UNITED STATES OF AMERICA CENTERS FOR DISEASE CONTROL

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SAFETY AND HEALTH

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NATIONAL INSTITUTE FOR OCCUPATIONAL

ADVISORY BOARD ON RADIATION AND WORKER HEALTH

92nd MEETING

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TUESDAY
JULY 16, 2013

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The meeting convened at 8:30 a.m., Mountain Daylight Time, in the Shilo Inn, 780 Lindsay Blvd., Idaho Falls, Idaho, James M. Melius, Chairman, presiding.

PRESENT:

JAMES M. MELIUS, Chairman HENRY ANDERSON, Member JOSIE BEACH, Member BRADLEY P. CLAWSON, Member R. WILLIAM FIELD, Member MARK GRIFFON, Member DAVID KOTELCHUCK, Member JAMES E. LOCKEY, Member WANDA I. MUNN, Member JOHN W. POSTON, SR., Member DAVID B. RICHARDSON, Member* GENEVIEVE S. ROESSLER, Member PHILLIP SCHOFIELD, Member LORETTA R. VALERIO, Member PAUL L. ZIEMER, Member* TED KATZ, Designated Federal Official REGISTERED AND/OR PUBLIC COMMENT PARTICIPANTS

ADAMS, NANCY, NIOSH Contractor

BARKER, CHRIS*

BARRIE, TERRIE

BURGOS, ZAIDA, NIOSH

CARROLL, STEPHANIE

CHMELYNSKI, HARRY, SC&A*

CRAWFORD, FRANK (CHRIS), DOL

FITZGERALD, JOE, SC&A

HINNEFELD, STU, DCAS

LEWIS, GREG, DOE

LEWIS, MARK, ATL

LaBONE, Tom, ORAU Team*

LIN, JENNY, HHS

KINMAN, JOSH, DCAS Contractor

MAKHIJANI, ARJUN, SC&A

McFEE, MATT, ORAU Team

NELSON, MARK

NETON, JIM, DCAS

RUTHERFORD, LAVON, DCAS

STEWART, JOAN

STIVER, JOHN, SC&A

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P-R-O-C-E-E-D-I-N-G-S

8:33 a.m.

CHAIRMAN MELIUS: Okay. Welcome to the whatever meeting we are, what is it?

MR. KATZ: 92nd.

CHAIRMAN MELIUS: Number 92 meeting of the Advisory Board on Radiation and Worker Health, and I'll turn it over to Ted for preliminaries.

MR. KATZ: Thank you. Right, welcome everybody, on the line as well. So let me remind folks on the line and we'll try to do this periodically, to please keep your phones on mute, except when you're addressing the group. Press *6 if you don't have a mute button. That will mute your phone, and then pressing *6 again will take your phone off of mute.

The other thing for everybody on the line is please don't, at any point, put the call on hold. But hang up and dial back in if you need to, because hold will disrupt

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the audio for everybody. Thank you.

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So the materials for this meeting are posted on the NIOSH website. That's all the presentation materials and some other background meeting as well, background reading materials as well. They're on the NIOSH website under the Board section, under the Meetings page for today's date.

So you can follow along that way. We're also, for the first time, running this meeting with Live Meeting well, as Live presentations. Meeting for the So the presentations, as they're being shown here, you should be able to watch them on Live Meeting, and that information, to log Live Meeting, is on the agenda, which is on that NIOSH website.

So if you log in there, you should be able to see the presentations as they're shown here, although if you have a problem with Live Meeting, again the presentations are all posted on that website, and you can just

1	pull them up and look at them yourself,
2	changing the pages yourself if you want to.
3	Okay. Public Comment tonight is
4	from 5:00 to 6:00 p.m., and as usual we'll
5	start with people, commenters in the room, and
6	then we'll have folks who want to comment on
7	the line. I have a couple of inquiries
8	already for people who would like to comment
9	from afar and that's great. Happy to have
10	you.
11	Okay. Let's go to roll call, and
12	with roll call, I will as I do roll call,
13	I'll address conflicts of interest that relate
14	to today's sessions. There are not that many.
15	(Roll call.)
16	MR. KATZ: That covers it for roll
17	call, and Jim, it's your meeting.
18	CHAIRMAN MELIUS: Okay, and the
19	first item on our agenda is a NIOSH Program
20	Update.
21	MR. HINNEFELD: Well good morning
22	everyone. Those of you on the phone this is

Stu Hinnefeld, Director for DCAS at NIOSH.

As is custom, I'll give a our short program update. There are a number of I'11 statistics here go through pretty quickly. If you have any questions on those, or any questions on those, or any questions at any time, please just let me know.

In terms of the program news, it occurred to me, as I was putting the slides together, we're in the process of essentially rebidding our dose reconstruction contract. It was originally scheduled to end at the end of April of this year. Some time ago, working with our Programs and Grants Office, we granted a six-month extension to the existing contract, to provide, essentially provide time for an orderly procurement process.

We are at the stage now where the RFP has been on the street for a while, and the proposals from potential, from the bidders were due last Tuesday. So the proposals are in. I am not on the Technical Evaluation

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Board, so the Technical Evaluation Board are the people who know about the responses, you know, who and how many.

work to do. So one or more entities responded to the proposal. So that will be proceeding. We'll see how that procurement, with any luck, will go a little smoother than last time, when we tried to rebid the contract and it was a very, very difficult procurement process involving a number of short term extensions. We're hoping we can get this one done in the autumn time frame, and just go ahead and make the award, and then move seamlessly into a new contract.

I wanted to mention chronic lymphocytic leukemia a little bit. Everybody knows we've added that as a covered condition, you know, essentially removed the radiation risk factor of zero from chronic lymphocytic leukemia, and by making a regulation change, a rule change some time ago.

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In doing that, we developed a dose model which was a relatively complicated dose model. Rather than having one target organ, you have target lymphocytes, which are distributed throughout the body, with some uncertainty in which organs they are located in.

So you have uncertainty an distribution on the location of your target organ, and you have an uncertainty distribution of the doses apportioned to those organs tissues, where the or lymphocytes might be.

So it's a pretty complicated arithmetic problem to put all those combinations together, and it's being built into our dose reconstruction tools for site by site. So we're still working through that. We made a lot of progress on completing those revised tools and rolling them out.

So many of the chronic, most of the chronic lymphocytic leukemia cases now

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have a way to be done and are being done. But there's still a handful that we're working through the models on, in order to complete all of those. So those are, those cases are not all quite available yet.

We continue to participate in outreach activities. Ι don't talk about outreach activities very much. But participate in a Joint Outreach Task Group with the Department of Energy and Labor, of you know, for this Department also for the and Former Workers program, Monitoring Program, and they are here, by the way today.

The Former Workers Monitoring
Program folks from here in Idaho are here, and
they're hopeful to find some additional former
workers for their program.

We've participated in joint outreach task activities this year in Chicago not long ago for Argonne and Fermi Lab, and somewhat earlier than that, we participated in

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the sort of an ad hoc outreach activity in Attleboro, Massachusetts for the site we know as nuclear, let's see, Metals and Controls, an AWE site up there.

That was really done at the behest of the Congressman from that area, Congressman Kennedy. So we participated in that. In addition, through our outreach contractor, we do dose reconstruction SEC workshops with affected populations, whether they be claimant advocates in Labor, a lot of local Labor officials.

We've done a couple of those in the Los Alamos area for, one for the Los Alamos building trades folks and one for the fire and security services. And then we expect to do a workshop, a longer workshop in Cincinnati, for collection of people from a number of sites toward the end of September.

So those outreach activities continue on as part of our work to the, with the claimant community. Also not on the

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slide, but something I think I probably should talk about briefly, is we've had occasion to examine our conflict and bias policy, conflict or bias policy recently, and our use of it and our behavior with respect to it, and historically our behavior with respect to it, based on some emails that go back a number of years.

Our conflict and bias policy has evolved quite a lot in the last ten years, since I've been on the program. We started with sort of a common understanding, that if you had worked in the radiation safety program at the site, you should not do a dose reconstruction from that site.

So that was essentially the starting, the starting block of this, and whether you should not do it or review the dose reconstruction. So that, you know, I'm conflicted at Fernald. I can't do those dose reconstructions.

Now it's gone beyond that. We've

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been, through experience, have recognized additional considerations that should be added into that process as we go forward. For some time, we identified what we call "key program functions," and decided that not only should you not do dose reconstructions; you should not do any of these key program functions that are for site-specific documents if you're conflicted at a site.

That's things like author of the Site Profile, worked on the SEC Evaluation Report, things of that sort. Then some time later, and I'm thinking this was on the order of three years ago, although I'm going by memory here on dates, and so I may not have the dates exactly right, on advice from -- we always get advice about these things from counsel and the Office of Ethics.

On their advice, we expanded the policy a little farther, to line up more appropriately, I guess in the eyes of the attorneys, with the language in the law and

1	the regulation that cover such things. That
2	gets into language that people like me don't
3	understand, things like a specific, or a
4	particular item with a specific party or a
5	particular what's the language?
6	CHAIRMAN MELIUS: Particular
7	MR. HINNEFELD: Okay. There are
8	general matters and there are specific
9	matters, and there are
10	So language that I can't remember
11	and don't understand.
12	(Off mic comment.)
13	MR. HINNEFELD: Yes. I know what
14	I can't do, and so it's in order to
15	interpret that, then it became a little
16	broader application, in terms of what people
17	are allowed to do, and we've to the point
18	where we are today.
19	So that if someone is to
20	participate even, you know, not author but to
21	participate in this, someone who is employed

by the program needs to be treated

22

like

someone who is not employed by the program.

So if someone wants, you know, if the people working on the Fernald, I'll just keep using myself, although I'm not a terribly good example because I've got an authorization to participate.

did But. if Т not. have an authorization to participate in Fernald, and somebody wanted to get my input about what happened at Fernald, then you know, somebody on my staff, then they would have to interview me, document the interview, just like they would with any other, any other former Fernald employee who's not employed by the project. So there's been this evolution of items could be done.

So that's also then affected how we've behaved. I mean many years ago, quite a long time ago, I attended some Fernald Work Group meetings. But then we adopted this latest policy that said you shouldn't participate if you have a conflict.

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Not only should you not have, you wasn't performing a know, I key program function, but I would go to the Fernald Work Then when the latest revision Group meetings. of policy comes out that says you shouldn't participate, well then I didn't. I stopped participating in the Fernald Work meetings, until I got authorization from the Office of Ethics to again participate.

So during this whole evolution, and because of the evolution, there were, specific because of the nature of the requirements, and because of some things some of our folks said in emails from a number of that maybe -- and the years ago, actually predates the effective date of the latest policy, but was sort of during the rolling implementation of the latest policy.

We needed to make sure that everybody was aware of what does this really mean, what does this policy mean in terms that we can understand? So we've put out a message

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to all the staff, saying this is what's expected.

Now naturally when you do that, you get questions. People think of well what about this situation and what about this situation, which you know, frankly I didn't think about when I wrote the message originally.

So we're working with the Office of General Counsel to arrange some answers and probably a briefing, a give and take sort of question and answer briefing period for the people that are affected, to make sure that we're implementing this correctly.

Made sure everybody knew that this is a company policy or an Institute policy, and it's to be complied with. Just like any other policy, if you don't comply with policies, you're subject to discipline. So that was part of the message we sent.

So that's the message that we've taken from that, and I think periodically we

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probably just need to remind ourselves about what this policy says and how we're supposed to behave, in accordance with it. So I'll be glad to answer any questions about that or any of the other news items I've got here.

CHAIRMAN MELIUS: Questions for Stu?

(No response.)

CHAIRMAN MELIUS: If not, I'll -I have some. First of all, just on the topic
you just brought up, and I'm not sure if
everybody on the Board is aware of the latest
round of emails. But I only get Stu, it's a
question of, you know, the timing of when
policies went in place.

There is one email there from an individual who says that states that he has, knows he has a conflict of interest, but still feels obligated to participate and provide information and a recommendation on the site that he has conflict on.

So I mean I don't think it's a

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1	question of when that was put in place; it was
2	a question of someone thinking that they were
3	not bound by any conflict of interest
4	policies.
5	I think that's a very serious
6	problem, and needs to be addressed.
7	MR. HINNEFELD: Well that's
8	certainly the intent of the message. The all-
9	DCAS message that I sent was that hey, this is
10	the policy and you're subject to discipline if
11	you don't follow it.
12	CHAIRMAN MELIUS: But you know
13	obviously, in retrospect, it should have been
14	dealt with at the time, because it wasn't a,
15	you know, an email just to one individual.
16	Others were aware of it.
17	Secondly, I think with these
18	emails, there's certainly a person who
19	information relevant to a Class Definition at
20	the site, which is the Mound Site, was not
21	brought forward, information that was
22	pertinent to that, until it was finally

revealed some months later and was brought to attention.

understanding from talking to Stu that once it was brought to his attention, then there was follow-up. But there was a period of months when information that essentially would have and did significantly change the Class Definition, was known to people in the program; it was not dealt with. That delayed, at least to some extent, maybe a month or two, maybe longer, the action.

I think that's also a serious problem. There were clearly people that objected to the -- didn't like the Class Definition, didn't like what was being done with that at that particular site, and were, you know, again taking steps to try to undermine that.

I think that seriously hurts the integrity of the program, and I'm hoping that will be addressed also. Now that individual

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is no longer involved in the program, so it's not someone we're currently dealing with. But I think it's a pretty serious situation, at least as reflected in those emails, which again may not be complete. There may be other information. But it certainly appears.

I think we'll be hearing from the petitioner involved in the public comment period, who also has views on this.

MR. HINNEFELD: Yes. I understand that, and I can tell you that we are trying our best to be diligent and make sure we see all the evidence. Now if it's not, you know, say in making sure that people are aware of their obligation to do that. They do that through performance, a review and performance intervention.

CHAIRMAN MELIUS: Yes, and I would add that in that same set of emails, the person, one of the people involved also made some disparaging remarks about Board Members who were involved in this, because they, this

Board Member disagreed with them, 1 2 asking questions, which is also, I think, not 3 appropriate. 4 MR. HINNEFELD: That's not appropriate. 5 6 CHAIRMAN MELIUS: The disagreement 7 can be, it may be appropriate, but expressing it and using that as a reason for taking some 8 of these actions, I think, also doesn't speak 9 10 well. Yes, and that was 11 MR. HINNEFELD: that's before in email 12 actually, come up 13 exchanges, and attitudes toward the Board and the quality of the research we present to the 14 15 Board. 16 CHAIRMAN MELIUS: Yes, yes. 17 MR. HINNEFELD: I can say that it's something we're aware of, and we're 18 19 attempting to address. You know the three of 20 us, four of us in the room, counting Josh from I think you know that we're Cincinnati. 21 trying to pursue and make sure we have a basis

for the things we put in front of the Board.

CHAIRMAN MELIUS: Yes, and appreciate that. But again, to make everyone aware. One other subject on budget. update Board is with I'11 the on the sequester, Stu and I had a telephone call, once Stu figured out sort of how much money they had. As you remember, the ORAU contract hit particularly hard was or disproportionately because of the, just nature of what could be done.

At that time, it was the, sort of, I won't say the target. It wasn't targeted, but it ended up it's basically bearing the brunt of the sequester, and Stu worked with them and he and I had a conversation, Ted was involved also, to talk about making sure that the -- we were in tune on terms of what priorities would be for handling that.

We prioritized obviously towards the outstanding Special Exposure Cohorts that needed to be actioned on and stuff that could

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be done in, you know, a reasonable time period and so forth, affected by the sequester, obviously keeping the dose reconstruction and other activities going.

I think that's worked out as well, though not to say that the sequester didn't have an impact on what could be done and certainly could slow down resolution on a number of items. I think we've had to, and we may continue to have to put off or delay, at least not get certain things done, Site Profile reviews and so forth, as quickly as we would may like to, because of budget issues.

The budget for next year is still up in the air, and we won't know in, I think until -- well, hopefully we'll know before October 1st, but we'll see.

MR. HINNEFELD: I'll give you 2 to 1 odds that we'll start on a continuing resolution. I'll give you 2 to 1 at least on that.

CHAIRMAN MELIUS: Yes, yes.

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MR. HINNEFELD: Yes, thanks Jim. I neglected to bring up the money issue. There's essentially no good budget news, but if you want to look for a silver lining, ORAU had to absorb a year's worth of sequestration cuts in six months, and this coming year, they'll be able to plan the cuts over 12 months. So it's an easier per month adjustment maybe.

