. We did not use in TBD-6000

the data with ventilation; we used the earlier stuff. That is only without localized ventilation.

MEMBER ANDERSON: And what about the residual period?

MR. ALLEN: The residual period is modeled based on what the surrogate data we were using for the operational period.

CHAIRMAN MELIUS: Dick.

MEMBER LEMEN: Well, you answered one question that I had, and that was what what were the dates of Henry asked, surrogate data. But I still don't feel in the Petition Evaluation Report that you explained why you picked the scrap recovery process, and why you went the surrogate data also how do the way, and measurement techniques on the surrogate data -- how would they relate to what would have been taken if you could have found data in this plant? other words, how does the data relate from the surrogate data to what's really going on in

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this plant? Because there are a lot of different operations, it seems like, that by just taking one scrap recovery process you are going to miss a lot. I just don't think you explained, at least to my understanding.

MR. ALLEN: Okay, well for TBD-6001, the scrap recovery process, the -- as I the airborne, the mentioned intakes dominated by the furnace operation, very much dominated by the furnace operation. furnace operation involved placing the material, material. scrap uranium furnace, heating that to oxidize the uranium and eliminate organics and then unloading it. And it appeared that unloading was probably the highest airborne-causing operation. Some of the material, or some of the samples, are Some GA samples. of the samples breathings on the samples. And TBD-6001 puts it together into a value.

CHAIRMAN MELIUS: Henry.

MEMBER ANDERSON: Maybe I just

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1	can't find it, but where is the TBD-6001
2	document? I don't see it on the O: drive.
3	Maybe it's buried somewhere.
4	MR. ALLEN: It should be.
5	CHAIRMAN MELIUS: I was just
6	trying to look it up on the DCAS website, and
7	6000, 6001 are hard to find. They don't fit.
8	They don't get indexed well.
9	MEMBER ANDERSON: Okay, if you
10	could show me where it is.
11	CHAIRMAN MELIUS: If I find it.
12	I'm looking right now.
13	MEMBER ANDERSON: We are starting
14	to do that here, too.
15	CHAIRMAN MELIUS: Dr. Ziemer or
16	David Richardson, do you have either of you
17	have questions?
18	MEMBER ZIEMER: No questions at
19	the moment.
20	CHAIRMAN MELIUS: Okay, David.
21	MEMBER RICHARDSON: Yes, I was
22	wondering if you could talk a little bit about

the chemistry that goes on with calcining uranium in a furnace, and I assume you end up with some variety of uranium oxides, right?

MR. ALLEN: Yes, you would end up with a variety of uranium oxides as well as some uranium metal itself that might be left over, not completely oxidized.

MEMBER RICHARDSON: And so how does the nature of the chemical forms of the uranium that you are getting through the calcining process that is kind of the starting point for the TBD-6001 scrap recovery process differ from the chemical forms of uranium that would be encountered at the Hooker facility where they are not doing it?

MR. ALLEN: Should be very The reduction pot linings is similar. the contaminated mag fluoride was from. in that, process uranium is converted, essentially uranium tetrachloride is reduced The contamination occurs to uranium metal. when the uranium metal is in a liquid form, a

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1 molten form essentially contaminates the 2 So you should have small amounts of lining. 3 uranium metal as well as plenty of oxides 4 heated around 1,200 degrees. 5 CHAIRMAN MELIUS: Henry, do you 6 still have questions? 7 MEMBER ANDERSON: Oh, no. CHAIRMAN MELIUS: 8 Mark. I found like 10 MEMBER ANDERSON: 9 10 references to the Work Group. CHAIRMAN MELIUS: Yes, go ahead. 11 12 MEMBER GRIFFON: Yes, Ι 13 wanted to follow up on the TBD-6001 Appendix AA. And actually this goes back to the 14 15 appropriateness of the surrogate model, 16 Page four in there, it talks about quess. basically the basis for using the surrogate 17 18 model. And it says the dumping operation was 19 assumed to be similar to the furnace operated Since that provides the highest air 20 trays. concentration for scrap recovery. To me that 21

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sort of is not the best rationale for using

1	that as a surrogate model. You were saying
2	that we assumed it's the same because it gave
3	the highest levels. I don't dispute it
4	goes on to show how high those levels are, and
5	it's a very dusty operation. I don't dispute
6	that. But again are we just bounding with a
7	high value or are they really similar enough
8	to be used as surrogate? That is sort of my
9	question there. And I don't even expect an
10	answer on that one.
11	The one question I did have was,
12	where was the slag material from the 43 to 48
13	time period? What companies were funneling
14	into Hooker at that time?
15	MR. ALLEN: Into Hooker?
16	MEMBER GRIFFON: Yes.
17	MR. ALLEN: All the material for
18	Hooker came from Electro Met. It was
19	reduction pot liners.
20	MEMBER GRIFFON: From Electro
21	Met, so it was all in-house. They weren't
22	getting anything from other plants at that

1	point?
2	MR. ALLEN: All the information
3	we have is it all came from Electro Met by
4	rail.
5	MEMBER GRIFFON: I think this may
6	end up and I think we might need further
7	reviewing and the TBD-6001 Work Group actually
8	might be a good place to do it.
9	CHAIRMAN MELIUS: We may be
10	getting there. But first I think if there are
11	no further questions right now, why don't we
12	hear from the petitioners.
13	Oh, I'm sorry, Jim.
14	MEMBER LOCKEY: Just so I'm
15	clear, in the original proposed Class, she
16	said furnace room, or the petitioner said
17	furnace room, but there is no furnace room.
18	MR. ALLEN: No furnace room where
19	they did the magnesium fluoride concentrating,
20	no.
21	MEMBER LOCKEY: So there is no
22	calcine?

1 MR. ALLEN: No. 2 MEMBER LOCKEY: Thank you. 3 CHAIRMAN MELIUS: Okay, 4 petitioners? Who wants to start for the 5 petitioners. And if you could identify 6 yourself and then --7 MS. GIRARDO: Can you hear me all right? 8 CHAIRMAN MELIUS: 9 Yes. 10 MS. GIRARDO: I wish I could have heard him better. I have a hearing problem 11 and I thought this would be better as far as 12 13 the acoustics go, but apparently it wasn't. And anybody with a mustache and a beard is 14 15 dangerous to me. It's very hard to -- you 16 need to look at the lips and the sound in order to put the two together. So I feel like 17 I was kind of outgunned with that. 18 19 I wanted to thank you for coming 20 into Niagara Falls. I hope you have a good time here. I am going to be 75 years old in 21

June, and of course I was born here in the

city, and my father worked for Hooker Chemical, and of course, it was Oldbury at that time, which was a company from England. And so we have a history with this company. And I want to say that we disagree with the evaluation of NIOSH based on use of surrogate data; I don't think that is fair. And I would like additional time to prepare a written statement, because we just got this over-50page evaluation, I think it was Thursday or Friday, and just had the weekend really to prepare. So I would prefer to have some more time to write a written statement. And Laurie Breyer is willing to see that the Advisory Board gets that.

And then I request that Sanford Cohen & Associates check out this evaluation to see what they think of it. And I thank you for your time. And again I say I hope you enjoy yourself in Niagara Falls. Thank you.

CHAIRMAN MELIUS: Thank you.

Anybody else? Okay. Thank you.

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We realize that there wasn't significant amount of time, but one of the to be able to gather reasons we came was information on what -- on the facility, and in order to help with our evaluation of it. you will have time to get additional information in, so it will be considered.

Any Board members have questions or further comments? We have it, it may be sort of arcane within the Board, we have this document, it's called the TBD-6001 that NIOSH has produced as a guide for a number of different facilities, the dose reconstruction. And we have a Work Group that is set up of Board members, a smaller group that concentrate on that -- that work on addressing issues related to TBD-6001. And I would think that this site we would refer to that Work Group for follow-up as they are beginning and doing their review.

Josie?

MEMBER BEACH: Well, I also

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wondered if it didn't fit in the Surrogate Work Group just because of those issues.

CHAIRMAN MELIUS: Well, I think we are -- we can decide, I don't know what the work load is for the 6001 Group relative to the Surrogate Group.

MEMBER ANDERSON: We don't know yet.

CHAIRMAN MELIUS: Yes. Yes. And do that, and so maybe let's start with the 6001 group, and then we can decide. The 6001 Group I think by assumption, both the 6000 and the 6001 are we'll be dealing with surrogate data issues, and that is part and parcel of the application of both of those documents. And so I think we would -- I think the place to start would be with 6001. Т think if we run into an issue of just the volume, that's why we split 6000 and 6001, that we would consider that. Also I don't know, John Mauro, have you reviewed Appendix What is the status of your review? AA yet?

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DR. MAURO: We have not. We have
reviewed TBD-6001 and have a number of very
significant comments. So the model I have in
my head is that first, the rock you are trying
to stand on is 6001, the basis for which there
is all this data that has been collected,
sorted and binned. And the degree to which
that was done well and captured the universe
of data that is associated with uranium
processing is sort of the first step in the
process. If it doesn't survive that process
it almost is, well, you can't really go to the
next step. If you don't have a sound 6001,
then you go ahead and use it, then it's almost
like you are not standing on a rock. So I
agree, the first thing is, while you are doing
this you are working 6001 at the same time.
Vou can't get they are hooked together

CHAIRMAN MELIUS: Yes.

MEMBER BEACH: And Jim, the ER also talked about AA, C and D, so are those three separate appendices to the 6001? I

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1	don't know.
2	CHAIRMAN MELIUS: You are going
3	beyond my
4	MEMBER ANDERSON: It is in their
5	references.
6	CHAIRMAN MELIUS: Yes, right, I'm
7	not familiar with that part of it. I don't
8	know if someone from NIOSH can help. Dave.
9	MR. ALLEN: There are several
10	Appendices to 6001 as well as 6000. I believe
11	this Evaluation Report referenced the Electro
12	Met, but I'm not positive. I think that is D.
13	MEMBER BEACH: D?
14	MR. ALLEN: Appendix D.
15	MEMBER BEACH: It wasn't in your
16	listing.
17	MR. ALLEN: I might have that
18	wrong. But that would be the only other
19	appendices to TBD-6001 that would be
20	referencing other sites. Is that your
21	question, Josie?
22	MEMBER BEACH: That's fine, thank

you.

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CHAIRMAN MELIUS: So yes, John, again, do you want to bring us up to date on where you are with 6001?

DR. MAURO: have reviewed We Electro Met, delivered our report. I think it showed up recently. So in effect, I think there were four or five appendices to TBD-So we have a situation. We certainly 6001. have to engage TBD-6001 on its own merits, and in the process of doing that -- now, I believe Electro Met doesn't depend that heavily on 6001. It stands more on its own data; I recall it has its own air-sampling data for example. there significant So are some differences in terms of the dependencies that Electro Met uses, which I believe is CC or C, and it sounds like certainly Hooker depended very heavily on TBD-6001. So they all sort of come together.

CHAIRMAN MELIUS: We are really just thinking that in terms of this SEC

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Evaluation, just wanted to make sure that SC&A had the relevant appendices under review or in process some way so that we don't get to the point where well, we still need to do one more. So if you can take a quick look at this SEC Evaluation and what is referenced there, just to make sure that we have got that covered, or talk to Dave who may be able to assist in that.

So does everyone agree that we will refer this to the 6001 Work Group? And I'll just add for the petitioner and other people who are interested, this Work Group will be sort of doing the initial review, they will assign SC&A for any additional work that is needed. There will also be a communication to the petitioner and other interested parties about when they are meeting, what's under consideration, any timing and any information that petitioner or other people can provide to the Work Group will be useful and considered and there will be an opportunity for you to

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comment and know what's going on. And I understand, I know you may not -- may have some trouble hearing some of what I'm saying, with the beard and mustache. I think Laurie can help also with that. But we appreciate you coming here today and taking the time and be assured that there will be follow-up on that, so.