You know, you never know what's going to happen in the future. We may lose more money. So that's it, plus we've saved money in other areas. The travel, the budgeted travel expenditures are quite a lot down thanks largely to Ted and the use of Live Meeting. We've had some attrition, and so our PS&B is down noticeably from last year.

So there's chunks of money that are programmatic money, and we're working with FMO to see exactly what we can get on the ORAU contract, and what we'll have. Of course, next year we just expect, everybody expects a continuing resolution, because the three

budgets, the President's, the House and the Senate are just not even remotely similar. So everybody expects a continuing resolution for next year.

CHAIRMAN MELIUS: And I would just add for Board Members, I think it's important that for, you know, Work Groups to keep some of the budget constraints in mind when you're, you know, assigning work, because whenever we look into something in a Work Group, it usually means work for SC&A, which is not been as affected this time.

But we don't know going forward, and also for, you know, NIOSH has to respond, or are you asking NIOSH to elaborate on something or whatever. I think again, I think everyone's doing fine on that. But do keep it in mind and, you know, at least if you have six things that need to be followed up on from say a Site Profile review, try to prioritize those, so that at least the more important ones, the ones that may have the most impact,

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will get done first.

That's not always easy to judge until you've done it, but I think we all need to do the best we can on that.

Finally, I have a question on, which actually came up after last meeting. But what is the notification now for -- how does word get out about these Board meetings? Because some people last time had felt that they, in the Augusta meeting, that they had not heard about it ahead of time.

MR. HINNEFELD: I checked on that.

CHAIRMAN MELIUS: Yes.

MR. HINNEFELD: The notification goes to claimants within a geographical area. I want to say it's 100 miles, 50 miles, something like that, who have active claims. In other words, the claim has -- they've submitted a claim, and it has not been sent back with the final dose reconstruction. So that's the notification list.

And you know, we've not made any

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1	adjustments to that for this meeting. I don't
2	know, you know, where else to go with the
3	notification.
4	CHAIRMAN MELIUS: But my
5	recollection, we used to do outreach to the
6	local newspapers
7	MR. HINNEFELD: Oh, I think we do
8	send those.
9	CHAIRMAN MELIUS: To the local
10	unions, the programs and so forth, the
11	screening programs.
12	MR. HINNEFELD: I don't recall if
13	we've contacted unions in the past. I believe
14	we still send a notice to the newspaper.
15	Whether they run it or not, I think, is a
16	newspaper's option.
17	(Off mic comment.)
18	CHAIRMAN MELIUS: Okay, because
19	for some reason, something slipped up and I
20	guess I don't know what happened.
21	(Simultaneous speaking.)
22	MR. HINNEFELD: there was a

1	particular guy that got up at Savannah River
2	and said hey, I didn't know. I should have
3	known. Well, he was not an authorized rep for
4	any active claims, and so he didn't get the
5	notice.
6	CHAIRMAN MELIUS: But he was a
7	petitioner, I thought. I thought that was
8	MR. HINNEFELD: No. The one who
9	complained about not being noticed is
10	essentially wants to function as an
11	administrative rep, or is an administrative
12	rep. But he has not been an administrative
13	he was not a rep for anyone with an active
14	claim.
15	CHAIRMAN MELIUS: Yes. We're
16	thinking different people then.
17	MR. HINNEFELD: Okay.
18	CHAIRMAN MELIUS: I'm talking
19	about a petition, petitioner.
20	MR. HINNEFELD: Okay. The
21	petitioner didn't know. I would have thought
22	the petitioner would have been told.

CHAIRMAN MELIUS: Yes. Well, I
don't know. But anyway, just make sure, and I
think that's critically important, because
there are delays now in setting up the
meetings and so forth, and we need to make
sure that word gets out as timely as you can,
given some of those constraints.
MR. HINNEFELD: Right.
CHAIRMAN MELIUS: Any other
questions, comments for
MR. HINNEFELD: Are there any
questions on any of the statistics? I didn't
run through those, but there's no real need if
you've got them all.
CHAIRMAN MELIUS: Yes.
MR. HINNEFELD: It's the same
MR. KATZ: Oh, and for Board
Members on the line, just Zaida has muted all
the lines. So you'll have to press *6 to come
off mute, in case you don't know that.

(Off mic discussion.)

1	CHAIRMAN MELIUS: Okay. Yes,
2	Brad.
3	MEMBER CLAWSON: Who is handling
4	the media outreach and stuff for NIOSH?
5	MR. HINNEFELD: Josh, Josh Kinman.
6	MEMBER CLAWSON: So like for this
7	meeting, where did it go?
8	MR. KINMAN: Every local TV
9	station
LO	MR. HINNEFELD: Josh, Josh. Can
L1	you speak into the mic?
L2	MR. KINMAN: I'm sorry. So the
L3	notices will go out to all of the, any media.
L4	I pull everything up that I can find within
L5	the areas. I've been finding that most of the
L6	time, there is very little media interest in
L7	Board meetings.
L8	They're shared with any outreach,
L9	and as far as the petitioner, we document all
20	of our interactions with petitioners, notify
21	all of them, and if I'll look into what
22	happened at Savannah River and find out if

1	that was the case and why. But generally, as
2	far as media, there's very little interest.
3	CHAIRMAN MELIUS: Okay, thank you.
4	Anybody else?
5	(No response.)
6	CHAIRMAN MELIUS: Okay. Thank
7	you, Stu, and now we'll hear from the
8	Department of Labor, which is
9	(Pause.)
10	CHAIRMAN MELIUS: Welcome, Chris.
11	MR. CRAWFORD: Good morning.
12	CHAIRMAN MELIUS: Tell Jeff we
13	miss him, but
14	MR. CRAWFORD: As it says on the
15	slide, my name is Frank Crawford, and I'm
16	delivering the DOL presentation today, in lieu
17	of Jeff Kotsch, who couldn't be here. These
18	are very tiny arrows, so okay.
19	It's quite a large slide
20	presentation, so I'm going to have to skip a
21	lot of the detail. I'm told that it will be
22	on the DCAS NIOSH site, for people who aren't

here. I won't go over the enactment of the EEOICPA, which I think by now everyone here is familiar with.

The case statistics, just one They're looked at in several little caveat. different ways, and some of the Part В statistics you will see will be only radiation-related cases, that is, cases handled by DCAS essentially, whereas some of our other statistics will be based on all Part cases, which includes chronic beryllium disease and silicosis. So if you see some obvious number discrepancies, that's part of what we're seeing here.

So to date, apparently we've had 163,912 cases filed, and we paid out over 9-1/2 billion dollars in total compensation. That's for the entire EEOICPA program.

We had 40,108 cases referred to NIOSH for dose reconstruction, 37,917 cases were returned by NIOSH, 32,000 with dose reconstruction and about 5,800 without a dose

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reconstruction. On the latter, probably many of those were recalled for SEC processing. But there are other reasons too sometimes a dose reconstruction doesn't get done. There are 2,191 cases currently at NIOSH, by our count.

Of the 32,000 cases returned with a dose reconstruction, we see that 26,000 of those have a dose reconstruction and a final decision, of which 9,300 were approvals and 16,800 were denials. Now these, I believe, are radiation-related cases, because of the DR coming back. The next slide has a different view.

of the Part B cases filed we see in this colorful pie chart, NIOSH really is only handling about 34 percent of the normal radiation-related cases. That other category is going to be primarily beryllium disease and silicosis. It's a pretty large category. I hadn't been aware myself of how big a part of the program that is.

Then we have SEC cases. Some are

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referred to NIOSH and some are not, and then some smaller part of RECA cases, 10 percent.

Of the Part B cancer cases with a final decision to accept, we have with over 12,000 payees accepted DR cases, and 1.28 billion paid in compensation. Of the accepted SEC cases, we have 19,363 cases, with 32,000 2.89 billion payees and compensation. So the SEC cases quite outnumber the DR cases.

Cases accepted based on the SEC status and having a PoC of greater than 50 percent, that is with a dose reconstruction done, 633 cases in that category, with 770 payees and about 95 million in paid compensation. The totals of all accepted SEC and DR cases, 28,619, with 45,000 payees and 4.26 billion in compensation paid out.

This is special for Ms. Munn. This time we went back to the top four work sites for the quarter. These are Part B EEOICPA cases, and we'll see. As Stu mentioned

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earlier, the only surprise on this list is one of our old favorites, Hanford, Savannah River and Los Alamos, with the additional of Metals and Controls.

Jeff and I asked about this, and it turns out that Representative Kennedy from Massachusetts had a town hall meeting specifically for Metals and Controls, which generated a lot of cases all at once. So that's why the quarterly ranking is so high.

The EEOICPA Part B cases, final decisions. We have 51 percent approved and 49 percent denied. These would include, Ι believe, the SEC cases as well. We have now a bar chart with percentage of new cases for DOE versus AWE sites, and we see there's -- if anything, there's a trend of more AWE cases in recent years, although still a great majority of the cases are DOE sites. Metals and Controls would be an example of the AWE sites.

DOL also participates in the Joint
Outreach Task Group, and these include town

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hall meetings and traveling resource centers. In the cases of small SECs, press releases are issued. From what we just heard Josh Kinman say, we don't know where they go from there.

Also, the outreach people have been hosting informational meetings regarding medical benefits, which is quite a complicated subject, provided under the EEOICPA. I think we've all seen this. The Joint Outreach Task Group has many departments and Members.

all monthly They're conference calls, and there's a JOTG meeting scheduled for September, tentatively. There's no chance of reading all of these meetings, but I will mention that this is for fiscal year 2013. That is from October 1st, 2012 through September 30th, 2013.

We've had SEC meetings at Hanford and Clarksville. We've had SEC and medical benefits meetings at Oak Ridge X-10. Also at Fermi National Accelerator Laboratory and Argonne National Laboratory.

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At Los Alamos, we also had an SEC and medical benefits meeting. We're up into February 20th now. Then in Knoxville, which was a little bit enigmatic, there was medical benefits meetings for physicians and health care providers. Presumably, this is for people who are living in the Knoxville area, and need information about medical benefits.

Then in Hanford, it was а hall for claimants and medical benefits just last meeting month. Santa Fe, Albuquerque and Grants, New Mexico had medical benefits meetings in June also, and then Jolingbrook, Illinois -- Bolingbrook, sorry, Illinois, had one also in June. This was a meeting of the Joint Outreach Task Group.

Now this week, it looks like, there was a Portsmouth, Ohio medical benefits roundtable. As you see, claimants' physicians and home health care providers will be attending these meetings.

Now we come to a slide on SEC

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Petition Site discussions, which was the ABRWH's agenda for March 2013. We see that Rocky Flats, Baker Brothers and Pantex are represented here on this chart. I won't go through the individual numbers, there are too many of them.

We can see that progress is being made, I think is the main message here. Also, same topic. We see figures for the Feed Materials Production Center, Idaho National Laboratory and Brookhaven National Laboratory, and here we go.

This slide, Employee Eligibility, just reviews the slightly different provisions under Part B and Part E of the Act, with respect to coverage, and we'll see the next slide, I think, with respect to survivors.

Both parts of the Act, B and E, cover DOE contractors and subcontractors. For DOE federal employees, however, only Part B covers them. They're not covered under Part E. For AWE employees, it's the same. They're

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covered under Part B, but not E. The same for beryllium vendors. RECA, however, employees there are covered under both Parts B and E.

Covered conditions. We see, again we're contrasting Parts B and E here. Chronic beryllium disease is covered under both Part B and Part E. Beryllium sensitivity, which is covered by Part B but only for medical monitoring. Under Part E, it's covered for compensation and health benefits.

Chronic silicosis is covered under both parts of the Act. Cancer under both parts of the Act, and any condition related to toxic exposure, as you might expect, is not covered under Part B but is covered under Part E.

Survivor definitions, just to make our life more complicated, are not the same for Parts B and E. However, there's some overlap. Certainly spouses at the time of death covered under both parts of the Act. Children under age 18, under age 23 if full-

1 time students or any age if medically 2 incapable of self-support, are covered under 3 both parts of the Act. Adult children, however, are covered only under Part B. 4 5 E has no coverage for adult children. 6 Benefits complicated, are 7 particularly for Part E. Under Part B, as we all know, I think by now, \$150,000 benefit can 8

"or the survivor." There's a plus sign here.

Under Part E, the impairment is measured as a

I think this should be

percent, and you get \$2,500 per percentage of

impairment. That's for the employee.

go to the employee.

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Under RECA, there's a \$50,000 benefit under Part B. Under Part E, there is a wage loss of 10 to 15 thousand dollars per year for the employee. There's also \$125,000 survivor benefit for RECA employee survivors. There's also a cap of \$400,000 for Parts B and E benefits combined.

(Off mic comments.)

MR. CRAWFORD: Then are there any

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Ι

1	questions?
2	CHAIRMAN MELIUS: Any questions of
3	DOL? Okay.
4	MR. CRAWFORD: Thank you.
5	CHAIRMAN MELIUS: Thank you. I
6	actually, come back. I have a question.
7	(Laughter.)
8	CHAIRMAN MELIUS: Actually, I have
9	a suggestion actually.
10	MR. CRAWFORD: Yes.
11	CHAIRMAN MELIUS: It seems to me
12	if Congressman Kennedy can get a big, you
13	know, large number of claims by holding a town
14	meeting near one of the AWE sites, that the
15	Joint Outreach Group should also consider
16	doing some of those sites, because a lot of
17	them have a large number of employees that are
18	there, and I think we've always sort of
19	assumed that maybe not many would be living in
20	the area, or they might be hard to reach, due

But the Congressman seems to have

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to the age of groups and so forth.

21

1	found some way of attracting them, and it may
2	be something to consider for some of the other
3	sites. There's a number of them around the
4	Boston area, I recall, Ohio and so forth we've
5	looked at that are, have numbers.
6	I think you did one for did
7	they do one for Cincinnati, General Electric?
8	I can't remember.
9	MR. HINNEFELD: We did a GE-
10	specific I think SEC meeting.
11	CHAIRMAN MELIUS: Yes.
12	MR. HINNEFELD: Just from that,
13	the Joint Outreach Task Group at DOL's urging
14	actually is considering a meeting in exactly
15	that area, the Massachusetts-Connecticut area.
16	I don't know if they're that's what they're
17	trying to decide, because there are a lot of
18	AWEs. There are also a lot of AWEs in the
19	Niagara Frontier in Western New York.
20	So that might be another place
21	they want to go. But this whole Metals and
22	Controls thing, I think, kind of brought home

1	to them that there are these sites up there.
2	Not only that one, but also Connecticut Area
3	Aircraft Nuclear Engineering Laboratory,
4	CAANEL.
5	CHAIRMAN MELIUS: Yes.
6	MR. HINNEFELD: You know, I think
7	it's Pratt.
8	CHAIRMAN MELIUS: That was again a
9	large
10	MR. HINNEFELD: Pratt and Whitney.
11	I think that's a Pratt and Whitney
12	CHAIRMAN MELIUS: Yes, Pratt and
13	Whitney is a large
14	MR. HINNEFELD: Yes. And so we've
15	heard some interest through our Worker
16	Outreach contractor from that site, from Pratt
17	and Whitney. So that kind of fits into DOL's
18	plan, to kind of go up into that area. So I
19	think they were planning to do that under the
20	Joint Outreach Task Group.
21	CHAIRMAN MELIUS: You know, like
22	again, Pratt and Whitney has a very active

union and active union retiree group there in Connecticut, and I'm sure some of these other facilities do. So I'm glad they're doing that. It was, you know, I think again, maybe not as many as the larger sites in terms of potential claims, but we've already done a lot of meetings at the larger, bigger sites.

MR. HINNEFELD: Right.

CHAIRMAN MELIUS: So good. Okay, thanks. Next, Department of Energy.

And while MR. LEWIS: Stu's queuing this up, just to clarify, one of the issues with the Joint Outreach Task Group for holding meetings by the AWEs is that Department of Energy, our main interest in the JOTG is for our Former Worker Medical Screening program, and the Former Worker Program does not cover AWE facilities. Ιt only covers Department of Energy.

So that doesn't at all preclude the other groups of having the meetings. But we would not be the driving force behind a

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1	meeting for an AWE. Okay, so
2	CHAIRMAN MELIUS: And Brad, can
3	you hold your questions, please? Let him
4	start.
5	MR. LEWIS: I'll try to address
6	some of them during my presentation, but I'll
7	leave plenty of time for questions.
8	Good morning, everyone. My name
9	is Greg Lewis. I'm with the Office of Health
10	Safety and Security within the Department of
11	Energy.
12	(Pause.)
13	MR. LEWIS: There we go. Okay.
14	So I'm here to talk to you about our role in
15	the EEOICPA Program. As most of you know, our
16	main role is to provide records. We provide
17	records and information to NIOSH and
18	Department of Labor, so they can reconstruct
19	dose and adjudicate claims.
20	We do this in primarily three
21	ways. The first is information related to
22	individual claims. So when someone files a

claim with Department of Labor or needs a dose reconstruction with NIOSH, they'll send a request to the Department of Energy site or sites where the person might have worked and, you know, we'll go scour our records and provide information back to them.