So what we will do is, we will have a brief presentation from the Linde Ceramics Work Group, follow up and do that. Then we'll take a short break. And then we'll start the public comment period. And we have agreed to start the public comment period focusing on Linde, so that there is some continuity in terms of follow-up. Okay.

LINDE CERAMICS (TONAWANDA NY) WORK GROUP UPDATE

MEMBER ROESSLER: While she is getting ready, we'll check my voice level. I don't have a mustache, but I want to make sure you can hear me. And also since we are

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disclosing ages, I turned 75 this past April, so we are in the same group here.

Again, while she is bringing that up I will mention that my purpose here today is to bring the Board up to date on the Linde Work Group activities, and also to, since we probably have interested people in the audience, to summarize it in a short period of time to let you know what the Work Group has been doing.

And I will also tell you that we will not be taking a vote today. The Work Group is not prepared to present to the Board information to take a vote. We still have some issues that need to be resolved.

CHAIRMAN MELIUS: While you are working on the computer, I actually have one request for Ted or somebody to take back is, can we get the TBD-6000/6001 put in the list of technical documents along with all the appendices so that they are easier to find. Separate folders both on the O: drive as well

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1	as on the DCAS part of the website. Because
2	it is extremely frustrating to find and then
3	to try to figure out whether SC&A has issued a
4	review is even harder. So relative to those,
5	the other parts, all the other Sites work
6	well. Just that one, because it's not a site.
7	
8	MR. KATZ: I know, I have similar
9	difficulty with the website. So we will
10	follow up.
11	CHAIRMAN MELIUS: Okay.
12	MEMBER ZIEMER: This is Ziemer.
12 13	MEMBER ZIEMER: This is Ziemer. If you do the appendices individually as sites
13	If you do the appendices individually as sites
13 14	If you do the appendices individually as sites that will work better, probably.
13 14 15	If you do the appendices individually as sites that will work better, probably. CHAIRMAN MELIUS: But then you
13 14 15 16	If you do the appendices individually as sites that will work better, probably. CHAIRMAN MELIUS: But then you have trouble getting back to the original; at
13 14 15 16 17	If you do the appendices individually as sites that will work better, probably. CHAIRMAN MELIUS: But then you have trouble getting back to the original; at least that's been my experience.
13 14 15 16 17 18	If you do the appendices individually as sites that will work better, probably. CHAIRMAN MELIUS: But then you have trouble getting back to the original; at least that's been my experience. MEMBER ZIEMER: Getting back to
13 14 15 16 17 18 19	If you do the appendices individually as sites that will work better, probably. CHAIRMAN MELIUS: But then you have trouble getting back to the original; at least that's been my experience. MEMBER ZIEMER: Getting back to the TBDs?

1	but then you sort of have to remember which is
2	in which.
3	CHAIRMAN MELIUS: Right, so
4	that's why there ought to be a better way.
5	MEMBER ROESSLER: Okay, thanks to
6	Dr. Poston. We are ready.
7	First of all, I'd like to point
8	out the Work Group members on the Linde
9	project in addition to myself are Josie Beach,
10	Mike Gibson and Jim Lockey.
11	And the team working with us from
12	NIOSH, and it's now called DCAS, Chris
13	Crawford and Jim Neton. Now neither Chris nor
14	Jim are here today, but Dr. Sam Glover is here
15	I think somewhere, in case we have any
16	questions later on.
17	The SC&A team is Steve Ostrow,
18	John Mauro, and Bill Thurber. And Dr. Ostrow
19	and Dr. Mauro are here.
20	Just a little background first
21	just to bring everybody kind of up to date on
22	it. Linde Ceramics Plant, a division of Linde

Air Products Corporation, was located in Tonawanda, New York. And I saw that area as we came up here in a cab. Linde Ceramics originally handled uranium products, used as dyes for ceramic tableware. And when I think of this I think of my kitchen cupboard where I have a large collection of Fiestaware. I'm assuming maybe this is the type of tableware that was made there.

Then because of the capability they had in 1942, Linde Ceramics contracted with the Manhattan Engineering District, and we call them MED, to process uranium ores to produce uranium oxide, also called yellowcake, and uranium tetrafluoride, also called green salt.

Okay, again a little background: in the 1930s, Building 14 was known as the Tonawanda Laboratory, and that is included in this study, owned by Union Carbide. They produced uranium, U-308 as a coloring agent for ceramic glazes. We already mentioned

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that. And at least 80 tons of U-308 were produced before this MED period began in 1943. The plant had production years but the years between, or the time between July 1st, 1949, and July 7th, 1954, there was no production. It was called the decontamination and decommissioning period.

Now there are three petitions of interest here, actually we are only going to concentrate on one, but just as a listing, and I'm not going to read through everything here, there is SEC-00044. This has been granted to the workers who worked at the plant between October 1st, 1942, through October 31st, 1947.

Another petition, 00154, has qualified for evaluation, but NIOSH has not issued an Evaluation Report yet for this period. So we are concentrating right now under -- on SEC-00107, for the period January 1st, 1954, through July 31st, 2006, called the renovation and residual periods.

This petition was received on

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March $3^{\rm rd}$, 2008. It qualified for evaluation on July $2^{\rm nd}$, 2008. I have already kind of alluded to this, but our Work Group has focused on the petition that has qualified for evaluation, and again, the period is January $1^{\rm st}$, 1954 through July $31^{\rm st}$, 2006.

And I'm going to tell you a little bit about our Work Group meetings, because there is not going to be time here to tell you about everything we did at them. But we did start in March, on March 26th, 2007, with our first meeting. The first set of meetings we had was to evaluate the Site Profile. And the way this works is that NIOSH comes to the meetings and presents the Site Profile. SC&A has had an opportunity to critique it. So we spent four meetings going over the critique. NIOSH made changes and we agreed on June 23rd, 2008, that the Site Profile Review was That doesn't mean that it maybe completed. wouldn't change in the future. But at that point it was completed.

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On July 8th, I think it was, as I mentioned before, the SEC petition qualified. And in the August teleconference of the Board our Work Group was assigned the task of going ahead now to evaluate petition. So we promptly met on September 2nd, 2009, to do this, and again along with NIOSH, talking about various issues SC&A, throughout these Work Group meetings which were in September, December, and January. brought out a lot of issues and discussed there seemed to them. Because be technical information that NIOSH and SC&A had to resolve, they had a technical call on February 23rd, 2010. The Work Group listened in but we didn't participate. Then on April 16th, 2010, we had our most recent meeting.

Now just to summarize a little bit about what we talked about, the potential radiation exposures there with regard to internal -- radon was one consideration present because of the residual contamination

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of surfaces by ores. Remember, this is not the production period but residual contamination remained.

Then during the renovation work in Building 30 in the 1960s, it's possible that there were airborne contaminants. And then also the airborne radioactive contaminants were evaluated for the residual -- the whole residual period.

With regard to external sources, gamma or photon or beta exposure from the residual uranium that contaminated the surfaces present. was Neutrons are not considered as a source of exposure to Linde personnel.

And just as a point of information, I want to remind people that even though some radiation exposure may have occurred during this time, and this is what is being evaluated, I want to remind you that dosed workers during this period was much less than during the production period.

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So with our first Work Group meeting in September where we were assigned to evaluate the petition, it was then that NIOSH presented their findings. And this is the statement from NIOSH. NIOSH found that the available monitoring records, process descriptions and source-term data, adequate to complete dose reconstructions with sufficient accuracy and so on. So this is where we started.

with started then the We discussions between NIOSH and SC&A to look at whether this was valid. And again, I can't go over all of this, but basically what happened at these Work Group meetings is that -- and I'm not talking about what the Work Group has decided, because we have not yet as a group had any vote. But SC&A accepted the NIOSH proposal for bounding the dose during renovation period and the balance of residual period. SC&A also agreed to accept the NIOSH treatment of radon, in other words,

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bounding radon exposures in the Linde buildings for this period.

I just thought I should put a note here that through all these discussions, though, there may be some changes. The dose estimates are probably going to be increased, and this may lead to Site Profile revisions.

Now, the reason that we are not yet able to vote today is that there are still some open issues. This came up primarily at our last Work Group meeting, and these open issues, and those of you who participated in the interviews this morning with SC&A know that this is what we are concentrating on, the open issues involve potential utility tunnel exposures. And this is a little bit hard to read, but I didn't want to leave out any detail on here, because this is what we are dealing with now.

The question is, can the tunnel dose be bound at or below 2.3 MAC, which in our jargon is Maximum Allowable Concentration,

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for the period through 1970. If this is accepted, then NIOSH must explain why they feel that these doses wouldn't exceed this particular level, in other words that they can really bound with this number. So they must show that the doses can be bound and the current open issues are looking at ventilation, composition of the tunnel walls, radon from the soil, et cetera.

Some other questions came up too in this discussion about placement and depth of the injection wells. I have heard that some of the workers discussed that this morning with SC&A. The hydrology of the area, the depth of the tunnels, the location of the sump pump discharges, and some other issues. And there is much information apparently available on this. So NIOSH, SC&A and others are looking at this.

So there is where we stand now; we have some open issues to resolve.

So what is our plan forward? The

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1	plan was to have more worker interviews since
2	we are in the area today. SC&A did conduct
3	those this morning. NIOSH is going to produce
4	a more detailed tunnel report. SC&A is going
5	to review this report. SC&A also has
6	available other data, is reviewing other
7	information about the tunnels. Once this is
8	all put together we hope it will be fairly
9	soon the Work Group will meet again. And
10	then our goal is to make a presentation to the
11	Board for a vote for the Board meeting in
12	Idaho Falls in August.
13	That is our plan, and we will see
14	how that goes. So I think at this point, I
15	can entertain questions from the Board and any
16	questions from the public or comments would
17	come later at the public session.
18	CHAIRMAN MELIUS: Anybody from
19	the Board have any questions at this point?
20	(No response.)
21	Thank you for a very concise and

That was excellent. And it's a

good update.

little after 4:15. We will take a break and we will try to start right at 4:30 with our public commentary. So the Board members and everyone else can stretch.

(Whereupon, the above-entitled matter went off the record at 4:17 p.m. and resumed at 4:34 p.m.)

CHAIRMAN MELIUS: We are going to focus on Linde. As I said, we had talked to the petitioners and that was their preference, agreed, since that -- I think, and we actually, the first one, we have a written statement from one of the petitioners that couldn't be here, so Ted was going to read that into the record, so we have it on the record. And then we will start the public comment.

MR. KATZ: Right. Thank you. Before -- also, there's a little spiel I have to give before every public comment session, generally about the redaction policy at NIOSH, which is, as many of you may know, all of the

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Board meetings are transcribed, including the sessions, public comment verbatim transcription, everything said on the so record is captured there. So, as a member of the public, when you speak, everything you say will be captured in the public record, including your name, including any personal information you might give, but we do redact from your statements any information you give about third-parties, other persons, to protect their privacy. So their names and identifying information about those individuals would be redacted from our And the full redaction policy transcripts. should be out on the table, and it's also on the website for your reference with the agenda for the meeting. So, just to let you know that up front.

Now, one of the petitioners, Linda

Lux sent in a letter just in advance of the

meeting saying she couldn't be here. That's

one of the Linde petitioners, but that she

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would like her statement to be read into the record with her being identified, so I have so identified her. And let me just read you her letter, and then we'll hear from the other Linde persons that are actually here, or might be on the phone, as well. So, from Linda Lux, May 17th.

"To Advisory Board on Radiation and Worker Health Members. Because it is not possible to attend this meeting, I would like to voice my concern in letter form as to why the Linde site SEC petition should be approved.

I have, for the last eight years, given NIOSH and DOL every bit of information available to me regarding the claim for my deceased father. I have provided unemployment records, multiple medical records, and two letters from Dow, Union Carbide's purchasers, stating my father's dosimeter records have been destroyed. Unfortunately, much of this very important information has gone

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unrecognized, and, obviously, unread by NIOSH.

My father worked in the computer department at Linde, so this puts him in the category of office worker. Office workers in the dose reconstruction at Linde receive an extremely low dose, despite the fact that I in the worker outreach had stated before meeting on page 120 of 126, it states that clerical "eight office and workers all developed cancer within a short time from one another."

In the dose amount given to office workers, it would be impossible to qualify for In my father's medical records, compensation. he stated to his doctor before this EEOICPA started, that he worked program ever extremely dusty conditions for a two-year time period. My father passed away from cancer in 1994 at 59 years of age, so I cannot ask him what time period it was. I believe he was referring to the 1960s remodeling jobs that going at the Linde site in the were on

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building he was working in. I do remember him coming home from work in that time period with a lot of dirty dust on his clothes, and an odor on him.

After receiving in 2006 the first dose reconstruction, I asked the Department of Labor to read the medical records regarding a second cancer and some lung brushings that were done and listed in the medical records. DOL called me back to say they were going to send this claim back to NIOSH to redo the dose, and add a second cancer, but I would need to get more information for the lung brushings.

I could not retain any further records from the doctor that did the lung testing just days before my father passed away because the doctor had retired and the records were only kept for 10 years after a patient's death until they were destroyed. If only NIOSH had read the records when they were submitted in 2002 and told me I needed more

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details, I could have received those records because it would have been in the 10-year time frame.

When I received the redo of dose reconstruction just two weeks ago, not only did my father's dose amount not go up with a second cancer added but it went down, I was told the reason was NIOSH has way down. the dose adjusted amounts to be more realistic. What I expected from NIOSH, at the least, to receive dose very was reconstruction that included both cancers and considered radiation as the only risk factor that matched the cancer my father had, but what I received was what looks like, to me -is a manipulated application of numbers to control the outcome.

If a true consideration of my father's cancer and other petitioner's cancer was caused by a work location, I would think it would be important to consider statements the workers have made at both the worker's

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outreach meetings, and in the medical records, as well as factors that go along with the listed cancers.

I don't see how it can be said that "best available science" is used when comments from workers are not considered. The Linde site has unique features that also must be considered, such as toxic chemicals poured into wells that overflowed with rain, toxic chemicals that were poured into the drainage system that also overflowed, toxic chemicals buried in the ground and poured into nearby water streams, construction and remodeling done to buildings during the 1960s and 1970s that were embedded with toxins while Linde workers stayed working in the buildings with no protection.

To say a person didn't have enough exposure to cause a certain cancer is an untrue statement. Any dose amount is too much and is enough to cause cancer in some people.

Every person's body chemistry is unique and

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1	can handle or fight off different amounts of
2	toxic substances or radiation before a cancer
3	sets in. These workers were unaware of the
4	radiation all around them, including in the
5	dirt and water outside, so they would not have
6	acted in a cautious way. To not consider
7	these facts and not include unique
8	circumstances to each worker is not a fair or
9	true dose reconstruction for many of the Linde
10	workers. It would then not compensate many of
11	the workers who should be compensated, and
12	they are who this program was created for.
13	Sincerely, Linda Lux."
14	CHAIRMAN MELIUS: Okay. We now
15	want to hear from the Linde petitioners. And,
16	Antoinette, are I'm not sure what order to
17	go in here. Yes, from there. We need to
18	activate that mic, I think.
19	MR. KATZ: Try again, Antoinette.
20	MS. BONSIGNORE: Thank you. The

first issue I'd like to raise is regarding the

petition that just qualified for Linde SEC-

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1	00154. I received a letter stating that DCAS
2	will not be able to meet its 180-day deadline
3	for two reasons, first being that interviews
4	were conducted today that may affect that
5	petition. And, secondly, that there were
6	documents that have been uncovered at the
7	National Archives that deal with the Linde
8	tunnels. And I don't know if Mr. Rutherford
9	is here, but if we could get some
10	clarification as to what those documents are,
11	and when they might be made available to the
12	petitioners?
13	CHAIRMAN MELIUS: Stu, can you
14	address that?
15	MR. HINNEFELD: Well, LaVon is not
16	here, but and I'm not conversant about what
17	those documents are exactly. We can provide,
18	certainly, that information to you during the
19	week. LaVon will be here tomorrow.
20	MS. BONSIGNORE: Okay. Thank you.
21	And the second issue deals again with the
22	180-day deadline for the release of Evaluation

Reports, and, specifically, with respect to the Linde petition.

My question to the Board is why is DCAS allowed to continually revise ERs as more information becomes available to them? And, many times, looking for additional information that will justify the denial or recommendation of a denial for the SEC petition. What permits DCAS to go beyond the 180-day deadline that is specifically prescribed in the regulations?

CHAIRMAN MELIUS: I don't think we can speak to that legally. My understanding is that it's not a binding time period, and I think there's also, I think in general in this there is the policy program, as new information becomes available, and it favors the claimant, that it is then incorporated into dose reconstructions. So there's been a general policy as new information becomes available to utilize that. I think as Board, we have concerns about the timeliness

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of response to these new information and sort of what the limit is. And I think NIOSH has concerns about that, also, and are trying to address it in order to make this more timely, so it's not an endless process. However, if information or a new issue comes up, such as the tunnels or something, then I think it sort of behooves us to try to allow time for the gathering of additional information.

MS. BONSIGNORE: However, if gathering of additional information over an extended period of time works to the detriment of the petitioners, if, in fact, the original issued was somehow incomplete, ER that was or deficient inaccurate, in some shouldn't that be the Evaluation Report that actually considers? if the Board And additional research is conducted thereafter, that would benefit petitioners or individual claimants, then that would be fine, but why petitioners penalized are the when been, original ERperhaps, would have

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1 considered by the Board to not be 2 satisfactory? 3 CHAIRMAN MELIUS: Again, the 180 days isn't binding, so it's a question of 4 5 judgment. I mean, I think we'll take that as 6 a comment. I don't think we can sort of fully 7 address it. MS. BONSIGNORE: Okay. All right. 8 Thank you. 9 10 CHAIRMAN MELIUS: Thank you. Okay. [identifying information redacted]. Is 11 12 [identifying information redacted] 13 speaking to Linde? I don't always have what -- okay. Mary Girardo, again. We'll do again, 14 15 and see what this -- those people may think 16 that we're only talking about Linde right now. Sandy Rykiel. Okay. If you'd like to step 17 18 to the mic, either this mic here, or you can 19 use the podium. And Ι apologize if 20 mispronounce anybody's name. With a name like Melius, I'm used to --21

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RYKIEL:

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CHAIRMAN MELIUS: You can go to the podium, whatever is better.

Thank you. MS. RYKIEL: I'm here to speak about my father, William Donovan. was an employee at Linde Division in Niagara He worked there for 36 years and 29 Falls. He was a chemical operator from 1942 to 1957 the cobalt plant. at He was an electrician from 1957 to 1961 in the cobalt plant. He was an operator of the furnaces, Operator A and D, and a foreman, and from 1961 to 1965 he was an electrician. From 1965 to 1970 in the Linde Division, mining the metals. He was a foreman, master mechanic from 1970 to 1978.

When I originally filed this claim, I had heard about it from Roswell Park.

My father did have prostate cancer. From my understanding, prostate cancer is one of your organs on the list of categories, but yet you use the bladder as a surrogate organ, as an

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1	internal surrogate organ. How can you use a
2	surrogate organ for the prostate? I don't
3	agree with this surrogate organ. I don't
4	can you explain that to me?
5	CHAIRMAN MELIUS: Yes. Stu, do
6	you want to
7	MS. RYKIEL: I understand what a
8	surrogate organ is; I just don't agree with
9	the way you're using it.
10	CHAIRMAN MELIUS: No, no. And I
11	think there's an explanation.
12	MR. HINNEFELD: Yes. I can do it
13	now, or we can do it later on.
14	CHAIRMAN MELIUS: Well, why don't
15	you do it now. It's a general question.
16	MR. HINNEFELD: Okay.
17	CHAIRMAN MELIUS: If we can answer
18	a question quickly, and it does not involve
19	personal information
20	MS. RYKIEL: Yes, that's fine.
21	CHAIRMAN MELIUS: we'll try to
22	do it. If not, we can also do some of these

in follow-up. I'll also indicate that all these comments are being recorded, and that actually the Board -- we actually have a Work Group that's looking to -- making sure that we do the follow-up for the comments and that they're collected and dealt with in terms of the information being used in sort of our future efforts on SEC evaluations and dose reconstruction.

MR. HINNEFELD: Okay. In this context, there are -- a dose reconstruction relies on data that allows you to convert certain measured quantities, like the quantity measured by a film badge to the dose received by some internal organ, whatever organ you're interested in. And the International Radiological Protection Commission on published a number of correction factors, in other words, ways to correct something like a dosimeter badge to specific organ doses. did not publish one for the prostate. an external dose, the method that's chosen for

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dose reconstruction is to choose another organ in close proximity to the target organ you're interested in. So the bladder was chosen as the organ in close proximity to the prostate the one where the dose would as essentially, the same as it would be for the prostate from this external source. I believe we use bladder for the surrogate for an external.

On occasion, you'll have the same issue with an internal dose where a particular organ that doesn't concentrate the radioactive material that's being ingested or inhaled does not really receive any particular dose, except from the blood that circulates through that So its dose would be essentially the same any other organ that doesn't as concentrate the radioactive material, but just it receives the dose from the blood circulating through that organ. So you would use, in that case, if your dose model doesn't include the exact target organ you're

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interested in, you would use some other organ of that same kind, that didn't concentrate the material, but just received the dose from the circulating blood. So that would be -- those were the two uses where an organ would referred to as a surrogate organ.

MS. RYKIEL: Okay. Well, father 44.78% for they rated my at They pay out at 50 percent. prostate. worked there 36 years and 29 days. He inhaled it; he was exposed to it; it was on his body. How can -- I don't know how you're coming up with these formalities. He had prostate cancer, he had skin lesions removed, he had a right breast mass removed, he was anorexic, he had malaise, he constantly short was breath, five heart attacks with open heart surgery with two aneurism repairs. TIAs, he had CVAs, he had congestive heart and his final thing that took him failure, down was respiratory arrest from pneumonitis, which is -- he also had kidney problems,

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1	decreased kidneys, profuse sweating, increased
2	blood pressure, diabetes and stroke. He was
3	exposed to mercury, plutonium, cobalt, and
4	uranium, as well as radiation and asbestos.
5	Thirty-six years, and I'm getting denied.
6	CHAIRMAN MELIUS: I mean, I don't
7	think we can speak to the actual dose
8	reconstruction. I just would say that the way
9	that the calculation is done to a great extent
10	is required what's required by law, and is
11	based on other studies that have been done of
12	cancer from radiation exposure. And those
13	other factors many of those other factors
14	that you mentioned are not things that are
15	taken into account because they're separate
16	from the other illnesses are separate from
17	the cancer and the radiation.
18	MS. RYKIEL: Okay. Then they
19	would fall under Part E then. Right?
20	CHAIRMAN MELIUS: Part E, and
21	that's
22	MS. RYKIEL: Okay. So when I

originally started this, and I -- this has been a long, ongoing, very tedious operation When we first started this, we reported here. everything that my father had, heart problems, and the whole gamut that I just read off to you. And now when I just spoke to Department of Labor, they have no record of They have no record of my father's this. medical records. You just told me you couldn't find his film badge, then you're telling me that there's not enough evidence. You're talking 1940s. This is 2010.