The second is for large-scale records research projects like Site Profile reviews, Special Exposure Cohort research projects, or the Department of Labor The third is to work with Exposure matrix. both Department of Labor and NIOSH to do research on covered facilities.

Our site point of contacts are the individuals out at each DOE site that manage and drive our records research activities. They coordinate with NIOSH, the Advisory Board, Department of Labor and all associated contractors. They set up tours, site visits, worker interviews. They identify subject matter experts on site that might be able to find the right records or answer the questions

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that these researchers have, and they also work with the workers on site on occasion.

Sometimes, you know, they'll point them in the right direction. They'll help them file their claim if need be, or they'll answer questions about, you know, where the records are or what Department of Energy is doing to pull those records. Just for reference, Richard Dickson is our Site Point of Contact for the Idaho National Lab.

So for individual records, 16,000 about records requests per Recently, these are all now going through our Secure Electronic Records SERT system, Transfer System, which I think I've talked about in some previous meetings. But that's really enhanced our ability to send receive requests from Department of Labor and NIOSH. It's also helped us with tracking and, you know, managing our responses.

Many times individuals worked at multiple sites, particularly in the Oak Ridge

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areas or, you know, with certain sites, with the labs in the Nevada Test Site, for example. So we'll often have to go to multiple different locations for one individual. Our records packages can often be hundreds of pages long, and in certain cases, they've been boxes and boxes of information just for one individual.

established You know, we have procedures at each site, and we often check locations. many different Αt site, one there's over 40 different places that someone could go, particularly if someone had a 30-Oftentimes, year career. as contractors changed or as technology changed, there will multiple different databases that be we migrated into.

So they might have to check one database from 1970 to '75, and then a separate database from '75 to '82 or something like that. You might also have to go to microfiche, microfilm, hard copy paper

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records. There's a lot of different places that we might go for records, and we have these search procedures and our Site Point of Contacts and their staff help determine, you know, where the right places to go for each individual are.

So with the large-scale records research projects, obviously those are driven by Department of Labor or NIOSH. So we respond to their inquiries, and try to work to answer their questions or provide them the information that they need.

We also, we review many of these due to classification, but we believe we have procedures in place that allow that to be done, for the most part, without causing any significant delays. I know records that, documents or reports that come through headquarters we turn around very quickly.

On occasion, due to the site's workload or staffing, it can take a little bit for large requests to get reviewed for

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classification out at the sites. But for the most part, you know, we do that without undue delays.

So I put up -- these are some of the sites. This isn't all, but these are some of the sites where we're supporting SEC research, and that's to varying degrees. provided of these we've most of information, and now it's at the point where NIOSH may be coming back for smaller, more targeted questions, and in some of them we're still providing quite a bit of information.

One of the ones I want to talk about up there, which I think I would get questions either way, is the Savannah River Site. I know that we have been really pushing the site to respond a little bit quicker than they have been. We're continuing to work with them. We've approached their management on numerous occasions and their management has been working with us to try to find a way around some of the challenges we've faced.

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In short, most of the issues are caused by some existing, you know, budget cuts and staffing issues out at Savannah River, but are over and above the EEOICPA program. Due to the sequestration and, you know, budget cuts down there, they've had a few different reduction in force, you know, episodes, and they've been down in staff at some of the key locations, particularly the records center, but also in dosimetry.

So we have been working with management. Mr. Podonsky, my boss within HSS, is very involved. We're going to be sending a formal letter down there within the next couple of weeks, as just another effort to try to expedite this.

We do think that at this point we've started to make some progress, just within the last couple of weeks. We believe within the week, a representative from SC&A will be upon the unclassified network within Savannah River Site, and able to do all of the

keyword searches himself.

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So we think that's going to be the key to resolving the request for unclassified keyword searches. We're also moving forward with setting up interviews. I know that SC&A again is working with the site, and they're starting to contact workers and, you know, see who's available for interviews and when they can do them, moving towards a site visit.

We have not been able to complete the classified keyword searches. That's one the things where staffing had significant challenge for the site. last week, they've told they've us identified a staff member to do those searches and, you know, we're following up to see how quickly they can do those searches.

We don't believe it should take too long once they get the staff assigned and on it. But we're going to see, and we also hope this formal letter will give that a nudge to move forward as well. So again, we're

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continuing to work with the site, but it has been a challenge to get that done in, you know, in a timely manner.

I know, I think the Board's expectation has been that it would be moving forward quicker than it has. So we're doing the best we can to get there.

I think I talked about document reviews a little bit earlier. Again at headquarters, the average turnaround time is about eight working days at the sites, and that's also because the headquarters documents are NIOSH reports, which are typically, you know, shorter, whereas the site requests are for source documents, and it can be many source documents, you know, boxes and boxes of records.

So they do take a little bit longer, but for the most part, we believe we've been able to do those in a timely manner.

Then the third function that we

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support is facility research. We update the database of over 300 covered facilities, beryllium vendors, AWEs and Department of Energy sites. The full listing is on the website with the link on your screen there, and for those on the phone, that's going to be in my presentation, page 13.

We are working with DOL, NIOSH and a few, you know, different facilities, trying to refine the years or make sure we have the correct years.

Outreach. Again, I think Chris covered the Joint Outreach Task Group. So I'll move forward, and I think he's, he was correct. I think September and October, the next meeting is tentatively scheduled for the Bay Area, both Livermore and Berkeley, and then we just had a meeting in Chicago about a month ago. We targeted Fermi and Argonne workers.

So the Former Worker Medical Screening Program is a program that my office

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funds and supports through cooperative agreement holders. It's a program that offers a free medical screening to all Department of Energy federal contractor and subcontractor workers from all DOE sites.

provide screenings close to We their location. We have certain cooperative holders that do agreement local programs around the larger DOE sites. But for the smaller DOE sites or for an individual that say has retired to Florida or moved out of the area, we have a supplemental screening program that contracts with local clinics throughout the country to provide these screenings.

The local screening programs for the Idaho National Lab for the are production workers, it's the Worker Health Protection Program, and there's representatives from the WHPP, they're as called, out in the lobby to talk to folks. For construction trade workers, or it's Trades National Medical Screening Building

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1	Program, and the contact information is on my
2	slide here.
3	So I think with that, are there
4	any questions?
5	CHAIRMAN MELIUS: Seeing none
6	oh Brad. What a surprise.
7	MEMBER CLAWSON: Greg, I know that
8	you do a lot of work on this and stuff like
9	that. But you realize what this makes it look
10	like with Savannah River. When we have to
11	have, we've been trying for how long to be
12	able to get in there.
13	My question to you is there
14	anything that the Board can do to help
15	facilitate or assist? Looking at it from the
16	claimant's standpoint, there's all this
17	information out there and we can't get to it.
18	It really, it really puts us in a
19	bad situation. I'm wondering is there
20	something as a Board that we can do to help
21	facilitate this?
22	MR. LEWIS: Well offhand, I would

say from the Board's standpoint as far as research or being accommodating or trying to work with us or work with the site to come up with creative solutions to get the research done, I think the Board has really done everything in their power.

Again, I said the representative from SC&A has gone down there to Savannah River, received training on their networks, general employee training, received a site badge, and is now going to be able to do all of the searches himself from offsite.

So I think in some ways, that will make it easier from now on, you know, for the Board to do this research. But on the other hand, it was a pretty significant hurdle to get this set up. So from that standpoint, I think the Board is doing everything in its power.

I would say, though, being a Presidentially-appointed board, you certainly have some influence and could, you know, let

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us know formally about your expectations. That's certainly an option. I don't want to encourage that one way or another, but that's still, that's certainly something you could do, that would be taken seriously at DOE, both by HSS and by the sites.

You know, but we are continuing to try to make every effort from my office, Health Safety and Security standpoint, to expedite this. So you know, I realize it's been a little bit longer than you would like, than we would like, but we continue to make every effort.

MEMBER CLAWSON: Well Greg, I was wondering if it would help to have a letter from the Board to Savannah River or to DOE headquarters, because really, and I don't mean to put -- this is beyond really a joke anymore, you know. They can come to it or whatever else.

But when we can't even get access to this classified information, it's fallen

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really quite fast. I just -- if there's something that we can do as a Board to assist, I think that it would be very beneficial for all parties. So if there's something we can do, I would appreciate it.

MR. LEWIS: You know from my standpoint, you know, I think all I can say, a letter would, you know, is certainly an option available to you, and I don't think would hurt. Again from my standpoint at HSS, we are already doing everything we can. But, you know, a letter is certainly an option.

MEMBER CLAWSON: And I do realize that you guys are. It just seems like to me sometimes it's falling on deaf ears, and I wonder who the contractor really is working for, because when they're not listening to their own boss, that's kind of hard.

CHAIRMAN MELIUS: I mean my understanding, and I'm not sure Greg's sharing all of this, but there's pretty good high level attention to this within DOE. Glenn is

really working hard to, Podonsky, to get this addressed, and I think we're already seeing some progress. I think we have a way of monitoring it, and I'm not sure being more formal about it at this time is necessarily helpful.

I don't think it would be unhelpful, but I just think that -- I think we're working to get it resolved, and it's certainly something we're keeping an eye on, and I've been informed quite regularly, as this has gone along and so forth. So I think, you know, I think we're making some progress.

I think the unfortunate thing about, you know, it's a big site. We have a wide, a large SEC potential there, a lot of data, which makes -- you know again, means that more information's being requested and so forth. Unfortunately, it's at a time when there are budget cuts going on within the federal government, that are impacting this.

I think what we need to do is make

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sure one, that we're doing this cooperatively, which I think we certainly are and everybody's working together on this, and secondly, that we're, you know, prioritizing appropriately in terms of requests. These aren't just, you know, sort of shotgun requests for tons of information that really isn't relevant.

But I think they are as focused as they can be, you know, given it's a large site and you don't know what's there until you've seen some of it and so forth. I think that's probably the main thing we can do, is make sure that our requests are appropriate.

At the same time, we need to do a thorough and credible job of evaluating the SEC, evaluating that site. So it's not something that can be done easily or quickly

MR. LEWIS: And first, I do want to say that the Board, their contractor and NIOSH have been very accommodating and very reasonable with the request. They've been very targeted and when we've asked some

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questions or said do you really need all of 1 2 this, know, they've you been 3 accommodating, trying it to narrow down, trying to work with us. 4 5 would So Ι say it's been 6 pleasure working with the Board and the 7 contractor. And then another thing, just to reiterate what Dr. Melius said. 8 Podonsky, my boss, is very aware, engaged and 9 10 involved. Pat Worthington and I briefed him last week, and I've been briefing him, and he 11 12 has been applying pressure and will continue 13 to do so, to expedite this. 14 CHAIRMAN MELIUS: Yes, and 15 certainly tell Glenn appreciate his we 16 involvement and effort. Any other questions, 17 comments? 18 (No response.) 19 CHAIRMAN MELIUS: Okay. If not, I'm sure Brad will 20 thank you very much, Greg. have a few more comments and questions before 21

you leave the room.

(Pause.)

CHAIRMAN MELIUS: I hope I need no introduction. State your name, right. This is the Jim and Jim presentation. We've been working with the SEC Evaluation Work Group with NIOSH cooperatively, to try to address the issue of sufficient accuracy, and we're pursuing that. NIOSH has done sort of two background White Papers that were helpful but really didn't sort of get to the core of the matter.

So at the meeting we had a few months ago, we asked NIOSH to draft up an outline of what they thought sort of the key issues were, in terms of approaching sufficient accuracy. That was a three-page, I believe it is outline. It's been in all that voluminous material that we were all sent before this meeting by the shortest, next the agenda.

I don't know if anything else made it under 20 pages or 50 pages or three pages.

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So hopefully you've had a chance to read it. I will sort of go through it a little bit. We're going to continue this discussion after our break this morning, because coworker models are I think one key area of this issue that we're going to have to deal with.

They're coming up. They're becoming more important. They're important for Savannah River Site evaluation; the they're important for Fernald, important for a number of sites we've dealt ongoing fashion, with in and an evaluate those is in some ways tied back to the whole issue of sufficient accuracy.

So I'm just going to go through a few quick slides, mainly summarizing the NIOSH outline. Again, remind all of the us regulation, radiation doses can be constructed with sufficient accuracy, if NIOSH has established sufficient the access to information.

Estimate the maximum radiation

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dose for every type of cancer for which radiation doses are reconstructed, that could have been incurred in plausible circumstances by any member of the Class, or NIOSH has sufficient established, has access to information to do it more precisely than the estimate of the maximum radiation dose.

So this is the regulation we've lived with for quite a number of years now, and continue to try to interpret that and apply it to all of the sites that come up, in terms of making some assessment on, particularly on SEC evaluations. Just to go through briefly through the NIOSH outline, there's a section there, what they refer to as preliminary steps.

But what probably are the most time-consuming, and the most important part of what we do, and probably the most influential is really what are the actual facts about the site. We then have to try to apply our evaluation and NIOSH applies theirs to those

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What information's available, what was done at the site, identifying the exposed populations, trying to understand better about information's available what on those populations. Probably on, my guess is 90 percent of the sites, it's relatively that information straightforward, and really what guides our assessment of, know, whether or not dose reconstruction can be done at that site.

It's time consuming, but it also usually gets us to the answer most readily, in terms of what we're trying to make our evaluation. Then, as it goes through the outline, divided into sort of two types of data.

One is personal monitoring data that is available, at least to some extent on most of the sites and covering a significant proportion of the people that work on some sites, not on others. So when personal

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monitoring data, what NIOSH is trying to do is number one, demonstrate that the highest exposed workers were covered; one can bound the dose.

That everybody at the site was monitored, which is probably a rarity, but not necessarily for external exposures; and/or that the monitor group included those that were, had the highest exposures. Again, the concept of bounding.

So if not everyone was monitored, than it's clearly important to understand who had the highest exposures and that there's information on them, on that. The monitoring method is important in a practical way obviously, and then NIOSH for a number of years applied coworker models for sites where not everyone was monitored.

Again, we'll be spending some more time on that after our break today. But the concept is that the coworker models, they must be inclusive. They need to cover everybody

and be able to be applied to everybody at the site, and second, that they need in some way to account for what's referred to here as stratification, but that there may be people that were monitored in a different way or worked in a different part of the site, or had different, did different tasks at a site or worked in different buildings at a site.

So that that -- that at least has to be evaluated. If a coworker model's going to be used based on personal monitoring or that stratification other data, becomes one of the important things that's looked at. Again, I think the one we most commonly dealt with has been the issue of construction and maintenance workers, and whether they are, essentially have the method of monitoring, as well as the distribution of the results of that monitoring as do the production workers, and that's often been a problem at many sites.

I'll just add that all this is

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also assuming we, in a practical way, that one has a way of placing people within those stratas, if there is stratification, really placing people within however one breaks down the coworker model and uses that. There can be more than one coworker model at a site, depending on where people work, and it can apply to different types of exposures.

In some, it may be very good for it. Typically for external exposures, it's much more data. So the coworker model is much easier to develop and to evaluate, whereas for internal exposures, there's often less data. Methods have changed over time, and it's a much more difficult endeavor to try to deal with that. That's typically where we found the most difficulty, in terms of dealing with Special Exposure Cohort petitions.

Now the outline also identifies some other types of data, and I'll sort of briefly summarize that. But obviously there can be air monitoring, source-term, surrogate

data available. It can be something that has to be evaluated for its sufficient accuracy.

These, at least in a practical sense, are much more problematic, because they usually indicate the personal monitoring wasn't available for that, though in some cases they can be used to fill the holes or gaps in the personal monitoring, or combined in some way, because for certain exposures one has personal and other exposures one doesn't have personal monitoring available.

Basically, what NIOSH's usual approach here is to develop some sort of summary of that monitoring, or develop a model based on that monitoring, that one has to account for the highest exposure; again, it has to be bounding, but at the same time has to be plausible.

For this type of data, that's maybe more difficult than for personal monitoring, because again you're using sort of indirect indicators of exposure under that,

and one doesn't have as comprehensive a set of data. So it can be much more difficult both to bound it, be sure you're bounding, but be sure that you're bounding at the same time, you're not being unrealistically high with that bounding.