CHAIRMAN MELIUS: Yes. I mean, we can't speak to --

MS. RYKIEL: How can we -- how do you expect us to find this information? And I was told that we are the ones that have to provide you with the information. The family has to be the one that provides you with the information. How can we possibly go back to the 1940s and get this information when they're only carrying medical records for 10

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1	years? It's impossible. You can't Union
2	Carbide, Linde can't even find my father's
3	pre-employment, while he was employed, or
4	post-employment chest x-ray. If they can't
5	find it, how can I find it?
6	CHAIRMAN MELIUS: No, I think we
7	recognize that's a problem, but, again, the
8	Department of Labor is the one that has to
9	determine what information they will accept
10	for proof of illness. And it is difficult
11	because it is such a time period, but that's
12	not something we can directly address.
13	MS. RYKIEL: All due respect,
14	though, it sounds like you're passing the
15	buck.
16	CHAIRMAN MELIUS: Well, yes,
17	that's true, because the buck
18	MS. RYKIEL: So you are passing
19	the buck.
20	CHAIRMAN MELIUS: Yes, the buck is
21	not here, the buck's with the Department of

1	MS. RYKIEL: Okay. So then why
2	are we even why are we meeting here then?
3	CHAIRMAN MELIUS: Because we're
4	meeting to get
5	MS. RYKIEL: Are you our
6	advocates?
7	CHAIRMAN MELIUS: Excuse me, let
8	me finish.
9	MS. RYKIEL: I'm sorry.
10	CHAIRMAN MELIUS: Okay. We're
11	meeting to gather information and to listen to
12	concerns. There are certain concerns that can
13	be addressed through this program. There are
14	other concerns that have to be addressed
15	through the Department of Labor.
16	MS. RYKIEL: And what
17	CHAIRMAN MELIUS: We do not advise
18	the Department of Labor; we advise NIOSH.
19	MS. RYKIEL: You are Department of
20	Energy?
21	CHAIRMAN MELIUS: No.
22	MS. RYKIEL: What are you?

1	CHAIRMAN MELIUS: We're part of
2	we advise the Secretary of Health and Human
3	Services, which is of which NIOSH is the
4	Agency.
5	MS. RYKIEL: That does the dose
6	recalculation.
7	CHAIRMAN MELIUS: Correct.
8	MS. RYKIEL: Okay. So, we're back
9	to the beginning again. They did the dose
10	recalculation of 44.78%. When you spit all
11	this information in, did they put in all of
12	this information and come out with this
13	outrageous number? I mean, 5.22 percent of 36
14	years and 29 days? This just doesn't make
15	sense, you guys.
16	CHAIRMAN MELIUS: Well, yes
17	MS. RYKIEL: There's something
18	wrong.
19	CHAIRMAN MELIUS: All I can say
20	is, we have a program that we don't we
21	can't look at individual cases. We do have a
22	program that reviews a sample of the cases to

1	make sure that they are done correctly and
2	makes corrections to that process.
3	MS. RYKIEL: Yes, but is that
4	sample that you're using the ones that you've
5	paid out to?
6	CHAIRMAN MELIUS: No. In fact, we
7	concentrate on those that are closest to 50
8	percent, but below 50 percent. We try to get
9	the ones that are the most
10	MS. RYKIEL: So then my father
11	should be in there?
12	CHAIRMAN MELIUS: What?
13	MS. RYKIEL: Then my father should
14	be in there, with the ones that you're looking
15	at.
16	CHAIRMAN MELIUS: I can't address
17	be among the sample that would be
18	evaluated, yes, but I can't
19	MS. RYKIEL: Sure. I understand
20	that.
21	CHAIRMAN MELIUS: Yes.
22	MS. RYKIEL: Okay. All right. So

now I've been directed to reapply under everything under Part E, including the prostate cancer.

CHAIRMAN MELIUS: Correct. And under Part E, the Department of Labor can take into account other factors, including the chemical exposures, for example, that your father may have had that could be related to the development of the cancer. Under this program, the Part B program under cancer, we address the radiation only able to are exposures.

MS. RYKIEL: Okay. But I --

CHAIRMAN MELIUS: So you have a facility like Linde or something where there were many other exposures that may be involved in cancer or other diseases, that's something that's taken care of under Part E.

MS. RYKIEL: Okay. And then my final thing I just wanted to say is I think it should be taken into consideration that thyroid disease, MS, and prostate cancer,

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1	we're one of the highest areas in North
2	America, we are the highest rated areas. And
3	it's probably because more than likely, I
4	should use your words not your words, but
5	the assumption, because you hear the word
6	assumption all the time, with the assumption
7	that these plants are the ones that did it to
8	all these men.
9	CHAIRMAN MELIUS: Yes.
10	MS. RYKIEL: So they should be
11	paying for the prostate.
12	CHAIRMAN MELIUS: Okay. Could
13	very well be, but I can't you know, the
14	Department of Labor is going to have to make
15	that determination under their guidelines.
16	MS. RYKIEL: Okay. Thank you very
17	much for hearing me.
18	CHAIRMAN MELIUS: Well, thank you.
19	Eleanor Tornabene, I believe. I may again,
20	I apologize for pronunciation if I was wrong.
21	MS. TORNABENE: Hello, I'm Eleanor
22	Tornabene. I live on Grand Island now. I'm

here on behalf of my husband, Sam Tornabene. He worked at Linde in Tonawanda from 1962 until his death in 1993. He died of lymphoma. I guess the -- from early on when he started at Linde, he carried a radiation detector. At that point, we should have realized there was really something wrong with this work area, but I don't know what happened to that detector, or who kept track of the information that they garnered from it.

I'm sure that his exposure was on a constant basis because he worked in the factory. He started out in janitorial, then went to maintenance, worked in several areas in the factory, and he finished his career as a top grade welder.

I think the bottom line here is that this facility was unsafe. They knew it was unsafe, and something should have been done a long time ago. Our initial claim went to workman's compensation in 1994, and we, again, had a claim to the federal government

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in 2005. Mr. John Lipsitz is the lawyer who's taking charge of this case and has been very persistent and very valuable to us on my husband's behalf. Thank you.

CHAIRMAN MELIUS: Thank you. And then, John Lipsitz, I believe you signed up.

MR. LIPSITZ: Good afternoon. Мy name is John Lipsitz, and I'm an attorney in Buffalo, New York, and I represent Eleanor And I asked to be able to speak Tornabene. because I think this case illustrates the apparent irrationality of the system and why I have over the past several years gotten so many calls from so many frustrated claimants telling me that they were being unfairly treated. And why I believe the only solution to this kind of unfair, inconsistent, and, apparently, irrational system is to grant the Special Exposure Cohort for the people at Linde who worked there between 1954 and 2006.

Sam Tornabene, as Eleanor pointed out, worked at Linde. He actually was there

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from September 1962 through September 1993. Specifically, he conducted renovation work in Building 30 for a six-month period during the mid-1960s. This is a building which was identified by a 1976 Department of Energy radiologic survey as the most contaminated building at the Linde facility.

renovation work involved, The among other things in Building 30, breaking up concrete for hours at a time, which exposed high levels of airborne Mr. Tornabene to particles. alpha-emitting dust The exposure he was subjected to over at least a six-month period of time in Building 30 is the radiation exposure which type of increase the probability for the development type of non-Hodgkin's lymphoma of the eventually developed and died from. medical evidence in this case does not reveal or suggest any other competing risk factor for this gentleman's non-Hodgkin's lymphoma.

The renovation work was very

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specifically and in great detail described at
a hearing which took place in the New York
State workers compensation court in
approximately well, the period from 1994
through 1998. And it's important for you to
appreciate that we were initially seeking
compensation before the enactment of the
Energy Employees bill by going to New York
State workers compensation court. And in that
proceeding, we produced both written reports
and testimony from a well-qualified
pulmonologist in the Buffalo area, who is
board-certified in pulmonology, who testified
that the route of entry for the inhaled alpha-
emitting dust particles was such that it came
into his lungs, migrated to the lymph nodes,
and created the conditions for the development
of non-Hodgkin's lymphoma, which originated in
Mr. Tornabene in lymph nodes that were
proximate to his upper chest cavity.

We also heard testimony from Mr. Tornabene's oncologist, a Harvard-trained

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medical doctor by the name of [identifying information redacted], who very clearly offered his well-considered opinion that the non-Hodgkin's lymphoma was caused by this exposure. But, perhaps, in terms of medical, or rather expert testimony, the most telling thing is that we had reports and extensive health testimony from а physicist, [identifying information redacted], testified at length and over a several day period, and under intense cross-examination, that this, indeed, was a competent producing cause of Mr. Tornabene's cancer because the levels of radioactive dust that he inhaled were clearly injurious.

Αt the hearing, which, again, lasted on and off over a period of about four years, we heard testimony from Mr. Tornabene's coworkers, including most notably the [identifying information testimony of Now this is a third-party, so I redacted]. suppose his name may be redacted, but he's

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here right now, and he's in this room, and he described, again, at length, under oath, and under cross-examination, as well, how Samuel Tornabene and other men working with performed renovation work in the 1960s 1970s, and specifically described with reference to the floor plan of Building 30 the heavy pieces of equipment that Mr. Tornabene had to break off from the floor with the use of a jackhammer, and then move to another location and reinstall, giving rise to large amounts of dust in the air. And, again, these are the floors that were later studied by site commissioned surveys by the government, particularly, I believe it was the Department of Labor, showing high levels of alphaemitting dust particles, both fixed And, of course, these were all removable. liberated by the process of the jackhammer.

These workers did this work without any protection and without any special work clothing. They would go home with their

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regular clothing covered in dust and then have their clothes washed at home. They performed many renovation activities of this sort.

Now this was the testimony at the Workers Compensation Board. The employer produced testimony, as well, from experts, notably an expert in radiation and illness, [identifying information redacted], probably very well known to some of you who, at one time, worked for the federal government in the Public Health Service.

[identifying information redacted] took the position that non-Hodgkin's lymphoma is not caused by exposure to alpha-emitting dust particles. This was the same position that was taken by another expert offered by employer, medical doctor from the а the University of Rochester. They didn't say that there was no exposure; they didn't dispute that the exposure occurred; they dispute that the exposure was massive. just took the position that exposure to alpha-

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emitting dust particles does not cause non-Hodgkin's lymphoma.

Well, that decision dashed our hopes, considerably, and several years later when the government passed the Energy Employees Act, we gathered our evidence up, all of the testimony, including the transcripts of the experts, their reports, the testimony that [identifying information redacted] gave, and we submitted that again, this time with a considerable amount of hope because non-Hodgkin's lymphoma had been listed as a radiologic or radiogenic cancer by the government following the of the enactment Energy Employees Act.

Well, it's instructive to learn exactly how we've been bounced back and forth over the past five years. The claim was filed in 2005. A dose reconstruction determination was made in 2006. It was -- the claim was -it was recommended that the claim be denied because it 50 percent was less than а

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probability, and that was in 2007. We asked for a hearing, which we had in 2008. The final adjudication branch in 2008, in October, issued a final decision denying the claim. And all of this -- all up to this point, all we got was 10.24% was the estimated Probability of Causation, not one articulated reason, or opinion, or statement, or document individual human being who actually assessed the facts of this case.

We attempted to get the claim -the denial reconsidered. filed We Request for Reconsideration in November 2008. That addressed and denied in 2009. was Again, in May of 2009, we requested that the claim be reopened. A Request to Reopen was rejected because, according to the District Director, what we believed to be a revised Site Profile for Linde that had been issued in November of 2008, was, in fact, not a revised Site Profile, didn't really constitute new evidence, and, therefore, didn't change the

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dose reconstruction method.

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Then we filed a lawsuit against the Department of Labor. We filed a very thick lawsuit against the Department of Labor in August of 2009, and one month later we got a letter from the Department of Labor saying well, we've changed our minds. We're going to annul the initial decision denying the claim, the subsequent decision denying reconsideration, and the subsequent decision denying your request to have this matter So now we'll go back to the drawing reopened. board, and that was last year in September of 2009. say the least, it To is frustrating procedure that we've been going through.

This is a case where the claim was denied not because there wasn't proof of exposure, but because the exposure doesn't cause non-Hodgkin's lymphoma. Well, now we know that it does cause non-Hodgkin's lymphoma, but, apparently, it's not enough

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because it doesn't cause non-Hodgkin's lymphoma in this particular case according to no one in particular.

The government has produced no experts. It hasn't disputed the exposure. The cancer is classified as radiogenic, and this is really just an object lesson in how the people that have made these applications, when they follow it to the logical extreme, when they are persistent and continue to do it, will be frustrated at one turn after another.

And, in conclusion, I'd just like to say that when you look at a situation like this and you say that you're going to use surrogate data in order to determine whether to accept or reject a claim with such specific exposure evidence that you're using that's highly irrelevant and not at particular this friendly, at least to claimant. Thank you very much for listening to this presentation.

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1 CHAIRMAN MELIUS: Thank you. 2 one question I had, was this a Subpart E or 3 Subpart B claim? 4 MR. LIPSITZ: B. 5 CHAIRMAN MELIUS: B. Okay. 6 MR. LIPSITZ: Thank you. 7 CHAIRMAN MELIUS: Thank you. 8 Nancy Mendola Haug, I believe. Haug, okay. 9 Sorry. Thank you for letting 10 MS. HAUG: me speak today. My father, Peter Mendola, his 11 12 account number is [identifying information redacted]. He died of colon cancer in 1977. 13 He started working at Linde in 1951, four or 14 15 five months before I was born. He was a dead 16 man walking before I even came into this world, so he had his dose reconstruction, and 17 we think -- actually, the only people that are 18 19 left in my family are my brother and I. 20 mother is dead, my sister died of MS, which

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is, obviously, one of the things when he

brought home all this dust on his clothes that

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could have been part of the cause. My mother had a thyroid problem, which could have also been part of the dust that came home with my father.

You did the dose reconstruction, and the problem I have with all of this is Linde poisoned my father, bottom line. poisoned whether he 26.9, or was 29.6%. Poison is poison, and that's what they did; they poisoned him while he was there. followed to know exactly what air he breathing, what water he was drinking, areas he traveled in, and it's good for you all to sit there and just tell us it doesn't because you did matter your reconstruction. And the bottom line is he could have been in areas that were more exposed, and not exposed. And he was such a wonderful person, and to have him die that horrible death, just not right.

You couldn't have possibly monitored all the areas that he went into.

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The impact on my family was tremendous when he died. And then I was told when we went for the cohort group to be put into E, that if I was 23 years old, I was 25 when he died, if I was 23 years old in school, or disabled, I would have been approved. Well, I was 25 years old, and that doesn't mean the pain of losing my father was any less than when I was 23. And I still needed him, I still depended on him. I'm sorry.

(Off the record comments.)

HAUG: Then I got the letter MS. indicating that I wasn't approved because I was 25. And my brother did send one statement that he would like -- he's out of town and was unable to make it, and he wanted me to say that, "No studies large enough or over enough time have been run to disprove that low-dose ionizing radiation causes cancer that makes the possibility that it causes or doesn't cause it at 50 percent either way, which still meets the criteria for inclusion." And the

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bottom line is we lost our father because he worked at Linde, and he had so much life left to him. And it appears, just to me, okay, I may be 58 years old now, but it's been a long time without him. I named my son Peter so I would say my father's name every day for the rest of my life because my father's name was Peter and that's the only way I can keep his history or love alive.

It appears that, you know, you're all here doing this, and it's like a big circle. And it's almost like you're waiting for us all to die off so that you don't have to compensate us.

(Applause.)

MS. HAUG: And at this point in my life, I've lost my husband. I'm a widow, too. That doesn't make any difference, but the bottom line is, we take care of ourselves. We can afford to live, and this compensation actually doesn't mean anything to me. My father was worth a gazillion dollars to me,

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not \$150,000. That's useless. Thank you very much.

CHAIRMAN MELIUS: Okay. Thank you.

(Applause.)

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CHAIRMAN MELIUS: Karen Mortensen Noonan.

NOONAN: father, Royal MS. Μy Mortensen, worked at Linde from 1957 to 1965. He died at the age of 43. He was extremely healthy, active man. He sailed in In fact, as a teenager he built a the summer. sailboat with his father. My grandfather wanted to sail back to his native land of Denmark. He skied in the winter. He was on the ski patrol in World War II and delivered to the troops. I don't remember ever seeing him sit down. He was always doing additions to the house. He built our garage. He went to night school to get his degree as engineer. After World War II, he served on many government projects before working at

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Linde. And, yet, even though he died of spinal cancer, the dosage reconstruction said it was only 5% of the cause of his death. And spinal cancer is very rare. It comprises only 2% of all cancers, and its primary cause is radiation. And, yet, this is the lowest one I've heard here today, was below 5%.

My mother never recovered from his She had three small children. death. 12 at the time, and she made it clear that it was hard for her. And he brought out the best When he died, my mother. my reverted to her family way of being very cold and hard to live with. She Excuse me. started this claim 10 years ago, and I had to take it over at her death five years ago. didn't live to be denied. Well, we denied several times. And I have to say that this was very upsetting for her, just to first even start the claim because she got out the old pictures of my father working at Linde. Everything brought back the old memories to

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1	her. And I, myself, every time we were denied
2	and I had to reapply or get new documents, I
3	was always tempted to just let it drop because
4	it was very depressing and sad to live through
5	this again. However, every time I'd say to
6	myself, I have to do it for him. He did so
7	much for his country, and for you to just deny
8	that this caused his death, I think, is
9	unfair. So I hope that you reconsider. Thank
10	you for listening.
11	CHAIRMAN MELIUS: Thank you very
12	much. [identifying information redacted].
13	MS. SHAFFER: [identifying
14	information redacted] is my mother, and I will
15	
16	CHAIRMAN MELIUS: That's fine.
17	Either the podium, or the other mic, whichever
18	is
19	MS. SHAFFER: My name is Kathleen
20	Shaffer, and I'm [identifying information
21	redacted]'s daughter. My mother has filed
22	claim on behalf of her stepfather, Jesse

Hendershot, who worked at Linde from 1945, July of 1945 until only March of 1946. died at age 68 in 1977 of bladder cancer. And now the surprise here is that in hearing all the other people whose relatives worked at Linde, and talk about the dose reconstruction, and how people had worked there for 30 years and very long periods, my grandfather worked there for only a very, very short time; yet, his bladder cancer was diagnosed, and his dose reconstruction came back, and I believe it's 26% dose reconstruction, which is the highest of any that's here. So, obviously, there some great discrepancy in terms seems to be of how the dose reconstruction is made because my mother's claim was denied on the basis that he worked there for too short of a time. Tt. didn't meet the time constraints, as well as it didn't meet the 50% criteria for the dose reconstruction.

So it appears that -- we were told that they shortened his time frame for his

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employment, and it was only a number of
months; yet, he was diagnosed with bladder
cancer, one of the covered cancers in the
action. And his cancer was due to working at
Linde. And, again, as another woman had
previously stated, you don't know where the
person was when they worked at Linde, you
don't know the food they ate when they were in
the lunchroom at Linde, could have been
contaminated. You don't know any of that.
And I guarantee that every single one of you
sitting at that table, no matter where you go
to any of these meetings on behalf of any of
these agencies, and any of these families,
that if it was available for you to go and sit
in Building 30, which has been discussed here
about Linde, as it being the most contaminated
of all the buildings, I don't think that any
one of you would sit there for five minutes,
let alone work there for six months or 36
years. And I think that everybody needs to
realize the fact that these families are here

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on behalf of the efforts that their loved ones did on behalf of the United States government, and they need to, again, get away from dose reconstructions.

isn't any -- it's There shoot. Some people can be exposed cancerous type of thing for five minutes. Look at all of the claims that have resulted from rescue workers working at Ground Zero on 9/11. Some people were there one day, some people were there for months. It doesn't matter if you were there for five minutes, you might get a cancer that will kill you. can be there 36 years, and get a cancer that will kill you. It's the stuff that's there; it's not the dose reconstruction; it's not the percentages; it's the fact that there were all cancerous conditions in of buildings, and the government needs to do its job to compensate these workers for the jobs that they did and the fact that their families these people's lives, as well as

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1 people themselves who lost their lives, and 2 their capability, unbeknownst to them that 3 they were taking any type of risk. Thank you. 4 CHAIRMAN MELIUS: Thank you. there anybody who wishes to speak to Linde, 5 6 have I skipped over? There are some people 7 here that are -- a few people are listed for Hooker, which we will get to, and some people 8 that aren't identified. I'm not sure. 9 I just 10 don't want to miss anybody from -- yes, sir? PARTICIPANT: Bethlehem. 11 We'll get 12 CHAIRMAN MELIUS: Okay. 13 to you, also. But I just wanted to finish up on Linde, try to group people together. 14 Ιf 15 not, I think we'd like to hear from -- I think 16 Senator Schumer's office has a representative And she can introduce 17 here. Laura Monte. herself. 18 19 MS. MONTE: Му name is Laura and I'm on Senator Schumer's staff. 20 Monte, And I have a letter that I would like to read 21 into the record. This is a letter that comes 22

from Senator Schumer and Senator Gillibrand.

And this letter goes to the Honorable Kathleen

Sebelius, Secretary of U.S. Department of

Health and Human Services, and Dr. John

Howard, the Director of NIOSH.

"Dear Secretary Sebelius and Dr. Howard, we are writing today on behalf of sickened nuclear workers and their families that have been denied a fair hearing regarding compensation benefits under the EEOICPA.

Over the last decade, regulations that have been implemented by the Department of Health and Human Services have fulfilled the Congressional intent of this landmark remedial compensation program representing a claimant favorable paradigm. end, the Linde Ceramic ΤО that Special Exposure Cohort Action Group submitted a petition for rulemaking to the Department of Health and Human Services on September 28th, 2009.

This petition outlined suggestions

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for needed reform to the EEOICPA administrative regulatory framework. The petition has been coordinated by advocates and stakeholders on the front lines from workers' representatives and workers themselves, who deal with the onerous and bureaucratic burdens forced upon sickened workers and families under the current EEOICPA process.

One of the primary issues raised in this petition deals with the inappropriate use of surrogate and/or coworker data in this SEC evaluation process. The SEC program was designed avoid the difficult and to technically complex dose reconstruction program in order to provide sickened nuclear workers with fair and equitable compensation under EEOICPA. The use of surrogate -- I'm sorry, I skipped that line. The use of surrogate and/or coworker data in the evaluation process reflects an analytical framework used by NIOSH that is designed to grant SEC petitions only as a last resort.

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This analytical paradigm unfairly limits the ability of workers to be granted relief from a dose reconstruction program that has become a bureaucratic and technically incomprehensible nightmare for the lay person.

The fundamental inability for sickened workers to understand the reconstruction program deprives these workers of their basic right of due process EEOICPA. The claimant or petitioner cannot understand why their claims are being denied by NIOSH and, ultimately, by the Department of Labor due to the inherent technical nature of dose reconstruction provided reports to Consequently, claimants. even claimants provided with the right are appeal denied decisions, that right is useless when someone is functionally precluded from understanding why their claim was denied in the first instance.