Then finally, and one that we probably have not dealt with it much, but it is the question of, you know, what is sufficient accuracy. How much, how accurate do we need to be? To date, I think our main pattern, in terms of how we've evaluated that, is that for, and this has evolved I think over the last several years for the program, is for situations where there's a very low potential for exposure.

I think, you know, the easiest example is the residual periods, where operations have ceased; there's some contamination on the site, but usually it involves a lower level of exposure. I think we've been willing to accept a lesser degree

of accuracy, in terms of the information, because one, we usually don't have as much information; it's usually less personal monitoring for example, or whatever.

But also, the range of exposures, the variability of exposures is probably going to be much less and at a much lower absolute value than would be found during production periods. But we've never really sort of looked at that in any very rigorous way, or tried to set a level.

For those of you that are new to the Board or not even that new, we wrestled with all of these same issues, and the issue of defining sufficient accuracy when we first started the Board, first started the program and when the first set of regulations were developed. We didn't have good answers then, and I'm not sure we have easy answers now.

But it's not -- all this is not a new issue, but it is something that I think is becoming more important as we deal with maybe

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some of the more difficult Special Exposure Cohort evaluations.

Then the other area where I think this is also going to become key is how much accuracy, you know, what is sufficient accuracy with the coworker models, which will be presented a little bit later today, this morning.

It's sort of how do we evaluate, for stratification? How much difference, how much differences there have to be between people in one building or people in different tasks, between construction workers and production workers, in order to say that a single coworker model is adequate and sufficient for that particular group of workers?

Does one need a separate one for construction workers or for people in Building A versus Building B? How we evaluate that and, I think as we'll see, look at percentages, when one starts trying to do that

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at a say more rigorous statistical approach, then what's the test of that? How much accuracy does that statistical test have to meet? What parameters do we have to use for that test?

So Ι think that's where we are now. The Work Group had a short conference call to discuss the outline. We want to, since it's a key issue, we cant to bringing it back to the Board for review and discussion. We'll continue. I think we can maybe talk some more after we've gone through the coworker issue, because that, as I said, I think is one major aspect of that.

Let me give Jim Neton a chance to fill in, if you want to, or make comments.

DR. NETON: I think Dr. Melius did a great job of summarizing what was in the three-page outline that we put together, and I don't know if I have too much more to add, other than I do think that this concept of small low level exposures and sufficiency

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accuracy in one consideration needs to be addressed at some point.

Because we have been behaving that way pretty regularly in the residual contamination period, and that's something that I'd be very interested in discussing further. The other pieces of the document really sort of follow out of what we've been practicing. It's sort of а practical discussion of how we've been behaving all along, using the hierarchical model and that sort of thing. I think that part seems okay to me.

CHAIRMAN MELIUS: Yes, and I think from our discussion in the Work Group with NIOSH, I think what we're probably aiming for is not a new definition of sufficient accuracy or, heaven forbid, a new regulation, but something that would, a set of guidelines like we've agreed on for reviewing SEC evaluation, reviewing sufficient accuracy.

We have them for surrogate data.

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We have them for SEC evaluations in general. Yet these are the parameters you need to look at and evaluate in reaching conclusions. Not strict criteria, because I think there are just many, these sites too are too complicated, the situations are too complicated, that to try to develop criteria, we'd spend lots and lots of time, and we already spend enough time doing this, all this work.

think So but Ι of as set quidelines, so at least we're consistent in the approach that we use, and that we consider what needs to be evaluated as part of doing Again, it wouldn't have specific that. criteria, and I think sometimes people look for those in looking at these guidance.

They're not that. They're guidelines for how to do the evaluation, what should be done. I think we would come up with a, hope to come up with the same for this. But again, we're looking for input from all of the

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Board. Again, it should be something that we're all comfortable with and that we all would find helpful.

So let me open it up for comments or questions. Yes, Gen.

MEMBER ROESSLER: You brought up
- by the way, that was really a very good summary of the outline and everything, and I think we want to keep this in front of us for a reminder. My question has to do with the questions that come up, like how much accuracy is sufficient. That's the basic question.

It seems like to answer a question like that and to keep consistency across the program, we ought to go back and look at those SECs that were denied, or also maybe the ones that were awarded, and try to see what we have done.

Is there a consistent pattern? Are there some things in the past, and you referred to what we've done in the past, and how important that is. But are we planning to

do anything like that?

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CHAIRMAN MELIUS: Well, I think we've already done it in the two White Papers, and those Ι think were, Ι believe Ted distributed them again to people. I think what we found is though they were helpful, they didn't really identify an approach to dealing with sufficient accuracy, and some of that is because they were so dominated by the circumstances at a particular site.

So the practical issues, you know, what monitoring was done, what information was available at the site, what they did at the site and so forth, really drove those decisions much more than -- you might think in retrospect, and I think what may be a way of approaching it, again thorium.

Thorium, you know, was the one that probably led to, you know, a number of SECs early on. But then, you know, as we get into some of the other sites, we found that had sufficient information there to be able to

do dose reconstruction.

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It was just a practical, you know, situation of what available was at particular site, that allowed us to reach those conclusions, rather than any sort sort of calculation principal or any or difference in approach on sufficient accuracy. I think what we, what may be worthwhile doing up with after we've come а guidelines, is then think back do these make sense in terms of our past? Do these capture our past decisions, and do that.

I mean I'll -- when NIOSH first gave us this outline, I sort of skipped over the beginning. I said oh well, we do that all the time. This is sort of the second slide I showed with, you know, what are the practical issues.

But when you think about it, those really are what drives so much of what we've done. It's our evaluation of the information available on the site, and being consistent in

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how we approach that, making sure we don't miss, you know, what happened in different buildings or something like that, that we have as complete information as possible.

But I think we can go back and do that. It just didn't seem to be, I don't think we had the right way of approaching it early on. Maybe we picked the wrong example with thorium. Maybe there are some other examples that would be, would have been better.

But right offhand, I think we all thought that would have been the best example, because we had, you know, SEC evaluations, sort of granted SECs, we had not granted SECs, and we had some tough decisions on those, where the Board wasn't certain what to do. But that appeared to be more due to information available. But that is something we can revisit. Henry?

MEMBER ANDERSON: Yes. It seems to me one of the other things that we've

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struggled with is, and it's sort of wrapped up into variability, and that's kind of the range of exposures, where I think we're comfortable in the residual period that by and large the range of the exposures is quite manageable, where if you have orders of magnitude differences, then you get into choosing a highest variable, and then say well, that will be bounding, that that bound then starts to press the plausibility issues.

So one of the key factors that I think we have to come to grips with, is there a range of exposures where unless we've got lots of data, it suggests there's, it's problematic where we start applying these various statistical things.

it's well, that seems Then too high, so let's just come up with something a little bit lower, and that seems -- and then we get into very much of a subjective decision That, I think, is really process. good description challenge. of So the

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variability in what is an acceptable range of variability, if we're going to apply some of these, I think, is really a critical thing to look at.

CHAIRMAN MELIUS: Yes. No, I agree, and then I mean I think we all have in the back of our mind, even though we may not have a number for health endangerment. We do, you know, think to what extent is this exposure going to affect, you know, dose reconstruction? Is it going to have an impact in that?

If it's a relatively low exposure and the variability is contained, you know, in a practical way by the circumstances, then I think we say well, it's not really going to, you know, affect -- very unlikely to affect individual dose reconstruction involved, and therefore we're comfortable because we --

We're comfortable with both the bounding and we're comfortable that even though maybe we don't have as much

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information, monitoring information, we have enough that we can feel that it's a fair approach, and that NIOSH is again, being a claimant, giving benefit of the doubt to the claimant in terms of doing that, but at the same time coming with an actual, you know, conclusion that we can do dose reconstruction with sufficient accuracy, for that particular exposure.

Yes, and there are circumstances where, on the residual where we have not, where there's construction going on or some other activity on the site, that we know that that range would have been much, could have been much higher. I can't see who -- Phil, yes.

MEMBER SCHOFIELD: The one word that really bothers me at some of the facilities, say like a Rocky Flats or Savannah River or Oak Ridge, is "plausible."

(Off mic comment.)

MR. KATZ: Phil, your mic's off.

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Phil.

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MEMBER SCHOFIELD: Oh, okay. All too often we've had personnel at some of these facilities have been close to the unplausible. So this definition of what is plausible and what is not, at some of more the complicated and the sites where the risk factor is much higher. I think we need to try and narrow that definition a little more stringently.

CHAIRMAN MELIUS: Well, I mean I think that's what we're sort of looking at is plausibility, because a really, is what bound, you know, we can come up with a high number and bound any exposure. I'm sure there's someone who will have а counterexample, but what we've encountered so far at these sites, we can always bound.

But is it a plausible bound, and that's what we really, I think, wrestle with that, and you're right. We need to come up with some better parameters on and consistency on how we do that, at least try to

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capture what we've done so far, so we all have the same understanding and can apply that. Wanda.

of MEMBER MUNN: Some these are certainly well-received. The comments problem, the basic problem that we have is we, in very simplistic terms, trying eliminate all of the technical information that puts layers of complication on what we're looking at, is the fact that we do not have a situation where we have cause and effect of what we're talking about.

We have a situation where we have an influence on the outcome of exposure, but the fact that simple exposure does not translate to either harm or benefit is, leaves us with a situation where we have no clear defining line. Without a clear defining line, where we can say anything below this kind of exposure for this different type of radiation, is not going to be of deleterious effect to anyone.

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But there's a point where anyone would agree that there is going to be harm above that limit. But that limit cannot be defined, despite the fact that we have over 100 years of experience in defining outcomes and recording outcomes. Until we can come to some grips in this body, of what we consider to be the gray area, then it's hard to see how we can address the question of sufficiency.

We probably cannot even agree on where the gray area is, below which there's not likely consequence and above which there certainly will be consequence. That's a very large gray area, and although I really appreciate the work that's being done in recent months, trying to pin this down better.

It turns out to be a statistical ball of tar, and for those who are not really well-versed in statistics, it becomes of definition, which problem although eliminated by our discretions and by the papers, still does not show a clear path on

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how we come to our decisions about the definition of those terms.

I'm at a loss to see how we're going to get past that, in the absence of cause and effect, and in the absence of clear, bright lines from which we can say here's our standard by which we have to make We're dealing in decision. an extremely amorphous area when we're trying to define these terms, and if we're going to insist on defining them clearly, in order for everybody to be in the same boat, then I guess saying what I suspect most people feel. is a thorny issue, and I'm not at all sure that we can resolve it.

I commend your efforts to try to get us to an agreement point. I'll be very interested in seeing how we get there, if we can get there.

CHAIRMAN MELIUS: We'll see. I just would point out that, yes, I guess two things. We wrestled with trying to define

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health endangerment when we first started here, and then for some of the reasons you stated, we decided that, you know, a quantitative definition was not going to be necessarily feasible to do and reach.

But also, I think we have to also remember that the risk, the evaluation of risk that we're feeding information into is the IREP model. That's what Congress set for us, and we're doing what Congress, you know, we're implementing this Act, is what NIOSH is doing and so forth, and they defined what, you know, the level of risk is through the -- and who gets, at least in terms of compensation for these workers, through the IREP models.

So it's what we're feeding into the IREP model is what we're focusing on, and can the exposure part of that be reconstructed with sufficient accuracy, to be then fed into the IREP model. So the risk determination for purposes of compensation are in some ways out of our hands.

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1	It doesn't mean that the bigger
2	questions don't, that you put out, aren't
3	still there. But we in some ways have a more
4	limited role than determining whether or not
5	this level of exposure, presuming we agree on
6	it and so forth, is sufficient in terms of
7	compensation. We have a more prescribed sort
8	of approach to that.
9	For Board Members on the phone,
10	Paul's part of the Work Group. I don't know
11	Paul, if you're still on, if you have comments
12	or David or others? Ted probably has them
13	muted.
14	MEMBER RICHARDSON: Dr. Melius?
15	CHAIRMAN MELIUS: Yes.
16	MEMBER RICHARDSON: Can you hear
17	me?
18	CHAIRMAN MELIUS: Yes, we can.
19	MEMBER RICHARDSON: Okay. I
20	appreciate the discussion. I was, and I agree
21	with what you've characterized with how we've
22	been operating, in terms of dealing with

settings where, for example, residual periods where effective dose we weren't more comfortable I think coming to where it was bound and why.

It's led me to think a little bit about, I think actually the language, sufficient accuracy is probably good if, you know, in the kind of more standard sense of accuracy as being how close is the assigned dose or distribution of assigned dose as to the worker's true dose.

I mean you want something that's accurate, from that sense. So it shouldn't be biased, and if there's also an aspect of precision there, how close are the agreement of the values in the distribution there. So that's, I mean that led me to think we have like a situation of residual period where we can, you have relatively good precision in applying those doses, because the range at distribution is relatively narrow.

You have a good precision and it's

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unbiased with a -- it's sufficiently accurate. But when the bounding gets large, I would imagine you've got poor precision, where the accuracy is low over a wide distribution.

So it might be that we -- I'm not sure if this is helpful or not, but as you start to think about the, you know, pulling the language of sufficient accuracy out into issues of bias and precision, I think that's getting at some of the discussion.

You want assigned values that are close to true values, and we want to be able to do that with little or no bias hopefully, and as the precision of those estimates get, of the distributions we're talking about get wider, the accuracy is less.

I mean I've gone around and around in my head trying to think about what we were talking about. We could try a little bit more to have a, something that ties -- but what I think what we were saying is something that ties the, what you described as the absolute

value of the assigned dose, which is kind of the precision of the range of the values from zero to the bounding dose, giving us a sense of if the precision is good or poor, and therefore whether the accuracy is good or poor.

CHAIRMAN MELIUS: No, I think that is certainly the way I've been thinking about it more, and I think the coworker model issue sort of brings that concept forward, because that's -- because we're trying to deal with that in more statistical approaches, and I think, you know, in terms of precision and bias and which at least us epidemiology people are used to dealing with are, become important in evaluating those.

MEMBER RICHARDSON: Yes, and we have decisions or discussions that turn on both of those, like in a sense are they bounding? I mean are the values, is there good trueness to the assigned values. But we also have discussions about the precision of

the distributions.

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think intuitively, I tend to feel like when we get to places where we're talking about bounding doses which have, you know, are not accurate, you know, in closeness of the assigned dose to the person in a relative sense. But as you said absolute sense, the magnitude of the kind of small, meaning dose is relatively precision is relatively good, and we start to feel comfortable bounding a dose there.

CHAIRMAN MELIUS: Yes, okay. Thank you, Dave. Paul, are you on the line and wish to comment?

(No response.)

CHAIRMAN MELIUS: Okay. Anybody, any other Board Members? Okay, good. So our plan is we'll talk more later and sort of moving forward here. I'll tell you what happened in the Work Group call. We went over the outline and said well, what's the next step? Should we develop the outline to a

1	full, more complete report and
2	MEMBER ZIEMER: Can you hear me?
3	CHAIRMAN MELIUS: Oh yes, now we
4	can. Sorry. Go ahead.
5	MEMBER ZIEMER: Yes. I thought I
6	was off mute, and it didn't seem to work. Yes,
7	this is Ziemer. Most of my comments have
8	already been said in the Work Group, and I
9	kind of agree that we're not really looking
10	for a bright line or a numerical bat or
11	anything like that. We're looking for a
12	process
13	CHAIRMAN MELIUS: We lost you
14	there, Paul.
15	MEMBER ZIEMER: Can you hear me
16	now?
17	CHAIRMAN MELIUS: Yes, we can.
18	MEMBER ZIEMER: Thank you. I
19	don't think we're looking for any kind of a
20	bright line or value, or necessarily even a
21	specific range in every case. It may be very
22	site-specific. But we have to have a process,

I think, where we are able to make the evaluation, have reached sufficient accuracy for this situation, and that's where we need criteria rather than numbers.

Keep in mind, for example, we do a lot of dose reconstructions where precision or let me keep it with accuracy, accuracy is not the issue. The cases where we already passed the 50 percent value with just the external, and we don't have an accurate dose. We have a dose that gives us a point where we can make an accurate decision on compensation.

That often happens, where we have cases where we have already accumulated enough dose to go over the 50 percent mark, without completing the total dose reconstruction. don't necessarily need we accuracy for individual doses. We need to make sure that we have done what is sufficient to get what I call decision would an accurate on compensation.

So we need to keep that in mind,

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whether it's with coworker doses and individual doses, it's the process that makes sure that have the information that's we necessary to make а correct decision compensation. So this is not a straight statistical thing. This is a combination of both the science and the policy.