To expect a claimant to be able to develop in advance an effective appeal is

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impossible if they do not understand why the claim was denied in the first place. This fundamental absence of due process undermines the essential purpose of passage of the EEOICPA. That is why we fully support the reform agenda outlined by the Linde Ceramics SEC Action Group.

We urge the Department of Health and Human Services to review and consider these reform measures without delay within the 10-year EEOICPA review currently underway at The men and women that have served our NIOSH. nation and were unknowingly exposed radiation at nuclear facilities around the country deserve to have their claims evaluated in a fair and equitable manner. Fairness and equity can only be achieved through clear implementation of a standard of claimant favorability. Moreover, fairness and equity only realized be when the regulations are applied in a manner reflects the remedial nature of this vital

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compensation program.

Thank you for your attention to this critical request. Sincerely, Charles E. Schumer, United States Senator, Kirsten E. Gillibrand, United States Senator."

CHAIRMAN MELIUS: Thank you very much, Laura. And then I believe we also have Bill Greeley from Congressman Higgins's office.

MR. GREELEY: Good evening. My name is Bill Greeley, G-R-E-E-L-E-Y, and I'm here representing Congressman Brian Higgins. I'd like to welcome the members of the Board to Western New York. I'm kind of proud to think that you're here today because I was one of the people that advocated for a Advisory Board meeting in Buffalo, and Niagra Falls naturally is the next best thing.

In January of 2005, I began working for Congressman Higgins after he got elected. On April 30th of 2010, I officially retired, but I made a commitment to the

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Lackawanna Action Group that I wouldn't abandon them, and with the Congressman and his senior staffer's permission, I would continue to work on behalf of the members of that Action Committee. With me today, this or evening, is the Congressman's District Director, a young lady by the name of Megan Corbett. And Megan has been my boss for the last four and a half years -- or five and a half years. So, I'd like to, on behalf of my Congressional colleagues in some of the other offices, thank you for coming and listening to these stories on behalf of all the workers that down with such a serious have come illness.

When I started in 2005, one of the first calls I received was from a gentleman by the name of Ed Walker. And most of you know who Ed Walker but Ed was, was an uncharacteristic type of guy. Не was He was a bricklayer, worked at tradesman. Bethlehem during that period, was

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remodeler, but he was a marvel. Ed could talk about this program and argue with attorneys, with doctors, with scientists, and with elected officials and their staffs. I used to just marvel at Ed. And Ed and I got to be very, very good friends. Practically every Friday afternoon, late in the afternoon when I was trying to sneak away from work, Ed would call and give me the week's rundown everything that had happened. And I'd like to think that one of the reasons that you're here today is out of Ed's legacy. He just was a wonderful guy. I enjoyed his company very much, and his wife is a real sweetheart. And I'm sure that tomorrow you're going to hear from her.

You know, this community a little over 10 years ago was glued to the television about a famous murder trial. And I can still remember the day when a piece of evidence, a bloody glove, was put on the defendant in the charge on his hand, and his attorney jumped up

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and said, "If the glove doesn't fit, you must acquit." Now you're sitting there thinking why is he bringing this up.

Well, I'd like to just point out that part of this program, when it's using surrogate data to fill in blanks for information that doesn't exist at a facility like Bethlehem is wrong. And just like that glove, the evidence got thrown out, I'd like to point out to you that this surrogate data, which doesn't fit at Bethlehem Steel should be thrown out, and an SEC has to be approved for the workers and the claimants at Bethlehem Steel.

(Applause.)

MR. GREELEY: Thank you. Now one of the things that I brought with me was a statement from the Congressman, and I think that Mr. Katz has passed it out to everyone. Brian would have loved to have been here tonight or tomorrow to address you, but the House is meeting in session, and there's going

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to be votes this evening and tomorrow.

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of the things that One the Congressman points out in this is that for the last five and a half years, he's been an advocate for the workers at Bethlehem. I could just read some of this, "Local steel workers and their families have suffered for decades from the toxic exposure to uranium dust at the former Bethlehem Steel Plant in Lackawanna, New York. I wish I could say that the supervisors at Bethlehem didn't fully know the risk that the workers in the uranium rolling facility were being subjected to in those early days of the Cold War, but, fact, huge gaps of monitoring data for the facility, mean will never really know we whether proper precautions were carried out, and carried out at all. In addition, reports of exposed -- or suggested, there are major inadequacies in the attempts to use surrogate data to reconstruct the toxic exposure."

One of the things that Brian

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points out is that the late Ed Walker made this issue the cause of his later life. about the intricacies of these knew more facilities than most, and he knew, before many others would admit, that the system set up to deal with the Bethlehem Steel workers was flawed and needed to be fixed. That's why he led the effort for Bethlehem Steel to be placed as a Special Exposure Cohort. His memory lives on today in the dozens of local families who have their come to express support for fair relief. As you deliberate, the Congressman

As you deliberate, the Congressman says, "I urge you to reflect on their concerns and the fallacies of the system that they have been subjected to, and make a favorable recommendation for their petition."

And just before I walk back to my seat, I'd like to just reiterate, "If the glove doesn't fit, you must acquit."

(Applause.)

CHAIRMAN MELIUS: Thank you, Bill.

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1	Thank you for your dedication even after
2	retirement, so appreciate it. We have some
3	more people listed. I'm going to go through
4	them first. I don't know if we have people in
5	the we have two more, also. But let me
6	start at the top of the list again.
7	[identifying information redacted]. Is there
8	a [identifying information redacted] here?
9	May have just sometimes people just sign
10	the wrong list as they come in, and so forth.
11	Mary Girardo? Okay. This only
12	has a first initial, so I don't know, someone
13	from Hooker, P. Scremmin or Scremm. Okay.
14	Scremmin it was, okay. If you'd like to step
15	to either mic. That one may be a little bit -
16	_
17	MS. SCREMMIN: Thank you.
18	Actually, some of my
19	CHAIRMAN MELIUS: You need to pull
20	it can you help her?
21	MS. SCREMMIN: Some of my
22	questions have been answered, and some I

realize were probably misplaced at a meeting of NIOSH because I wondered about the Special Cohort for Lake Ontario Ordnance Works. And if anyone here knows why that -- to be in the Special Cohort, the employee had to have 250 days of employment. And, also -- so, what is the magic about 250 days?

Secondly, there was a deadline of December 31st, 1953. Should I address this question to Department of Labor, or can someone from NIOSH answer the question?

CHAIRMAN MELIUS: Well, I think we can answer the 250 day question. When the ordinance was originally passed by Congress for the Special Exposure Cohorts that were included in the ordinance, the law, they used to qualify under the Special -- those Special Exposure Cohorts had to have worked at least 250 days at the facility, I believe, for three of those cohorts. So when the regulations were written, the 250 days was taken into the regulation for facilities. Unless there was a

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1	very acute, very high exposure, then there can
2	be exceptions to that.
3	MS. SCREMMIN: Okay. All right.
4	Was it just an arbitrary number of days, or
5	did it have anything to do with the amount of
6	radiation exposure, or medical problems, or
7	was it just 250 days, an arbitrary date?
8	CHAIRMAN MELIUS: In some ways,
9	it's arbitrary.
10	MS. SCREMMIN: Okay.
11	CHAIRMAN MELIUS: It represents
12	one year of work at a facility.
13	MS. SCREMMIN: Yes.
14	CHAIRMAN MELIUS: And there would
15	be some an opportunity for significant
16	amount of exposure.
17	MS. SCREMMIN: Okay. So, to be
18	included, it would be an all or none. The
19	employee had to have 250 days, or he was
20	the employee was not included in that cohort.
21	CHAIRMAN MELIUS: Correct. That
22	is adjusted for overtime, and people worked

1	weekends, so there's some adjustment can be
2	made for it, for longer work schedules.
3	MS. SCREMMIN: All right. Why the
4	end date of December 31 st , 1953?
5	CHAIRMAN MELIUS: That part I
6	can't answer directly because I think it has
7	something to do with operations. But does
8	anybody Stu, are you still here, or someone
9	to
LO	MR. HINNEFELD: As I recall, our
L1	evaluation indicated that the from that
L2	point forward there was sufficient data and
L3	the dose reconstruction was feasible. That's
L4	my recollection, but I'm speaking from memory.
L5	CHAIRMAN MELIUS: Yes. Someone
L6	will follow up, and that may still be under
L7	consideration. We just don't recall.
L8	MS. SCREMMIN: Second question,
L9	maybe the NIOSH representatives can answer,
20	what, if any, weight in the decision making,
21	what, if any, weight was given to smoking in
22	the employee's history?

1	CHAIRMAN MELIUS: That would
2	depend if it was being done as part of the
3	Special Exposure Cohort, there would be none
4	direct. I mean, that's
5	MS. SCREMMIN: I don't understand.
6	CHAIRMAN MELIUS: Yes. Stu, do
7	you want to explain the
8	MR. HINNEFELD: If the claimant
9	smoked and the cancer involved is lung cancer,
10	there would be some adjustment to the risk
11	number, and, therefore, the Probability of
12	Causation outcome as a result of the dose
13	reconstruction. For any other cancer, there
14	would be no effect from smoking.
15	CHAIRMAN MELIUS: Right.
16	MR. HINNEFELD: For any other
17	cancer, other than lung cancer. For lung
18	cancer, there is I'm sorry, she didn't
19	understand what I said.
20	CHAIRMAN MELIUS: Yes.
21	MR. HINNEFELD: For lung cancer,
22	there is an adjustment to the risk in terms of

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how -- essentially, how high the PoC goes per unit of radiation dose based on the person's And that is based on the smoking history. epidemiological evidence that is available for of the occurrence cancer in exposed populations. CHAIRMAN MELIUS: Yes. The

CHAIRMAN MELIUS: Yes. The science that is used as the basis for the calculation takes into account -- that into account.

MS. SCREMMIN: Thank you very much for the opportunity.

CHAIRMAN MELIUS: Okay.

MS. SCREMMIN: And I will say that over the past nine years, I have certainly had aood education in bureaucracy. а Unfortunately, I have not gotten education in radiology or the effects of radiation. All I have learned is how much I don't know and how I do not understand any of the charts that were sent to me. So, I thank you very, very much.

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CHAIRMAN MELIUS: Well, it's -- we understand it's complicated, and I will say that NIOSH is making some efforts to make some of that information more intelligible and easier to understand. It is difficult. We know how frustrating that can be.

Amy Witryol, I believe it is.

MS. WITRYOL: My name is Amy Witryol, and I live in Lewiston, New York. And, first of all, I'd like to thank the speakers that I've heard for very thoughtful and very intelligent remarks.

Secondly, I'd like to clarify a comment, I believe by Mr. Greeley. Niagra Falls is not the second best thing to the City of Buffalo. We think Niagra Falls is a pretty darned good place to be, and the best place to be with those choices. And we also appreciate the diversity of opinion, perhaps, in that view, because we love Buffalo, as well.

I'd like to read comments first by a friend, and then, secondly, add my own

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comments. These comments are from [identifying information redacted], who formerly lived in Youngstown, New York, which is just to the north of here, just beyond Lewiston, and presently lives in Sheboygan Falls, Wisconsin. She's а chemist and engineer with an enthusiasm for historical records and, might I add, an excellent well. [identifying analyst, And as information redacted]'s statement is as follows.

accept the "Please following may be relevant information which to the exposure of Hooker employees at the former Lake Ontario Ordnance work site, known to us as the LOOW site. Keep in mind that there are investigation serious gaps in the of radioactive contamination conducted at this site by the Atomic Energy Commission, the Department of Energy, and, currently, the U.S. Army Corps of Engineers, Buffalo District, which fully addressed in this are not

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1953, the U.S. Atomic Energy Commission contracted with Hooker Electrochemical Company to construct operate a boron-10 isotope separation plant at the LOOW site. Although boron-10 is nonradioactive, Hooker workers were working on a contaminated site and were likely exposed to excess radioactivity. In addition to boron-10 production, Hooker personnel were also employed in the cleanup of radioactive contamination the LOOW site at and the disposal of nuclear reprocessing waste burial and burning.

Boron-10 production took place in the former LOOW power plant known as Building 401, which was found to be contaminated with radioactivity and now, just this year, is scheduled for demolition by the Army Corps of have identified Engineers. Past surveys significant radioactive contamination lockers 401, workers' in Building the

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question of whether workers had the potential to bring contamination into the building should also be investigated.

The boron-10 plant was operated from 1953 to 1958 by Hooker. It was placed on standby for six years, then reactivated and operated from 1964 to 1971 by the Nuclear Materials company, NUMEC.

The plant was housed in Building 401, which was originally the LOOW power Between 1952 and 1953, immediately plant. prior to the construction of the boron-10 plant, Building 401 was used to store nuclear reprocessing wastes from the separations process research unit, SPRU, at the Knolls Atomic Power Laboratory, which we refer to as KAPL, in Schenectady, New York. The wastes were highly radioactive, and contaminated plutonium and mixed fission products were included in these wastes.

An estimate of 408 curies of mixed fission product waste containing traces of

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plutonium were shipped to the LOOW. The amount of plutonium waste sent outside of that contained in the fission product waste has not been estimated. As part of Hooker's contract with the Atomic Energy Commission to operate the boron-10 plant, Hooker was also contracted to maintain adjacent storage areas on the LOOW site.

In this maintenance capacity, and as part of a 1953 effort to clean up the site, Hooker personnel were involved in storing, handling, and burying radioactive material at the LOOW. These activities were carried out under Hooker's contract with the AEC," which I'll provide to you in a list of documents in a package I have to provide you with these "Inadequate storage conditions for comments. the KAPL Schenectady waste at the LOOW led to concerns about safety and a desire by the Atomic Energy Commission to dispose of The AEC enlisted wastes. Hooker Electrochemical to assist in disposal of the

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KAPL wastes, and directed KAPL to assist Hooker in the disposal operations.

At the end of 1957, Hooker workers began preparing over 1,000 drums of KAPL waste for shipment to Oak Ridge for disposal. first shipment of the KAPL waste to Oak Ridge from LOOW took place in January of 1958. Hooker Electrochemical personnel were directed to burn the combustible KAPL wastes but encountered problems." Again, reminder, KAPL the nuclear waste was reprocessing waste from Schenectady, the naval reactor in Schenectady.

AEC specified environmental "The sampling before and after the burning of KAPL waste, as well as urine analysis for Hooker personnel handling the KAPL wastes. The results of these two monitoring exercises have not been published. In September 2009, the Engineers released Army Corps of scoping document to the public requesting input associated with the on any issues

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proposed demolition of Building 401 on the Niagra Falls storage site. The scoping document failed to address the past potential exposure of Hooker Electrochemical personnel and the Nuclear Materials company workers who worked in contaminated Building 401.

Requests for the Army Corps of Engineers to publish all associated documents detailing the nature and extent of the radiological contamination in Building 401 have not been met, and it is concerning that evidence of radiological contamination, which may be of value in determining worker exposure will be lost when Building 401 is demolished."

Again, that's scheduled for this fall.

"One radiological survey from 1955 has been located, which records americium-241 being present in Building 401. In 2007, the U.S. Army Corps of Engineers released remedial investigation report for the Niagra which Falls storage site recorded the detection of plutonium-239 in a Building 401

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That includes the comments from [identifying information redacted], and I have for you 15 references of documents dating back to the 1940s in support of these remarks. I would now like to offer some of my own remarks.

disconcerting very today when we still have an opportunity to determine what the radiation exposure is, that workers going back to the 1940s, `50s, `60s, and `70s may have been exposed to, but also with the community living here today may be exposed to during the demolition of these I don't want to take time away facilities. from other people who want to speak tonight to go into detail about the 7,500 acre former Defense site that we have located in the towns of Lewiston and Porter, that in many ways is very closely tied to the Defense Department and Department of Energy operations that took place in Niagra Falls and in Buffalo.

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also, for example, the repository of some of the wastes from the Linde site, as well.

But what I will say very simply is we have multiple state and federal agencies involved at this site, and should your work, or referral of this information to any other board involve soliciting information from a state or a federal agency, may I urge you to contact [identifying information redacted] for the completeness and accuracy information, because having spoken with boards citizen advisory and restoration advisory boards all over the United States, when it comes to contamination from weapons production, many of these sites are bigger and badder than what we have, but in talking with these advisory boards, they tell me that bar none, when it comes to agency conflicts of interests, that the Lake Ontario Ordnance Work site, at both the state and federal level, because of private and government activities, has no peers.

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1	So thank you for coming to Niagra
2	Falls to hold this meeting, and I join others
3	in wishing you a very pleasant visit, and hope
4	that this Board honors the sacrifice made by
5	all of the people who were spoken about today.
6	Thank you.
7	CHAIRMAN MELIUS: Thank you. And
8	we also appreciate the the information is
9	always helpful. John Martino. Is John here?
10	MR. MARTINO: First of all, I'd
11	like to thank you for reopening my claim,
12	which has been laying dormant for three years,
13	and that of several other people here.
14	I have two points that I want to
15	make. I had thyroid cancer that I had it
16	operated on, it metastasized the lymph glands
17	up the side of my face. I had two growths
18	removed from my right ear, which is not on my
19	record. My ear is 80 percent shot. I wear a
20	hearing aide.
21	When I had my cancer diagnosed, I

was in Roswell for 10 days.

22

They came to me,

I didn't go to them and ask these questions. They asked me where I worked, and I told them at Hooker Chemical. They tried to get information back then. They tried. I didn't ask them to, but they told me it was caused by radiation. They checked me after my surgeries every six months for a year, then every year after that for 10 years. It came back in two years. That's not on my report.

had to qo on to treatments, knocked it out in another lymph gland, I had another scar here, another scar there. then monitored me for another eight years and says I'm okay, until they found that I had a blood cancer called monoclonal gammopathy, which in my letter here is not covered. they checked me every three months. I have an oncologist, hematologist. I go every three months, then I go every six months. three times I was threatened with chemo, my numbers, my IgM factors went way up above the danger point, and he says well, let me wait a

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couple of weeks, we'll take the blood again. Mine go up and down, up and down. But he says they keep checking me these past two years. People think I look pretty good. I lost 18 pounds in the past year and a half.

Why I'm saying this is because administrators and managers are processors of information. We're like computers. We process the information given to us, garbage in, garbage out. We get good information, we get good results.

I worked in maintenance at Hooker Chemical. I noticed in your literature that you talked to a couple of yard birds, you called them, and a couple of engineers. The maintenance people went in every building that they have. We crawled on the beams. We sprayed the dust off of us with air hoses. We had cloths, we didn't have masks in those days. They came with masks about five or six years later, but used to wrap towels around us because of the dust, and spray our hair, and

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stamp our feet. We crawled on the beams on the dust that laid up there in those buildings for years, and years, and years. God knows what it was, but that's the information that I'm saying that maybe you don't have that kind of information, or what kind of illnesses some of us have.

I used to be a 34, these are 32s, and they are falling down on me. And my doctor says, John, what's happening? happening? We've got to check you again. they're watching my numbers. Ι mean, doctor here knows what IgA, IgG, and IgMs are. And I keep them, too. I watch them, too. And when he says that I have to have chemo, I say what? And that's not even recognized. And it came from exposure to radiation. Tt.'s the literature on that disease, That's one of the causes, but that's not covered.

I get around good. It hasn't stopped me that way, but I know what's coming

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down the road. But I was very happy, I smiled when they said they're reopening my claim, but I got a little bit upset when they said that's not covered. But the other things I have, I say geez, I'm glad I had those. Those are covered. Thank you. Yes, it's funny. I know, I laugh, too. You have to look at the good side. I tell my wife, the glass is half full, it's not half empty. Maybe I'll get lucky.

CHAIRMAN MELIUS: I will tell you that when the Board here, we are just starting our review of NIOSH's report, so this is the first we heard about it was today. We just got it a week ago when everybody else did, if you got it when it was released. But one of the things we do pay attention to, careful attention to, and are very cognizant of are the maintenance workers. We understand that at many, many facilities that the exposure for maintenance workers is different. You do work in many different parts of the facility, and

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we always want to pay attention to make sure that's taken into account in whatever -- however this is being evaluated, both the SEC as well as the dose reconstruction.

MR. MARTINO: I ran wires into manholes there. Where do you think the water was washed in when they washed the buildings down, in the manhole, but we pulled cables down there, we ran wires. We crawled on beams, and all over the place, and dug holes, and ran conduit, and stuff.

I know, that's the information that maybe needs looking into. But my claim is in, but I'm talking for some of the other guys, construction guys that have their -- aren't recognized in this time around again. We've all been denied several times. This gentleman back here has been denied several times, too, and he's a contractor. And that's the information that is not getting processed properly and may be overlooked.

CHAIRMAN MELIUS: Yes. We have at

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1	least one member of this Board who can
2	probably share stories with you about crawling
3	around a facility, and so forth, at least more
4	than one, so it is appreciated.
5	MR. MARTINO: Well, then Mike will
6	understand what I'm saying.
7	CHAIRMAN MELIUS: Yes.
8	MEMBER GIBSON: Journeyman
9	electrician, 22 years
10	MR. MARTINO: There you go.
11	CHAIRMAN MELIUS: Yes.
12	MR. MARTINO: You crawled on many
13	beams, too, run that conduit, and get that
14	dust. That's what I'm saying. I'm glad mine
15	is reopened, and maybe somebody else will get
16	lucky here. Thank you.
17	CHAIRMAN MELIUS: Thank you. We
18	have a Donald Allan here from Bethlehem?
19	Donald.
20	MR. ALLAN: Good afternoon. My
21	name is Donald Allan. My father's name was
22	Boyce Allan. He was a steel worker, Bethlehem

Steel. He started working there in 1952, and like so many African Americans, he came up from the south for the American dream. Well, his dream has turned into a nightmare because I lost my father. He worked in that plant. In fact, two summers he had me work there. I'm glad Bill is here sticking up for the Lackawanna guys, Bill, we need you, and tell the Congressman I said thank you very much.

You know, he worked at Bethlehem Steel, and he had me come out there two And that's when I made up my mind summers. that the plant wasn't for me because that plant was short for plantation, trust me. Ιt was dirty, it was nasty. He went in Gate Three every day from 1952 until he retired in 1979. He went to Bethlehem Steel every day, some days he worked double shifts. I remember him coming home with dirt all over his face, bringing his clothes home, and mУ washing them. By the way, mУ mother [identifying information redacted] years old

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right now, and she has all kind of health problems because she washed those clothes that my father brought home.

Now, for this, or anybody to say that Bethlehem Steel is not a part of this, you need to go and see that plant, you need to go in there. I've been to every meeting going back, we've been through two Presidents, two Presidents. Okay? Clinton went through this, then Bush, he just, you know -- I'm not even going to get into that. But we went through 16 years, Presidents, terms, two nothing was done. People were denied, people I think this whole thing is were dying. waiting for everybody to die, but let me tell you something, we got families. And you know what, I've told my sons, you keep on this, and we're going to fight the fight to get this done, to get their people the right due on You sit here and say the buck don't this. It doesn't stop with you. stop here. whoever the bucks stop with, they should be

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here tonight.