You know, we already know that in cases where the uncertainty is great, that it tends to favor the claimant. So we need to keep those things in mind as we think about sufficient accuracy.

CHAIRMAN MELIUS: Thanks, Paul.

Jim Lockey.

MEMBER LOCKEY: Jim, when I was listening to David in your review, just for my clarification, if in the residual periods, in moving forward, where we're getting the lower exposure situations. Is it the group's thought that maybe more precision should be, precision in relationship to the maximum dose or the highest range, plausible concentration

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perhaps is more important than precision under that, or is it a combination of both?

CHAIRMAN MELIUS: I think it's a combination of both. It's sort of the range that we're operating in, and the absolute value of that range, of that. I'm not sure we've gotten to a point where we're -- whether we really evaluated that part of it, because I'm not sure how that extends beyond that.

I think we're all comfortable with the residual period approaches given, you know, the circumstances most commonly found there. I'm not sure how we then, are we all comfortable? Are we ready to extend that out? How do we extend it out beyond those ranges?

MEMBER LOCKEY: So if we had, if we felt very comfortable that the bounding dose was very precise, that in itself would not be enough, because we don't have, we may not have enough precision in relationship to the lower exposure situations under that bounding dose, maximum bounding dose?

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1	CHAIRMAN MELIUS: Yes. I think
2	some of how does it apply to that
3	population, because if the bounding dose is
4	very high, the range is high of exposure with
5	the population. You can have a precise
6	bounding dose. It only may apply to how
7	well, is that sufficiently accurate for
8	everybody else in that exposure circumstance,
9	everyone else in that building or whatever.
10	MEMBER LOCKEY: So that it breaks
11	down to job position, job task, and would it
12	fall under that?
13	CHAIRMAN MELIUS: Yes, yes. I
14	think we have an example coming up. Anybody
15	else? Okay. Thank you. We'll, I guess we
16	can talk more about moving forward later after
17	we've done the coworker, and we have some
18	Board work time.
19	We are on our break, and since we
20	are a little bit past, and since, if I told
21	people to come back in 15 minutes they might

not anyway, why don't we plan on what -- we're

not really scheduled. So a 25 minute break. So yes. So come back at quarter of. Is that fair? Okay.

(Whereupon, the above-entitled matter went off the record at 10:22 a.m. and resumed at 10:49 a.m.)

CHAIRMAN MELIUS: Okay. We'll get started again. I think LaVon. No, keep him on his toes. We're going to talk about coworker models, and Jim Neton and Arjun will be speaking, and between now and lunch, and based on, is it OTIB-53? And then there's an SC&A review of that, which everybody received ahead of time and is memorized, so the quiz will be later. So go ahead, Jim.

DR. NETON: Testing. That's good, thanks. Okay. Thank you, Dr. Melius. I thank you for setting the stage with the sufficient accuracy. I think you're absolutely right. A lot of what I'm going to talk about might fit into your, especially with the coworker model, approach that NIOSH has been using for quite

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some time now.

I think we started, TIB-19 was the first procedure or whatever that we put in place back in 2005, and we've had various iterations of coworker documents since then, specifically to address more and more detailed and sophisticated analyses, including censored data, number of data sets and most recently stratification of data in Report 53.

So I'm going to try to get sort of an overview of where we were and where we ended up with 53, to get the conversation going. But I would like to acknowledge that a lot of this work was done by some very talented statisticians that included Tom LaBone, Nancy Chalmers and Daniel Stanescu. Tom and Nancy are with ORAU and Daniel's on our staff.

So just I'm going to have a few slides of background, just for those maybe who are new to the Board, and just to make sure everybody's on the same page. The reasons for

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using coworker data are pretty obvious. The workers were unmonitored, and more significantly, they were potentially exposed.

I mean there are a number of workers that are never monitored, and the potential for exposure can be very small or zero. So we need to keep that in mind, as we move through in this discussion.

for workers that But were monitored and who were potentially exposed, the data could have been either lost destroyed, and this one was envisioned in the Act. Or, as Dr. Melius talked about earlier, monitoring methods were not reliable. The data couldn't measure what they purported to something to that effect, measure, the the complex, neutrons early on in maybe something like that.

Or lastly, available data insufficient to complete a dose reconstruction. You may have a few data points, but a person's career spanned 20-30

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years, and those few data points aren't sufficient, so you need to rely on some other source of information.

We have developed, I think since 2005, at least a dozen coworker models for internal and probably an equal number for external dose at various sites, and the data has come from any of these four different sources.

Preferably, we end up using the cover facility databases that we can obtain, which would include the urinalyses results, the GLD measurements, the film badge measurements, that sort of thing, and those are the best sort of data if they're complete and accurate and have very well identified. We prefer to use those.

But lacking that information, we have used epidemiologic study data that was collected at ORAU for the Center for Epidemiologic Research, or even CEDR, the Comprehensive Epidemiologic Data Resource that

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lists epidemiologic data sets in a deidentified fashion, which is kind of important in this discussion.

But we've used that, and also there's a TIB out there that talks about using claimant data. If we don't have any other sources and we have a fairly robust set of claimant data, and we can demonstrate, at least statistically, that the claimant data not а biased sample of the population, we've even used that in the past. So there's a large number of locations where we can obtain data for these models.

The general approach is to look at the data of the moderate population, and of course we looked at the pedigree of the data, determined the measurements reliable, established if the monitoring population is represented in the work force. That's pretty much the key on this slide, is that we're the highest exposed people monitor, okay. That gives us a good handle.

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Or, at а minimum, we're the representative, a representative sample of the workers monitored. If it's a representative sample that was monitored, then we have a fairly reasonable accurate model. Τf t.he highest exposed workers were monitored and we're applying a coworker model, then we'll probably have a slightly biased high estimate of the workers. We'll talk a little bit more about that later.

The statistical approach for doing the coworker evaluation is well-described in Procedure 95. That basically says you review the data and apply a statistical distribution, which is fit a log-normal distribution. It's been well-established log-normal distributions are applicable to environmental and occupational data, and group the data as appropriate.

Sometimes, when the data are sparse, one has to go to monthly, annual and sometimes up to three years' worth of data to

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fill a gap. Then one generates a summary of statistics, evaluates a fit to the data. It's a pretty straightforward regression analysis that was done early on these procedures, and like I say, those early models that we developed did exactly just that. I'll show you a couple of examples.

As Dr. Melius alluded to earlier though, the external coworker models are much more straightforward to apply to internal. There's a variety of reasons for that. Many, many more people were monitored for external. There's little interpretation required in the internal world, as I'll talk about a little bit later.

The excretion values that you obtain really don't provide information as to what a person's intake was. It has to be converted to some sort of an intake to be meaningful. That's really the trick, and I want to really emphasize that in my discussion here.

This is just an example, I think it's probably too simplistic, but a coworker distribution for external dosimetry of an untransformed data set where, you know, it's pretty clearly log-normal. I've truncated the distribution at 500. It extends way out.

But to get that nice little characteristic log-normal shape, I left it truncated at 500. But you get the idea of what I'm talking about, and many of the data, most of the data we have looks similar to this.

If one transforms, takes a lot of the data and plots it on what we call a Zscore plot, which is a standard normal variate with Z-score of zero being central а а estimate of the data set, and then the values right to the left and the in units essentially standard deviations, one normally obtains a plot that looks similar to this for external data.

Here, we'll have a geometric mean

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of about 130 millirem, and a fairly large GSD. I mean you can see here's six and a half. Interestingly, this is an aside, if you see where that site of administrative dose limit is, that tailing off at the top, this is very frequently seen in external dosimetry results, where, as workers approach the site administrative limit, they start pulling them out of the workforce and fewer and fewer measurements are obtained in that area.

But internal is what I really want to talk about today, because I think it's the most complicated thing, and is really the subject of TIB Report 53. In the internal world, we often have multiple bioassay results per monitoring period. They'll have a routine monitoring program, but sometimes workers are sampled more frequently than others, based on the potential for exposures or even in cases incidents, where there's one will multiple samples.

In that situation, if you're

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trying to model one year's worth of exposure, you can have many bioassay results for several people, and few bioassay results for the remainder of the population. That tends to skew the results, if one uses all of those data.

In fact, the bioassay results from one individual are going to be correlated, because it's the same individual being sampled repeatedly, and that violates the presumption of statistical independence of the data. So given that, I'm going to talk about this a little later.

In Report 53, there's this one person/one sample concept that we've adopted, that heretofore had not been applied in most of the coworker models. As I mentioned earlier, the raw data must be converted to intake and then dose. So you can have 1,000 bioassay results in one year that say .03 picocuries per liter are being excreted on average by this population.

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But what does that mean in terms of dose? What is the intake the population was breathing in? That's critical here, and if you're going to build a coworker model for external, the exposure pattern has to be presumed. If you measure a film badge, you measure a film badge and it represents that exposure period.

Here, what is the exposure potential? Acute, chronic, mixed? In this situation, we talked about this a long time ago. We have defaulted in these coworker models to chronic exposures, and we believe it is a claimant-favorable approach, to assume that the geometric mean of the distribution represents a constant. Everyone would have a constant excretion that was unmonitored at the geometric mean of that distribution.

So this is just a summary of what possible calculations go into a coworker model. In the first box on the upper left, of course you have the urine data, the raw data

that we obtain from some database. In the second box we have the, what we call the one person/one sample urine data.

In 53, we've adopted an approach where if you're modeling one year, you will take the average value of the person's urinary excretion in that one-year increment, and use that as one of the data points in the coworker distribution. In our opinion, that actually is more reflective of intake than any other method you could use, such as using all of the data by itself in that one-year increment.

If you think about it, it's almost like -- you take the average value of the person's urinary excretion in that one-year period, and multiply it times a day, the monitoring period, you have picocurie per liter days. An integrated estimate of that person's exposure over that one period. It just makes perfect sense to us.

The third box on the lower left shows that we will take, using the one

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person/one sample urine data, the 50th and 84th percentile of the data. That's generating a log-normal distribution plot, and then the subsequent in Box 4. You take the 50th and 84th percentile intake rates calculate geometric and geometric mean standard deviations.

That's where the rubber meets the road. That's where you're converting a bioassay excretion value, an average bioassay, a 50th percentile bioassay excretion value into some chronic intake over a period of time. Of course, and in Box 2, 3 and 4 is where we can actually look into the, peer into the inner workings of the models, and see how they behave.

We can't do anything in Box 5, which is person-specific intakes and doses. That gets into some very -- that was coworker model by cancer type and all of that sort of thing. It's just not possible. Of course, the final outcome is Probability of Causation.

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As Dr. Ziemer mentioned earlier, the real question here is do we have a model that provides an accurate compensation decision for a worker?

Okay. This is just an example of the probability of distribution for a single year for urinary excretion. Here you have the standard normal quantiles on the X axis. My geometric mean line's a little off; I wasn't perfect in lining it up.

But you can get the idea that the geometric mean of this distribution is .7, with a geometric standard deviation of 4. Fairly large, but this is fairly typical. In this case, N was 332 bioassay measurements. Little N was 196 uncensored, one person/one sample uncensored bioassay measurements.

So you get the feel here, and I want you to remember this graph, because I'm going to refer back to it a few times. But this is the distribution that one would see in the monitored population. You can see that on

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the Y axis, the quantities vary from .2 up to basically 20. So you've got a couple of orders of magnitude variability in the distribution here.

So just to refresh, this is a distribution for a single year. This is what we would calculate for one year, and this is what would go into the intake model. But on the next slide I want to point out to you, there's a number of points here. I think there's 14 points here representing different years.

So if the previous slide represented one year, where am I going here, wrong way. If the previous slide represented distribution for one year, that is represented by say the first dot right here. That would be the geometric mean of that distribution on this curve.

Now we would take subsequent years of data and fit, and plot them here as well, and then fit the best intake retention curve

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we can. It's essentially a linear square regression analysis through these data points, and this is where the rubber meets the road. This is the intake that's going to be assigned for, to this cohort, these unmonitored workers over a period of 14 different years.

So if there's a difference in any one of these little points, it may not make a practical significant difference in this curve that's fit, because if you see the values here, the curve predicts that the chronic intake is something like .96 DPM per day, with a standard deviation of .22 DPM per day, a fairly substantial error.

This is above and beyond the uncertainty associated with the individual coworker models. That needs to be kept in mind. So what we would do here is we fit this 50th percentile distribution, and then re-run calculation using the 84th percentile distribution to establish the GSD of intake for this entire 14-year period.

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So this is where Report 53 talks about what's the practical significance of any differences in the individual annual coworker models. This needs to be kept in mind.

So the application of the coworker model, as I mentioned earlier, is based on the potential that the monitor worker -- with the potential, the exposure potential for unmonitored worker that we're trying reconstruct. The person would receive either distribution, i.e. the full the 50th percentile with the geometric standard deviation as the input parameter, or the 95th percentile of the distribution.

So for this distribution, would either, when it's converted to intake of all and dose, course, but it's proportional, it's not that the person gets just the 50th percentile. The entire distribution is input into IREP as well, and that is sampled as representative of our best of the worker's intake for estimate that

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So it's not accurate to say that they're just getting the 50th percentile. We take advantage, well we acknowledge the uncertainty in the data itself, set and incorporate that distribution. So that's an important point to remember.

So each situation is evaluated on a case-by-case basis, and I don't want to get into the judgment that's used there. I talked about that in the past, whether you have administrative workers versus clerical versus chemical operators, that sort of thing, and that's taken into consideration.

But there is the issue that Dr. Melius mentioned of potential stratification of the data. That's where Report 53 has been issued, and it's our attempt to statistically, provide a statistical framework, which one can for analyze data sets potential I mentioned earlier stratification. Report 53 introduces the concept οf

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person/one sample, OPOS as we call it, and I've got a slide here that basically summarizes what that means.

I mentioned earlier, minimizes issues related to correlation of data. It minimizes issues related to one person driving the distribution. I think there's one data set we have. There could be 50 samples from one person and then 100 samples from another person. That person's samples would drive the distribution, totally not appropriate.

So to use the data, the concept of the maximum possible mean was developed, which is oftentimes you have a combination of censored and uncensored data. Censored data of course just meaning that the data report is below some limit of detection.

So I have three examples here to sort of point out how this one person/one sample would be calculated, given different scenarios. In the first example, you have four data points of 10, 3, 5 and 6. We would

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simply just take the mean of those data points and they would be put into the distribution as 6, the average value of those four.

In the second example, if you have the same data points but the 3 and 5 were censored, the report is less than values, you would still report the data as 6. That's where the concept of the maximum possible mean is. It probably wasn't a 3 or a 5, but it certainly is no higher than a 3 or a 5. So we're just going to assume, for claimant-favorableness, that it was that.

In a third example, if you have all censored data, it would -- the average is still 6, but it would be reported as a censored data point, using in the database as a censored datapoint as less than six. That's a pretty simple statistical calculation, but that's very important as to how we treat the data.

So to get to the issue of stratification, the monitored population is

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really, has got to be a conglomerate of a number of subgroups. You know, you take an entire 500 point data set. There's going to be different subgroups in there.

As I mentioned earlier though, the single distribution can be applied if the highest exposed workers were monitored, or workers were sampled representatively, representative workers were sampled of the workforce.

You took -- it wasn't biased in some particular means, such as only the lowest exposed workers were monitored or something of If you do, however, suspect that nature. stratification, it be statistically can evaluated, and Report 53 introduces the concept of the Monte Carlo permutation test and the Peto-Prentice test, and I'm just going to briefly describe those today.

The Monte Carlo permutation test has some assumptions that the data can be described by a log-normal distribution, which

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we know to be the case, and that the data is not heavily censored. No more than 30 percent of the data should be censored.

believe that it do is an important criterion that the data, if you're going to stratify data, it has to be based on some a priori criterion. You can't just go looking for differences data mining, "aha, I find a difference saying because statistically, if you do enough calculations, you're going to start finding differences that really aren't necessarily based in any reality.

The a priori criteria could be as simple as, and this shows up Savannah River construction workers versus non-construction workers at Hanford Area 100 versus 200 Area workers. But we feel very strongly that one needs to have an a priori criterion before you start investigating stratification. You have to have some reason to believe why they're different.

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For each of these strata, you calculate the geometric mean and the geometric standard deviation. So you have a delta. You have a difference between the geometric means and the geometric standard deviations of the two strata, okay. Stick with me here. It gets a little more complicated.

You calculate, as I mentioned, you calculate the difference between the two, and these differences comprise one data point, with an X-Y coordinate. You're going to have the geometric mean on one side, geometric standard deviation on the other. You plot a single point.