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This man just got finished telling you what he's going through. That's travesty. And for somebody to say that they couldn't get in the plant -- one meeting I was at, this gentleman stood up and said well, we can't get in Bethlehem Steel to review it. Don't tell me about the feds because if I committed a crime in my house, you'd find a way to get in there, so don't tell Bethlehem Steel in and do the can't go testing.

For you to say, for this Committee, or anybody else to say that the men that worked at that plant at Bethlehem Steel did not suffer, my father was a scrapping big guy. When my father died, he weighed 148 pounds. Okay?

I had to put him in a nursing home because he could no longer get around. His lungs were shot. He wasn't a smoker, either. His lungs were shot, his liver. He couldn't

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go to the bathroom by himself; he had to wear a bag.

I mean, if you think that plant didn't do that, if anybody can sit there and say that Bethlehem Steel didn't harm these men, and that the lady that got up and said about Niagra Falls being a good place, I'm glad you're here, no matter where you were, I was going to come. But I called some people to be -- they couldn't get here to Niagra Falls. I found some people, and I called them, people had passed away.

And that's what I'm saying to you, I don't know if this is a thing to wait for everybody to be gone, and then you don't have to pay anybody. I mean, it's ridiculous. will this, Т tell you get past the bureaucracy, get past the government stuff, and do this with your hearts. Think about the people and their families and these widows that are here, and their families, and do the right thing. And that's all I've got to say.

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1 And, Bill, again, thank the Congressman for 2 sticking with us, and tell him I said hello. 3 Thank you, Billy. 4 (Applause.) Thank you, 5 CHAIRMAN MELIUS: Mr. 6 Allan. Mr. Brooks, Larry Brooks. 7 MR. MIDDLEBROOKS: Yes, it's Middlebrooks. 8 Middlebrooks, 9 CHAIRMAN MELIUS: 10 yes. 11 MR. MIDDLEBROOKS: Good evening. My wife is not here, and I started the ball 12 13 with this for my mother-in-law. I did all the leg work. First, I was told they had no proof 14 that he worked at Bethlehem Steel, so we got 15 16 pay stubs, we had his badge, we had all these different documents to show that he worked 17 I went out on Route 5, they couldn't 18 19 give them me a anything. I had to send to 20 Then they sent me things. Pennsylvania. we had to prove all these medical things. 21

went to Buffalo General, got every document

that he ever had when he went in there.

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us, what they They sent through to find out if we qualify or not. Over 50%, over 50%. We talked Oh, you should be getting representative. your money in 30 days. Thirty days come by, didn't get anything. Call them again. okay, there might be a little delay, 60 days. Call them again, didn't get anything. thing you know, we had a meeting out on Cheektowaga, Four Seasons Hotel. We was out My brother-in-law, which this is his grandfather I'm representing, he had a packet that we passed to Louise Slaughter, Hillary Clinton, and some other people there that we gave them to.

Louise Slaughter, anyway, they identified themself as Louise Slaughter's office called and said there was a mistake, that we didn't qualify, although these professional people told us that we did. Over 50 -- I got the paperwork at home. We still

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have that.

And the other thing is, Bethlehem Steel -- I work in a plant. I work in a plant right now. I worked there in Republic Steel. You walk back there, dust is everywhere. I worked in BOF, dust go everywhere. When you're pouring that steel, when you in the cast shop. I work at Dominican Brass now, same thing, dust go all the way through the plant. You don't have to be working at one special area to get that.

Now the thing is, also, I remember we had dead fish. We had to clean up the lake. We had seagulls there at one time. Nobody ever identified why they was dead, why they was out there. We just had to clean it up. I wonder why.

I wonder here now, are you our opponent, or are you here to help us, because it seems like every time we give you what you need, you'll find something in there to say oh, there it is, no, you're not qualified.

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Cancer is cancer. I don't care, you know -- that's what they die -- have you seen these people when the final stages start coming in, like he said, losing all this weight, can't make it to the bathroom, can't take care of themself. These are men, these are strong men, and they lost their pride because they needed help to do certain things.

Now what we need you to do is to look in there and say oh, there it is, he qualifies. Find something in there to make us qualify, instead of finding something in there to make us not qualify. We need you all to get on the job. It seems like every time you all come here, we go through the same thing. We sit here, we discuss this, you all bring a group of people, where you can answer any question it is, if not, you take it back, you dissect it, you come back with the answer why we don't qualify.

All of these men and people worked out there. The City of Lackawanna, you think

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1	that dust didn't go up in the air? They
2	should even be here. You just pertaining to
3	the people that worked there. What about the
4	outside contractors that came there for the
5	vending machines and different stuff? There's
6	a whole lot of people that you're missing
7	here. But still, again, you don't want to
8	take care of the people that's there. Thank
9	you for your time.
10	CHAIRMAN MELIUS: Thank you.
11	Anybody else? We've run through the list. I
12	don't know if there's anybody else that would
13	like to comment, or on the phone?
14	MR. BEYERLEIN: Yes, sir. I'm a
15	reporter from Dayton, Ohio. And I was
16	wondering if any action had been taken yet on
17	the Mound SEC petition?
18	CHAIRMAN MELIUS: No, there has
19	not been any action yet. We're having to
20	confer. There's some issues about how to
21	address that. Maybe by tomorrow sometime.

MR.

BEYERLEIN: Okay. How can I

1	learn what the outcome of that is?
2	CHAIRMAN MELIUS: Yes, can you
3	give us a number where we can contact you, and
4	then we will make sure that you hear?
5	MR. BEYERLEIN: Certainly.
6	CHAIRMAN MELIUS: Name and number,
7	I guess.
8	MR. BEYERLEIN: My name is Tom
9	Beyerlein, spelled B-E-Y-E-R-L-E-I-N.
10	CHAIRMAN MELIUS: Okay.
11	MR. BEYERLEIN: I'm with the
12	Dayton, Ohio Daily News. And my phone number
13	is [identifying information redacted] .
14	CHAIRMAN MELIUS: Okay. We'll
15	make sure somebody contacts you tomorrow and
15 16	make sure somebody contacts you tomorrow and let's you know what the decision is or where
16	let's you know what the decision is or where
16 17	let's you know what the decision is or where things stand tomorrow.
16 17 18	let's you know what the decision is or where things stand tomorrow. MR. BEYERLEIN: Very good. Thank
16 17 18	let's you know what the decision is or where things stand tomorrow. MR. BEYERLEIN: Very good. Thank you, sir.

the phone that wishes to comment? Yes. Okay. Fine. We have someone from the audience. If you'd get up and -- since we don't have your name written out, it's important that you identify yourself for the record.

MRS. MORTON: I'm Mrs. Morton from Niagra Falls, New York. I just want to try to get something through my head that I don't understand. My husband died with cancer. suffered for months. Okay. Then government comes out with this oh, everybody is going to get all this money. They came looking for us. We didn't go looking for So I go through all the paperwork. them. It'll be what, nine years now? And I get all the paperwork, and the first thing that happens well, they lost it. Luckily, I had enough sense to have copies. Okay. So that went on and on. And three years ago, I got denied.

Okay. I got my denial letter today, and this fellow that I know gets his

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acceptance the day later. He filed in 2004, I filed in 2001. My husband worked at the Hooker for 40 years. He worked all over the place. This fellow also worked in the certain area with him. He worked there eight months. He got a thing like was up on that board, that's probably the gentleman they talking about, for 76 point something radiation. My husband got 4.5, 40 years 8 months working in the same area.

Now how did the dosimetry come up with these figures? And I see this gentleman every day enjoying life, having his coffee, going on his trips. I'm happy he got it, but my husband has been dead 20 years. I'm 83, still working. I just don't see where that was fair.

CHAIRMAN MELIUS: I don't think we can provide a complete answer to that. I will say two things, one is it does depend on where a person worked and the type of exposures they may have had and what was recorded for their

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1 exposures. But, secondly, that under -- your 2 husband worked at Hooker? 3 MS. MORTON: Hooker, 40 years. Then this is the 4 CHAIRMAN MELIUS: 5 first the Board here has been involved in 6 Hooker, so we will be reviewing the Hooker 7 site and looking at how these are done and seeing if everything is being done correctly, 8 appropriately. And should there be a Special 9 Exposure Cohort there in response 10 11 petition that people made. 12 MS. MORTON: I gave up. Ι 13 rejected twice. CHAIRMAN MELIUS: 14 Okay. 15 MS. MORTON: I haven't -- I don't 16 have the energy to keep fighting this. suppose -- the last thing they told me was to 17 resubmit it. Do you know how many hours and 18 19 days and months and years that I spent looking 20 for this stuff? I mean, I'm not complaining. I'm healthy. I work every day, but it just --21 22 when he told me that, I just -- I was happy

1	for him. I just couldn't figure it out
2	because they worked in the same areas.
3	CHAIRMAN MELIUS: Yes, I can't
4	understand that either.
5	MS. MORTON: Eight months, eight
6	months to 40 years. And they say well, the
7	dosimetry, and this and that. Well, how do
8	they figure it?
9	CHAIRMAN MELIUS: Yes.
10	MS. MORTON: Well, thank you very
11	much. Maybe I should call and reopen it.
12	CHAIRMAN MELIUS: Okay. Well,
13	think about that.
14	MS. MORTON: Thank you.
15	CHAIRMAN MELIUS: Anybody else
15 16	
	CHAIRMAN MELIUS: Anybody else
16	CHAIRMAN MELIUS: Anybody else that would like to speak?
16 17	CHAIRMAN MELIUS: Anybody else that would like to speak? MR. RAMSPOTT: Dr. Melius?
16 17 18	CHAIRMAN MELIUS: Anybody else that would like to speak? MR. RAMSPOTT: Dr. Melius? CHAIRMAN MELIUS: Yes.
16 17 18 19	CHAIRMAN MELIUS: Anybody else that would like to speak? MR. RAMSPOTT: Dr. Melius? CHAIRMAN MELIUS: Yes. MR. RAMSPOTT: Are you opening

MR. RAMSPOTT: I was trying to be polite to the local folks. My name is John Ramspott.

CHAIRMAN MELIUS: Yes.

St. MR. RAMSPOTT: In Louis, Missouri, and I'm calling regarding the General Steel Industry site in Granite City, Illinois. And I'd like to respectfully that the Board take under request consideration tasking SC&A with the detailed General Steel review of Industry's **NRC** provided FOIA 2010-0012, and ask the thing, that SC&A be tasked to review the White Paper on portable radiography recent sources at General Steel Industries prepared by David Allen, May 2010.

The reason I'm asking that this happen, the topic did come up in the recent week or so ago Work Group meeting for General Steel Industries, and if I understood correctly, that was going to be possibly discussed at this meeting, and offered to the

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full Board as an option that they be tasked. We believe there is pertinent new information, which originally we were told didn't exist, and it turned up in the form of, I guess, of new information regarding 1,015 pages source materials, safety procedures, monitoring at General Steel Industries that, [identifying information apparently, redacted]'s FOIA, second or third FOIA request was successful in obtaining.

And after a very brief review that we've seen by NIOSH, we believe there's pertinent information that was overlooked that should be included, and we would look forward in assisting in that possibility. And I, of course, appreciate your time. I didn't want to interfere with what you have ongoing locally there, but thank you for your time.

CHAIRMAN MELIUS: Thank you. And we'll be hearing from Dr. Ziemer and the Work Group either tomorrow or on Friday, and they'll update us on that.

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1	Anybody else on the phone? Okay.
2	If not, thank you all for coming tonight and
3	for your comments. And we'll be adjourning
4	now, and we'll be reconvening tomorrow
5	morning, I believe at 8:15.
б	(Whereupon, the above-entitled
7	matter went off the record at 6:19 p.m.)
8	

NEAL R. GROSS

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