Now what you do is you take the entire data set and you randomly pull distributions out of that data set. Say you had 300 samples. 200 were from one strata and 100 were from another. You would sample 100 random points out of that data set, calculate a geometric mean of standard deviation, and plot it.

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Then you would take, what did I say 200? You take the 100 data points and do the same thing, and you keep resampling this distribution 10,000 times, until you generate a plot of the possible distribution of all of the differences between the geometric means, the standard deviations within that one data set.

That gives you sort of the universe of possible issues, and what you have here is one of these plots that has a 95 percentile confidence envelope. It's an ellipse, and the line drawn around the points there is where 95 percent of the data fall.

Τf the difference that you calculated between the two strata in the first place falls within that ellipse, as shown here, at the 95 percent -- falls within the 95 percent confidence band, then concluded that the two strata that you attempted to evaluate are not statistically different.

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I just have an example here of one where you do the same calculation. This difference over here, this point on the right, is way outside the ellipse. Therefore, it would be concluded that the strata are statistically different.

Sounds complex, but it's fairly easy to perform on computers to get these results, and it's visually, it's pretty visual too. I mean it gives you a nice feel for how the, where the data are going.

Now the benefits is that you can easily compare whether the different strata are different obviously. But the limitations does require some a priori decision on the distribution of the data points. Here, we assume they were log-normal, which is pretty reasonable.

It doesn't work, though, if the data are heavily censored. You end up getting too many random draws of zero. Essentially, you can't -- the censored data is going to

come up zero, and they just can't plot the points.

In this case, our statisticians have researched, and it has been determined that the Peto-Prentice test is the most powerful test that can be used to compare two strata in this situation.

The Peto-Prentice is really a sophisticated version of a rank order Wilcoxon rank order test that we're all familiar with, the range values. I don't want to belittle it. It's much more sophisticated than I'm going to present here, but essentially the same thing as a Wilcoxon rank sum test.

It's a non-parametric test. In other words, priori distributions no а You merely rank the data points in assumed. the distribution. It can handle censored data. It's built to do that, and you can compare whether the strata are different at some p-value. Here we chose the .05 level of significance.

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We've done some testing and for cases where both the Monte Carlo and Peto-Prentice are applicable, they typically lead to the same conclusion. So it seems to be a reasonable test. So just to give you an idea, let me go back to this graph here.

This is the distribution of all the samples. So in the Peto-Prentice test, what one must do is you rank the samples in this order. It's a cumulative probability plot. It's a survival curve basically. But in this case, it will be a cumulative probability plot, and you take adjacent data points on the curve, add them together and subtract one, and you get a value.

So you take the first data point, add it to the second data point, subtract one, get a value, and you do that all the way through the distribution. So you get a series of values. You also, though, keep track of which data points came from which data set. So if I generate all my series of values here, if

I had more data in the high values from one strata, you can imagine you get a much higher number, because the higher values are appearing from that strata in the higher end of the distribution.

So it gives you a way of looking at are the data grouping in some particular manner in these distribution of samples. It's kind of very nice, interesting test. Okay. So moving on with the graph, here are two graphs of samples that we've tested using the Peto-Prentice test, and on the left you have the data were combined into a single data set, as we've talked about earlier, evaluated, and in this particular instance, in the Peto-Prentice test we concluded data were not significant.

The p-value I think was -- you can't read it very well, but it's like .17 I believe, and the data are -- you can say that they look different, but they're not statistically different by this test. In a second set, the data points, the data are much

1	more further apart, and as concluded here, the
2	data are significantly different at the .05
3	level. P-value is very, very, very small.
4	So that's used for when you have a
5	very highly censored data set, as opposed to
6	the Monte Carlo permutation test.
7	So in summary, I just point out
8	some of the obvious things I've been talking
9	about here, is that we believe that coworker
10	models can be used to reconstruct doses. But
11	one needs to be mindful of why the workers
12	that are being reconstructed weren't monitored
13	in the first place.
14	I mean you really have to come up,
15	come to grips with that scenario. I mean, you
16	know, if construction workers are different,
17	yes and they're a little higher, but then what
18	were the exposure potentials for the non-
19	monitored workers in that construction, in
20	that construction group?
21	You've got to be careful and

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applicability

1	representativeness, of course, quality. We do
2	need to be mindful of stratification. We also
3	believe this one person/one sample approach is
4	the way to analyze these data sets. It really
5	makes a lot of sense to us, and the
6	stratification can be evaluated as we propose,
7	using these standard Monte Carlo permutation
8	tests or the Peto-Prentice test.
9	I think that's all I have to say.
10	I'd be happy to try to answer any questions.
11	If not, I think I have some crack
12	statisticians on the phone, phone a friend as
13	they say.
14	CHAIRMAN MELIUS: We'll keep the
15	phone muted as a real test. But Jim, you do
16	get the prize for the best graphics so far.
17	DR. NETON: Oh, thank you.
18	CHAIRMAN MELIUS: The Monte Carlo
19	permutation and LaVon, you'd better get to
20	work, come up with something here.
21	(Off mic comment.)
22	CHAIRMAN MELIUS: So questions for

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that?

MEMBER KOTELCHUCK: Dave Kotelchuck. Your Monte Carlo test uses, assumes a log-normal distribution. But when you evaluate your OPOS data points that make up the distribution, you use an arithmetic mean rather than a geometric mean, which is to say a median. That may not be very different, but with limited numbers of points, those would differ.

DR. NETON: Yes.

MEMBER KOTELCHUCK: Why do you do

DR. NETON: When one normally calculates an intake, it's a weighted least squares analysis of the data, and if one boils down the calculation for an intake value, it ends up being the sum of the mean value of the excretion values, divided by the mean value of the intake retention fraction.

That's how one would calculate an intake, and so this mean OPOS value is really

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1	sort of a surrogate for an intake value, if
2	you want to think of it that way.
3	MEMBER KOTELCHUCK: That allows
4	you to work backward.
5	DR. NETON: Right. If I had three
6	data points sampled on a person, I would take
7	the average value of the urinary excretion,
8	divided by the average value of the intake
9	retention fraction, and I would get my intake.
10	That's what's and that, believe or not,
11	ends up being a weighted least squares
12	analysis.
13	MEMBER KOTELCHUCK: Okay, thank
14	you.
15	CHAIRMAN MELIUS: Other yes,
16	Gen.
17	MEMBER ROESSLER: This makes me
18	wish that I'd taken more statistics, a lot
19	more statistics. A simple question. If the
20	Peto-Prentice test is always better than the
21	Monte Carlo, why don't you always use that?
22	DR. NETON: I don't know if it's

1	always better.
2	MEMBER ROESSLER: Okay. I had got
3	that impression.
4	DR. NETON: I mean I think we get
5	similar results when we compare the two. But
6	I'd have to defer to our statistician as to
7	why one is more preferable. I think if you
8	assume I think taking advantage of the full
9	knowledge of the distribution of the data
10	would give you a better statistical test, is
11	my opinion, for the Monte Carlo permutation
12	test.
13	Once you start assuming that
14	there's no distribution and such, you lose
15	some power, I think, in your calculation.
16	CHAIRMAN MELIUS: That's correct,
17	Jim.
18	DR. NETON: Thank you.
19	MEMBER ROESSLER: Then I have one
20	more question. When you're talking about the
21	highest exposed group and then the
22	representative group, what criteria do you use

to decide whether a group is representative?

DR. NETON: Now that's the \$50,000 question or \$64,000 question, however you want to phrase it. That requires a lot of digging into the data sets themselves, as to -- I think it's time-dependent. If you look at some of the earlier sites that have been added for SEC already, the data, I would say, were not maybe representative.

But as you get more closer into it's my opinion that the programs time, started to more and more monitor the highest exposed workers. I believe that's true. Т think early on maybe there were sort of cohort model, cohort exposure evaluations, where they would sort of sample a person from the workforce, looking at the highest exposed workers.

But it's a very judgmental thing.

One needs to look at the data set very closely to determine that. One thing I didn't mention is you notice these large GSDs on these

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know, of So, you one the criticisms of this test is that you can't see very small differences. Well that's true. mean we prefer to let the data speak for themselves and say if I can't see a small difference between two data sets that have very large similar GSDs, I think that's sort of obvious that you can't do anything with I mean that's just the way statistics works.

So to presume that they are different at the get-go to me just seems sort of a violation of basic scientific hypothesis testing. But just my opinion.

CHAIRMAN MELIUS: Yes, Bill.

MEMBER FIELD: I was curious that you had the opportunity, and maybe there's data sets that aren't even available, that are pretty complete, to actually assess the validity of the coworker models by self-censoring, and then making comparison using

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these coworker models.

DR. NETON: Yes. We've actually - that's one of the commitments we have for
the ten-year review, is to try to do that, and
we did take a data set of tritium results at
the Savannah River Site, and did some
preliminary analyses and completed it.

At the end of the day, I'm a little bit concerned about interpretation of some of the data. But the original estimates that we came up with demonstrated that the 50th percentile seemed to be fine, with a full distribution for workers who we expected would be in that category, and the 95th percentile worked fine in the other direction.

But you know, that's N equals 1. Tritium is sort of like the easy one, you know, the low-hanging fruit. So I'm not sure how you would, you know, even if you could find and do this for two or three sites and say yes, it looks pretty good, you're always going to have the doubt in the back of your

mind well, does it really work for Site X or Y or a more complicated site.

I think right now, the big question in my mind is the construction trade workers, because there, it seems like you have at least the ability to define who was a construction trade worker and who wasn't. As Dr. Melius pointed out earlier, many of these other sites, you really don't know.

I mean you have an idea that this guy was working here, but you really only know that he was working there for that particular year, and maybe he changed jobs. So it becomes very problematic to identify and segregate people in most cases except maybe construction workers is a unique example.

CHAIRMAN MELIUS: Henry.

MEMBER ANDERSON: Yes. Just kind of on the issue of stratification, it seems that it's, you know, it's all heavily dependent on statistics, and you know, a level of statistical significance. We've all had

relatively small databases, and the statistics says oh, it's not statistically significantly different when you look at it.

You know, you do, your one where "oh, this isn't significantly said you different" changed the scale, and they're really quite different visually. So you know, is there any thought about, you know, what is the level of statistical significance? You're using .05, which is just а convention basically.

could, You it that the seems impetus here is there to say aren't significant differences, so therefore we can, you know, one size fits all for the, know, everybody was the same and you talk to the workers and they say well geez, you know, that's crazy, that we were all very different and we did this different. We're in different facility, different things.

But the measurements, you know, statistically seem to be similar. So you

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might think in terms of, you know, what is that p-value or whatever you're using, to say that isn't stratified or not. You could loosen that.

I mean we often on the epi side, you know, when you try to say what goes into your multiple logistic models, you put them in, if it's .01 or .02 or things like that. So you know, how the level of certainty you're asking for to declare that they are different, you know, makes a difference.

DR. NETON: You raise a good point and, you know, 95th percentile, of course, is standard convention, which is what we've adopted. But I'd also like to point out again, I can't over-emphasize the intake retention model that we developed, that is really -- ideally, I think the statistical test should be done at that intake retention level.

You put the data points in there, and you look for differences between those

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models. But you can't always do that, because it's a somewhat more subjective analysis.

So the idea here with this looking at the data sets themselves and comparing statistical differences at the one person/one sample distributions was to say if we can say up front that there's no difference here, we don't need to go back and look at the practical significance in the model. It's a kind of two-part test.

Because really, the intake that you're assigning is the important thing. I showed you that. Fourteen years' worth of data is 14 years' worth of bioassay points, 50th percentiles. If a few of them are different, I'm not sure it's going to make any statistically significant difference in the intake retention fraction, intake retention function that we apply.

So but I hear what you're saying.

I think you're probably going to hear something similar to what you're describing

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from the next speaker.

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CHAIRMAN MELIUS: I would just add that Ι think it's also to your one person/one sample also ignores error in that. I mean you're taking a mean of that and you're not really addressing the error that's in that, you know, the variability in person's testing. So that can be problematic to do that.

Brad, do you have a question, and then we'll move on to the next presentation.

MEMBER CLAWSON: Yes. Genevieve already took the \$50,000 question, but part of my, and I'm not a statistician or a speaker, but the problem that I see with this is it all comes back to the data and the reliability of the data that you get.

At so many sites, as you said, very well put, that in the later years, you started to see more of a representative sample. The highest people started to become sampled. But you go back into the earlier

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years, and we're looking at very, very small amounts of sampling.

Some can call it event-driven or, you know, you can say well that means it would be the highest exposed. Sometimes they used those to just get a baseline of what some people were getting and what the other people got.

What I really see on this is it comes down to the integrity of the data in question, and the representativeness of the sample of exposed workers was conducted. It all comes down to this, and I'll be right honest. In the early years, we haven't seen too many sites that were that way.

DR. NETON: I 100 percent agree with you. One thing I was going to do is to look at the early years, and which sites -- many of the sites that have large data sets like this are already SEC in those very early years, for really not reasons related necessarily to the coworker model, but because

1	of other issues.
2	But nonetheless, they are SECs,
3	and so they're, you know, that issue has sort
4	of been dealt with in a different way. But I
5	do believe for the more recent years, the
6	data, as you say, are much better and can be
7	used.
8	CHAIRMAN MELIUS: Board Members on
9	the phone have questions?
10	MEMBER RICHARDSON: Yes, this is
11	David Richardson.
12	CHAIRMAN MELIUS: Go ahead, David.
13	MEMBER RICHARDSON: One question I
14	had was you have those great slides that are
15	showing the difference between mean and
16	standard deviation of the distribution of the
17	two groups, and I wondered what is closest to
18	standard deviation for let's say Stratum B.
19	What is closer to the standard deviation in
20	that second stratum?
21	CHAIRMAN MELIUS: I'm sorry,
22	David. I couldn't get your question. You

1	were breaking up quite a bit.
2	MEMBER RICHARDSON: I have a
3	starting question. What factors influence the
4	standard deviation for what are the strata
5	that you want to look at?
6	DR. NETON: What factors influence
7	the standard deviation? I really don't know.
8	I mean it's what the data said it has in it. I
9	mean if I knew all the factors, I guess, that
10	contributed to it, I could tease them out.
11	But I mean a priori, I would have no way of
12	knowing why there's a GSD of 4 versus a GSD of
13	5.
14	In some cases, it is true when
15	they start reporting the data that are below
16	the lower limit of detection, you in effect
17	end up having two distributions.
18	There's an entire report that
19	deals with that, I think it's 44, that talks
20	about a normal distribution underlying the
21	very low data, which is a distribution about
22	background, and then you have this

superimposed log-normal distribution from the truly exposed workers, and you combine those two and you end up with like extremely large GSD on face value, until you do a maximum likelihood evaluation and you can tease out the two distributions.

But aside from that, I don't know of any other way to ferret out the factors that influence the standard deviation.

MEMBER RICHARDSON: Let me put it another way. Let's assume that you had two samples drawn from the same population, and one sample was four times bigger, and our question is are those, do those two groups, those two samples, do they arise from the same population or are they a mixture of two normal, two different normal populations?

DR. NETON: Right.

MEMBER RICHARDSON: It would seem to me that the fact that one sample was four times larger than the other one would influence the standard deviation.

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1	So if we did a test to say do they
2	come from the same underlying population or
3	not and a null hypothesis since they have the
4	same means and the same standard deviation, we
5	might reject that simply because one was
6	bigger.
7	So if one was one-fourth the size,
8	then the log of the geometric standard
9	deviation for the smaller one would be about
10	twice as large as the one
11	DR. NETON: Yes. I guess that's
12	not intuitive to me, that that would be the
13	case. But I'd have to think about that,
14	David.
15	MEMBER RICHARDSON: I mean the
16	standard deviation is a function of 1 over N.
17	DR. NETON: But if it's the same
18	distribution and you just have a smaller
19	sample size, yes, yes.
20	MEMBER RICHARDSON: So I guess,
21	you know, I mean I'm thinking about that
22	scatter plot, the null hypothesis is that

3	hinge on the fact that, you know, that this			
4	one worker subgroup is estimated as precisely			
5	which means we have as much information abou			
6	it as the other, let's say larger group. I			
7	don't know why that would be my starting nul			
8	hypothesis.			
9	DR. NETON: Is Tom LaBone on the			
10	phone? He was possibly going to be able to			
11	join us by phone. If he could maybe entertain			
12	that question? Tom, are you on? Are you or			
13	mute?			
14	(No response.)			
15	DR. NETON: Okay. Well, I guess I			
16	can't answer the question off the top of my			
17	head. So we'll have to take that under			
18	consideration.			
19	CHAIRMAN MELIUS: Any other			
20	questions from Board Members on the phone?			
21	(No response.)			
22	CHAIRMAN MELIUS: Okay. We'll			
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those two samples have the same mean and the

same standard deviation, and that seems to

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1 move on, but don't go away completely, Jim. 2 Arjun will present SC&A's review of OTIB-53. 3 (Off mic comments.) Thank you very 4 DR. MAKHIJANI: 5 I'm really a surrogate for much, Dr. Melius. 6 Harry Chmelynski, and I hope he's on the 7 phone. (Laughter.) 8 CHAIRMAN What is 9 MELIUS: а 10 surrogate? Don't we have a policy on this 11 about surrogate, surrogate data? We should have surrogate experts. 12 13 (Laughter.) I just wanted to 14 DR. MAKHIJANI: 15 put that caveat in. No, but I did review this 16 But what I'm presenting report. essentially Harry's work. So Dr. Melius, you 17 talked about sufficient accuracy and Jim Neton 18 19 went over the broad concept of coworker 20 models, and we've reviewed a number of those

other reports, Report 75, Report 44, Report 95

I think it was.

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This particular presentation review focuses Report 53, which is on essentially when do you compare two groups of workers, and how do you conclude whether their drawn from the measurements were same distribution or not?

So the essential question is can you join all the data into one coworker model, or do you need more than one coworker model, and how do you decide that question? 53 came out at the same time Report number of Savannah River reports, which used specifically the method in Report 53 comparing construction workers with nonconstruction workers, and that's a central question of SRS as you know.

So I'm not going to read all of the slides. I'm just going to give you an overview of some of the things we concluded, and Harry, please feel free to jump in.

One of our central conclusions was that when you're comparing two groups of

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workers, the sampling protocols of the workers should be same. That is if you're routinely monitoring one set of workers, the other set of workers should also be routinely monitored, and then you can compare the distributions and say well, one was much more exposed than the other. They were drawn from different distributions.

But if their monitoring protocols were different, then you can't really be comparing those saying two and are the distributions the same not, because orpriori, their monitoring protocols different. In the specific case of Savannah River Site, a number of those reports, including Report 56, which was I think, I think Report 56 was for trivalent, I can't remember.

Anyway, Report 56 and 58 deal with specific radionuclides at SRS, and both 56 and 58 say that construction workers had potentially a different bioassay monitoring

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protocol. We talk about whole body monitoring protocols, which is a whole different and more complicated issue actually.

being So is routinely one monitored and other not, then we don't feel those can be compared. So the comparisons at Savannah River Site are in question, at least when urine data are being used in that regard. So there's a question of representativeness of the data. So we have people who monitored. How do they -- I'm sorry for the unclarity of the slide. How do they compare with the people who are not monitored, characteristics what the of the are unmonitored population?

So to give you a specific example, if pipefitters were not monitored, were they the same as construction workers in general? Were they the same as the general monitored population of construction and non-construction workers?

So there's a question of

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representativeness of data. We feel that based on our prior work that we submitted to the Board and the Work Group, that at least at Savannah River Site, which we've examined in detail, it was necessary to compare subgroups of construction workers, because different construction workers have different exposure potentials.

I might mention here, Dr. Neton mentioned, you have to select a priori whether there was a difference in the groups or not, and to some extent we can see that the types of work construction workers did, some carpenters or electricians and pipefitters were different types of work, that may have had different exposure potential.

So there's some technical underlying reason to look at groups of construction workers, and that was borne out by the examination of some of the data that we already presented to the Board in previous reports.

So you have to actually at Savannah River Site, we show that you have to examine by job type and area of work. So you have to do multiple pair-wise comparisons. So one, simply saying construction workers versus non-construction workers may not be enough.

Now we don't know whether it would be enough at some other site. It may be perfectly okay at some other site or not. The detailed examples that we have done in terms of analysis are from the Savannah River Site.

So I'm just using those to put the caveats on the report, not saying generally it would be necessary to do it. But it should at least be examined whether there are groups of construction workers that are different from each other.

Once you start parsing one group of workers into subgroups, then you run into data problems, because you have to have a minimum of 30 samples for each category. It's not necessary that 30 samples may be enough,

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as I will show. But you have to have a minimum of 30 samples in each category.

That practical into you run hurdles, because non-construction workers were frequently. monitored But with more construction workers, you do into run monitoring problems.

Another problem is that when you aggregate the data, you're not averaging into one person, one sample, a large number of data points over a single year, and sometimes NIOSH aggregates over two years or three years even. But we know from interviewing workers that somebody sometimes may start out as а construction worker, and may be then hired by the contractor and become a non-construction worker.

So within the period of averaging, their job designation may change, and yet we didn't see that NIOSH has a method of actually sorting out individual getting in and an worker's data. Perhaps they do it in

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practice, but we haven't actually examined in detail, you know, the entire compilations. But we haven't seen it actually done.

That would take a fair amount of work to actually go and find examples of workers whose job designations were changed. But we do know that job designations did change, and this is a little bit of a problem that needs sorting out. It may be a non-problem if job designations didn't change very often relative to the number of data points.

So we have some problems with the power concerns, which was Finding No. 8. Now there are a number of ways to do comparisons. So you can start out with a null hypothesis, as Jim Neton says, that you assume that they are the same, and if you don't know anything in advance, this is a pretty reasonable way to start out the comparison.

We say we're going to assume the same unless proven that they are different, and that is the approach of Report 53.

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Confidence level of 95 percent is set, that you don't want to falsely reject the hypothesis that they are the same, and you want to be very sure of that. But that's called a Type 1 error.

other But there's the type of error, that you may falsely accept the null hypothesis that they are the same, when in fact underlying distributions the are different. So that is a problem, because these two types of errors are in tension with each other, if you don't have a sufficient amount of data.

To pick up Dr. Richardson's question, the dramatic standard deviations and the relation of those standard deviations with the geometric mean ratios comes into play. So you may actually run into a problem, where even 30 samples may not be enough, and I will illustrate that. Let's see.

So again, I apologize. Let me go to the graph. It might be easier to see it.

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So this is a chart that illustrates this problem. So here, you have this Type 1 error, and this is the power level that is set, 95 percent confidence, and these strips show the Type 2 error rate.

So you can see in this third dimension is the geometric standard deviation. As the geometric standard deviation increases, and remember now the ratio of the geometric means is fixed. So this is a very simplified calculation. The number of non-detects is fixed. The ratio of geometric means is fixed.

So we're only examining the influence of the geometric standard deviation on the error. If you keep the Type 1 error at five percent and the geometric standard deviation increases, you are falsely accepting the null hypothesis. That is, you're saying they're the same, when they're not actually the same.

If your geometric standard deviation is small, then your Type 2 error and

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Type 1 error can both be controlled, and you're in very good statistical shape, but not always. And of course, if you have a small number of data points with high variability, you're often going to wind up in the region where your geometric standard deviation is large.

And as we have examined, I don't have ready examples for you to present, but I think you will see, as we complete our work on Savannah River report, that this is a pretty big problem in practice. But this a simplified example, and real life actually gets much more complicated than this.

So we examine what is the effect of small sample sizes. Let me show you this table. Sorry, there's a table here. So this is a table of neptunium data from Savannah River Site, number of all posts, you know, these consolidated one person/one sample by year, and you can see except for the 60's, in fact we went into the 70's, '74, they're only

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ten OPOS data points for construction workers.

Throughout the period, you actually don't meet the minimum number of data points that you need. They're never more than 30 in the entire '74 to 1989 period. The practical effect of that is you could have the geometric means being different in 1974 by as much as 3.8 or a different test, as much as 3.5, and you would still say that the distributions are the same.

So you wind up in territory that could be very claimant-unfavorable, when you're saying you're going to ascribe doses to construction workers based on all monitoring data, which is dominated by non-construction workers, because they were the most frequently monitored, and for -- you could be off by as much as a factor of 3.8, 10, 12, 15, 11.

You could be off by a very large factor. So you could be very claimant-unfavorable, and this is shown in the chart. So this chart kind of illustrates when you set

a Type 1 error rate that is at this 95 percent level, you have this whole gray region basically of indecision.

You could wind up in a situation where you're saying yes, they're the same, when applying that hypothesis could result in a very non-claimant favorable dose or intake calculation.

So this is a very significant problem, because in practice you run into these data limitations, and very often when you have even more than 30 data points, if your geometric standard deviation is high, then your Type 2 errors that is falsely accepting the null hypothesis can get out of control.

If you relax the Type 1 error from 95 percent to 90 percent, because there's a tension between these two errors, you can reduce the Type 2 errors. But you know, it depends on how big your -- your geometric standard deviations and geometric means are in

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certain relation to each other, you can't fix this problem without a sufficient number of data points.

There are alternative approaches available. You could do a different test. We started with a test. NIOSH started with a test that will assume they're the same. Of course, you could start with the opposite test, which is more or less the same thing. You have opposite definitions of Type 1 and Type 2 errors.

But you could also start with a test saying non-construction workers were typically more exposed than construction workers, and you could test that hypothesis using the same set of data. You could start with the other tests, saying non-construction were less exposed than construction workers, and you could test that.

Typically, that second test would be more claimant-favorable if what you're examining is construction workers. The first

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1	test would be more claimant-favorable if what
2	you're examining is non. So it's not a priori
3	given; at least we don't agree that a priori
4	you should set a null hypothesis that says
5	they're the same, and examine whether they're
6	different or not.
7	In fact, some of the data indicate
8	that they are different, so there's no a
9	priori reason to assume that they're the same
10	and test that hypothesis. So that concludes
11	my presentation. Harry, did you want to jump
12	in and say something supplementary?
13	DR. CHMELYNSKI: Yes. Arjun, can
14	you hear me?
15	DR. MAKHIJANI: Yes.
16	DR. CHMELYNSKI: Okay. Earlier in
17	the morning here, there was a discussion on
18	sufficient accuracy, and I believe the latest,
19	the last figure, which is Figure 2 on page
20	ten, shows from a statistical perspective what
	11

we mean by sufficient accuracy. It's really

the width of that gray region.

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In other words, how much different do they have to be before I know I can tell them apart? I see no information on that sort in Report 53. The reason the tests are adopted is, as stated, that they were powerful tests. But we don't know how powerful they are, given the kind of variabilities that we have with the GSDs being high, and also how well the data themselves are reflecting the distribution.

Earlier also this morning, we had a discussion about OPOS. One of the problems with OPOS values are if you complete an OPOS value with one person's data for the year, and all we had was a couple of samples, you get a highly variable estimate of the OPOS mean.

On the other hand, if you have another worker who had a lot of samples, part of a regular protocol sampling, and in fact they may most likely be the non-construction workers, then that OPOS value is estimated much better.

So there's a basic problem of heteroscedasticity that's introduced once you start using OPOS values, and then trying to conduct tests on these values, where you're assuming they are independent samples, just doesn't make much sense to me, because that's not what they are anymore.

So the gray region isn't the really hard question here, I think. It's how far apart do they have to be before we're going to say they're different. I have yet to hear anybody answer that question. So we don't know what we're trying to do here with the test.

Granted, we could always say, hey,

95 percent will tell me whether they're
significantly different. Well yes, but if you
don't have enough data, you're always going to
say that they're not significantly different,
so the test really doesn't tell you anything.

The right way of doing these tests is to define how big a difference you have to,

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that you want to be able to detect, and then calculate the sample size that will allow you to detect that size of difference when there is a lot of variability.

Thirty may be enough, certainly no less than 30. But once you get up into the GSDs, as shown in the other figures, once you get up into GSDs of 5 and 6 and you're comparing these populations with a 95 percent confidence level, it's very hard to show that they're different.

that's not very claimant-To me, favorable. What we're saying is that the construction workers have to prove they're different. Now they don't even -- they don't know that we're asking them to do this, of But what we're going to say is oh, course. you guys are all the same, unless somebody can produce sufficient data to show you're different.

Well, we already know there isn't very much data. So you know, the idea of

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starting out with that null hypothesis that they are the same leaves me very unsettled. I guess that's the gist of what these 15 slides are trying to say.

DR. MAKHIJANI: Well, a lot of Harry's remarks relate to the analyses we've done on numbers at Savannah River Site. It may not always be true, but it's certainly true at Savannah River Site, that because of the nature of data, very often you don't have enough data.

The result is very claimant-unfavorable. If you conclude they're the same and apply that coworker model in all the years that we're looking at to apply this data, '74 to '89, you would be applying a result that would be very claimant-unfavorable. So that in this specific instance -- now it may not always be true.

So there's the caveat. This is what we've examined so far, and are continuing examinations into thorium or along the same

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lines, but we haven't finished. So we have some significant problems with the practical implications of the OPOS approach. I mean we don't, we don't disagree with the idea that if you have an incident and somebody sampled many times, that you have to take that into account somehow.

But in practice, I think applying the OPOS approach as proposed in the comparison of the two workers doesn't seem to well, work in of claimant very terms favorability, among other problems.

DR. CHMELYNSKI: Then Arjun, I'd also like to add that there's two topics we haven't addressed on these slides. One of them is the use of the regression on order statistics, and in our review of PROC-95, we indicated that there is a problem with the ROS method, because the data you're using in it are auto-correlated and heteroscedastic.

The auto correlations are quite high, around 0.6, 0.7, something like that. So

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because of this, the diagnostic statistics we see for those log-normal distribution fits really are not supported by the data. In other words, even when we say that there's a confidence level of 95 percent, that may not be the right answer.

The second issue here that we haven't talked about is that the use of OPOS - I'm sorry, in the first five pages, as we talked about how the data set should be derived using the same protocol, I think all those conclusions apply regardless of whether you use OPOS or not.

Now OPOS introduces another dimension to the problem of comparison, but again, if you're using data that's collected under one protocol and then trying to compare it to another set of data that's collected using a different protocol, I just -- I don't see how a statistical test is going to tell you anything. I guess I'm done.

CHAIRMAN MELIUS: Thank you.

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Questions for -- go ahead, Dave.

MEMBER KOTELCHUCK: I have to say that in terms of claimant favorability, using a one-sided test really doesn't make sense. It is, I mean as you know, in a lot of scientific studies you always use the two-sided test. It's the hardest thing to prove, because you would really like to be confident of the result.

But in this case, we know that -we believe that the construction workers
probably have a lower, they should have a
lower exposure than the people who are the
non-construction workers. So even with a 95
percent probability but a one-sided test,
we're much more claimant favorable.

I wouldn't take a position about 90 versus 95, except in terms of what you said. It would ease things. But at a simple level, a one-sided test would be an improvement in terms of claimant favorability, and I don't think that it would involve --

1	Jim, I don't think that it would involve major		
2	changes in the work, in the tests that you		
3	had proposed, right?		
4	DR. MAKHIJANI: The one comment I		
5	would have is just to refer to the last thing		
6	that Harry said, is that the comparisons are		
7	based on an underlying assumption that the		
8	monitoring protocols were the same.		
9	MEMBER KOTELCHUCK: Yes.		
10	DR. MAKHIJANI: And so a lot of		
11	the problem that we're having with the		
12	monitoring protocols, we either know are not		
13	the same and, as we've looked at whole body		
14	counting data, we're not able to establish a		
15	monitoring protocol because, in some cases,		
16	monitoring was quite infrequent, and you know,		
17	protocols are supposed to be annual.		
18	It's very complicated to establish		
19	a monitoring protocol sometimes. In case of		
20	urine data, you can actually talk about it,		

MEMBER KOTELCHUCK:

and sometimes you can't.

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That

I agree.

1	is the major problem. Nevertheless, a one-
2	sided test would be an improvement, and it's
3	fairly simple to do.
4	DR. MAKHIJANI: Yes. That was one
5	of our recommendations, that NIOSH should
6	examine whether a one-sided test would be
7	better.
8	DR. CHMELYNSKI: What I might
9	interject here is that the Monte Carlo
10	permutation test may not be amenable to
11	turning it into a one-sided test. Certainly,
12	you can do that with the Peto-Prentice test.
13	But I'll leave it to NIOSH to decide how they
14	would do that with the Monte Carlo permutation
15	test.
16	MR. LaBONE: Hello? This is Tom
17	LaBone. Can you hear me?
18	CHAIRMAN MELIUS: Yes, we can.
19	MR. LaBONE: I'm sorry. I could
20	not master the *6 back when Dr. Richardson
21	asked the question, and I was wondering if I
22	could address that now.

1	CHAIRMAN MELIUS: You sure can. Go
2	ahead. Sorry.
3	MR. LaBONE: Okay, okay. A number
4	of the concerns that are being raised, I think
5	if you think about this, these are
6	retrospective studies. We do not get to plan
7	the data that we get. We're presented data
8	and we were asked to make the best statistical
9	analysis we could of each data set.
10	So everything Dr. Richardson was
11	saying is correct, is that if you happen to
12	have a smaller construction trade workers
13	or a smaller number of individuals, then you
14	have issues with that. That's basically what
15	we have.
16	The other point is that there's a
17	common thread here is that failure to reject
18	the null is not equal to higher doses for the
19	construction workers. Is that that's not
20	necessarily true at all. But you might come
21	away from this conversation that it is.

So for example, the coworker model

built from the combined data set might actually give higher doses to the construction trade workers. It depends where are they relative to the other subgroups of workers.

The other thing is that a lot of the issues being raised are generic problems with null hypothesis testing, this failure to reject the null, the issues with power and so forth. There are other ways of handling this that can get around that. But what we have to do is you have to define what is practically significant.

I think Harry alluded to this, and this is a very difficult thing to establish. take these data it's Ιf you sets and neptunium-237 in urine, what is the practically significant difference in the concentrations of neptunium in urine?

We looked at that, and we just can't come up with a way to generically do that for every data set we look at. But that would get around the problems of this null

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hypothesis testing, where you fail to reject the null, which is basically an unacceptable answer apparently.

So that was just -- and the last thing was again, the OPOS process is not an ideal solution. But we feel that it actually solves more problems than it creates and, again, this can go into some more technical details. But those are a couple of issues I wanted to point out.

CHAIRMAN MELIUS: Thank you, Tom, and we appreciate how complicated this is, and you're right, that you have to make -- we're dealing with retrospective data, and trying to make the best we can from it. It's difficult, and I think it's also difficult to do this knowing sort of, Ι think without Harry described, what gray area are you aiming for, and what parameters are you trying address here.

Those are not defined, and that makes it even more difficult. Thank you,

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Arjun.

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DR. MAKHIJANI: May I make a couple of comments? I agree with Tom. I wouldn't want you to go away with the impression that, in applying this formula know, you construction workers and non-construction workers will always come up with something that's not claimant-favorable. I believe I actually did say that during my presentation.

It's just that in these examples, and with the specific data sets that we've looked at from Savannah River Site that is the result, and partly it may be the result because the monitoring protocols are different.

If you look at the neptunium report that we have submitted from the same data set, you'll see some considerable discussion of this very point, that are you coming up with higher results because the monitoring protocols were different, or were the monitoring protocols deficient and missing

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routine doses.

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So there are many permutations and combinations of this problem. I would also agree with Tom, and then we said this when we looked at external dose data for Savannah River Site in the context of a different procedure, that applying this procedure, you could give higher doses than the working conditions warranted for groups of some construction workers.

If you look at the report that we submitted on tritium some time back, a couple of reports that are referenced to you, you'll see that that is actually the case. It can also vary by period. The point here of saying that you should, you need to parse construction workers into subgroups is because t.he nature of their work was actually different, and their exposure conditions were different, and the data actually show that their exposure conditions were different.

So while this formula may be

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claimant-favorable for many construction workers, it will also be claimant-unfavorable for many other subgroups of construction workers. Whence the need to parse construction workers into groups, and the need for a lot more data. As it is, we don't even have enough for construction workers as group quite often. Sometimes you do actually have enough data.

CHAIRMAN MELIUS: Thank you. Any other comments, questions? It's getting towards lunch time, I can tell. Yes. What I would like to do is let's talk when we do our Board work, why don't we come back and talk about next steps then? I have some thoughts, but I want to talk this over with NIOSH and a few other people before I put my foot in my mouth or something here.

I will say even though Arjun, I like your three-dimensional power graph there, I still think that Jim wins the graphics prize

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1	DR. MAKHIJANI: I yield.
2	CHAIRMAN MELIUS: Or whoever came
3	up with that. So that's what someone's going
4	to have that's still the top graphic for
5	this meeting. We'll take lunch and we'll
6	return at 1:30.
7	MR. KATZ: Just one administrative
8	thing for Board Members. Those of you that
9	haven't sent in your updated ethics form that
10	was requested, that I sent to everybody,
11	either email it yourself to the email address
12	that they give you in that, or you can fill it
13	out here.
14	It doesn't take any time to fill
15	out really, and sign it here and give it to me
16	and I'll give to Zaida or we'll get it there
17	somehow, if you can't scan it in yourself,
18	whichever.
19	(Whereupon, at 12:09 p.m., a
20	luncheon recess was taken.)
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2 AFTERNOON SESSION 3 1:41 p.m. CHAIRMAN MELIUS: Okay, if we can 4 5 return to business here, reconvene. Ted, do 6 you need to do attendance or anything? 7 MR. KATZ: Yeah. Let's just check to see who we have from the Board on the line, 8 that's all. 9 10 Ziemer, are you on with us again? 11 MEMBER RICHARDSON: This is David 12 13 Richardson. KATZ: David, welcome. 14 You MR. sound really nice and clear now. 15 16 MEMBER RICHARDSON: Thank you. Ι feel clear. 17 Very good. 18 MR. KATZ: 19 Ziemer, are you on the line too? 20 (No response.) Okay, maybe I should 21 KATZ: 22 just check. Dick Lemen, are you on the line?

(No response.)

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CHAIRMAN MELIUS: Okay. But we have a quorum and we can proceed. Okay. This will afternoon be the LaVon Rutherford presentations, so hang on. We're ready, and we'll start with an update on the Rocky Flats petition. I believe LaVon will present and Mark will comment, and then we'll hear, open up for questions from the Board, and we'll also possibly hear from the petitioner.

Rocky Flats SEC Petition Update

MR. RUTHERFORD: Okay. Thank you, Dr. Melius. I'm going to talk about the Rocky Flats petition evaluation, where we currently stand, what's -- and what we're going to get done here in the future.

A little reminder. We completed our Evaluation Report and issued it on September 5th of 2012. We presented that to the Advisory Board and the public on September 18th, 2012 at the Advisory Board meeting in Denver.

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At that time, the Board made a determination at the meeting that we needed to do additional review, needed to send it to the Work Group, SC&A and the company or the Board's contractor to look at, and we needed to do some additional interviews and discussion and document review.

So our follow-up efforts that we conducted since we presented the evaluation, we've done data capture, both classified and unclassified, Los Alamos National Lab, OSTI, Office of Scientific and Technical the Information, EMCBC, which is Environmental Management Consolidated Business Center, and Legacy Management. We did those Denver. We again as I said out in Los Alamos, and yeah, Los Alamos and at OSTI.

We also had secure discussions, and I say secure discussions, because these were classified discussions over different things internally. We also secure interviews and we also had unsecured interviews, roughly

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19 interviews that were conducted. We discussed during those interviews not only the tritium issue, which was the main focus of the evaluation, but we also discussed things like the neptunium and other things that had come up during our additional data captures and review that became open issues, that we felt needed further follow-up.

Both the Work Group and SC&A and ourselves internally felt we needed further follow-up. Then we also did some additional dose reconstruction modeling. If you remember back in the evaluation, when we had presented the evaluation, we had come up with a bounding exposure of roughly 700 millirem, that based on the 1973 incident for tritium.

During our presentation of that, we committed to look at that, to see if we could come up with a little more of a precise analysis.

So to give us a status on where we are since these additional efforts, we have

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completed or we have five White Papers that we've worked on, two of which have been completed, and I'll discuss them a little Those five White Papers are made up further. of follow-up efforts on the tritium issues; evaluation of petitioner concerns about data falsification and/or data invalidation Rocky Flats Plant Building 123, based on worker allegations; a White Paper on U-233 and thorium strikes; and а White Paper on neptunium.

The final White Paper working on is other thorium activities, which came about from our additional data capture reviews and interviews. The first White Paper on tritium, we actually issued our report after we had done the data captures, interviews, went back and looked at all the additional information, and updated we basically our position on the tritium exposures at Rocky Flats.

We issued that White Paper on the

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25th of June. We provided that to the Work 26th and Group the the petitioner on unfortunately did not get that until 7/3, July 3rd, because it was at ADC review. At that point, we had made a point to -- a note that we recognized that we were giving very little time to the petitioner, very little time to the Work Group and SC&A to prepare responses for the Work Group meeting. So that was pointed out.

Based on that, the discussion was that we would present that paper at that Work Group meeting, and then allow additional time for the Board or Work Group and the petitioner and SC&A to review that information, and have a follow-on Work Group meeting at a later date.

The second White Paper that we presented was on the data falsification and data invalidation. This actual White Paper was brought about by a petitioner, by one of the co-petitioners, who had identified a

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document that presented a potential falsification or validation, lack of validation of data at Building 123, just based on worker allegations. It was an interview that was conducted by the FBI and the EPA.

We issued our report on that June 25th. We provided it to the Work Group and the petitioners on July 3rd. We presented that to the Work Group at the Work Group meeting on July 8th, and again we pointed out that we recognized the short review time of that information, and based on that, the Work Group intended to do a more detailed review and have a follow-up at a later meeting.

There are three other White Papers that we're working on. I'd say thorium strikes U-233. This was an issue that was actually, it was evaluated during the first petition at Rocky Flats, SEC 30 I believe, and we went back and based on some additional discussion during our classified interviews, and some of the reviews that we'd done, we

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decided that this needed to be revisited.

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had indication of additional thorium strikes that weren't previously identified. So we committed to develop a White Paper for that. That White Paper is very close to completion. It's not only going to look at the exposure from the strikes, but it also looks at the U-233 exposure. Again, this is very close to completion. We plan on having this out later this month.

Neptunium. This is another issue that actually came up during our classified interviews and discussions, and also during our document reviews. We went back and we looked at the transcripts and what had been done in the previous evaluation for neptunium, and felt like that issue really had not been thoroughly vetted.

So we committed to Work Group that we would put together a White Paper on the neptunium. We have come up with some additional issues on that; however, we do plan

on having that complete by the middle of August, in support for a Work Group meeting prior to the October Board meeting.

"Other Thorium Issues" is another White Paper that we're working on. This is again another item that during our classified reviews and some of our interviews, some additional items that we felt had not been really looked at closely previously and during the previous evaluation. We felt it needed a little more of a thorough look.

together, we've put So working on another White Paper. It's called "Other Thorium Issues." It's basically looking at the other activities outside of thorium strikes associated with thorium. We included, based have on the Work Group suggestion, the magnesium thorium discussion that was brought up by Terrie Barrie. So we are going to include that in that White Paper.

I think that's it for me for an update on where we are. Actually, out of the

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Work Group meeting, I will make a couple of comments, and I'm probably stealing some of Mark's thunder, but I apologize.

We are, had some things come out of the Work Group meeting. We are going back, and we're going to look at some information out of Pantex, to determine, based on some of the modeling that we had done, whether changes in the Pantex program had occurred after the 1973 incident at Rocky Flats. So we're doing some additional work there.

We're also doing some additional interviews on the data falsification/data classification. We're going to interview some of the former workers that worked at that time period, to if better see we can get а understanding if there was any change in their analysis techniques between before and after the raid that occurred in 1989.

There were a couple of other items that came out, action items that came out of the Work Group meeting that we are following

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1	up on as well. That's it.
2	CHAIRMAN MELIUS: Thank you,
3	LaVon. Why don't we hear from Mark, and then
4	we'll open up to questions.
5	MEMBER GRIFFON: And I don't
6	really have much that was a good summary by
7	LaVon. I didn't catch whether you mentioned
8	the timing for our next scheduled meeting. Did
9	you
10	MR. RUTHERFORD: No, I didn't. The
11	date is September 12th or 17 th . 12th,
12	September 12th.
13	MEMBER GRIFFON: September 12th,
14	yeah. So the hope is by then we'll have a lot
15	of these items that you mentioned in complete
16	enough form, that we can
17	MR. RUTHERFORD: We are, I think
18	we're putting a lot of priority on getting
19	these done. I know that the focus is for the
20	Denver meeting in October. So we're going to
21	put a lot of effort towards that and hopefully
22	get it done, and a little better time frame

1	for the petitioner and the Work Group.
2	MEMBER GRIFFON: Yeah, and we set
3	that time to allow, hopefully to allow SC&A
4	enough lead time to also review all this
5	stuff, because the last meeting we had, it
6	ended up being a phone call meeting, because
7	we were kind of the documents didn't get to
8	us with very much lead time. So hopefully
9	that's resolved, yeah.
10	MR. RUTHERFORD: Yes.
11	CHAIRMAN MELIUS: Is that a
12	promise?
13	MR. RUTHERFORD: You know, it's
14	hard to promise when it's just hard to
15	promise.
16	CHAIRMAN MELIUS: Okay. Do Board
17	Members have questions?
18	(No response.)
19	CHAIRMAN MELIUS: If not, I'll
20	start. I'm just a little puzzled by the
21	presentation. I'm trying to understand. Are
22	we making progress, I mean in terms of

1	addressing issues or not? This is all sort of
2	what's happened and what's been produced. But
3	the only content I sort of see is from some of
4	the petitioners' comments, which you'll hear
5	in a second.
6	I mean we'll come back after that,
7	okay, if that's how you want, would rather do
8	it. That's fine. I just didn't, I'm just
9	trying to understand where we really are with
10	this. I thought we were sort of farther along
11	or that we were resolving things.
12	MEMBER GRIFFON: I mean I guess
13	we're in the midst of, you know, we got some
14	assessment of these issues, but we got it at a
15	very late stage, right before the Work Group
16	meeting.
17	MR. RUTHERFORD: Right, right.
18	MEMBER GRIFFON: So we had sort of
19	a preliminary discussion on the Work Group
20	call, but SC&A's really got to come back with
21	a little more analysis on those issues, and

see if we --

1	MR. RUTHERFORD: Yeah. The time
2	period on the papers, I mean, was our fault. I
3	mean I should say it was from our end and, you
4	know, we had initial reviews of the White
5	Papers that came through. We identified our
6	concerns, you know. We had the sequestration
7	came through and dealing with all that.
8	I'm not pushing it all on that.
9	I'm just saying that we wanted to get out a,
10	you know, a product that we could live with.
11	So it was, it took a little time.
12	CHAIRMAN MELIUS: No. We've
13	already concluded it was your fault.
14	(Laughter.)
15	CHAIRMAN MELIUS: No matter what
16	you say. I guess let me put this in a way
17	that's a little bit more fair. You know, and
18	this is both for NIOSH and the Work Group. Do
19	you think that we've, through these additional
20	White Papers, have identified key issues that
21	will address and I won't say close out the

SEC, but will make significant progress for

the time for the October meeting.

I mean you can't judge what else is going to be found when you're still collecting data and reviewing stuff. So I don't think it's fair to ask that. But at the same time, are we going to be ready to make a recommendation by October?

MR. RUTHERFORD: I think the only White Paper that's going to be the holdup is the Other Thorium Issues. Yeah, you can see it's a September date, and it's just whether we can get all the information pulled together on that one or not.

The other White Papers, I don't see a problem in getting them together. I think I'm going to, you know, we're working with Joe, Joe Fitzgerald at SC&A. We're going to make sure he's involved and any of the other Work Group Members that want to be involved with the interviews that we conduct for the data falsification.

We'll make every effort we can to

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2	get the papers out in support of a Work Group
3	meeting, that we can try to get some
4	resolution.
5	CHAIRMAN MELIUS: Okay. Mark, you
6	have anything?
7	MEMBER GRIFFON: Yeah. I mean I -
8	- that's why we set this Work Group meeting to
9	September, was because we wanted some
10	significant progress, you know, and something
11	to report in the October meeting. I mean I'm
12	a little, I am a little concerned about the
13	thorium, and I think that's a big issue
14	obviously, the whole Other Thorium and you
15	know. I'm not sure we're going to be able to
16	resolve that in time for the October meeting.
17	But I think we'll definitely make
18	progress. We have to make some progress.
19	CHAIRMAN MELIUS: Again, explain
20	to me what the Other Thorium issue is?
21	MR. RUTHERFORD: Well, I'll just
22	yeah. I won't get into details, but there

have it -- at least answer everything, and

1	were other thorium activities that occurred,
2	and we're trying to determine the scale that
3	they occurred, what they occurred, and that I
4	don't think were previously addressed in the
5	last evaluation.
6	CHAIRMAN MELIUS: Okay. That
7	makes sense then. Okay, okay. Thank you.
8	Other Board Members? Board Members on the
9	phone with questions?
10	(No response.)
11	CHAIRMAN MELIUS: Okay. If not,
12	then I think the petitioner has comments, or
13	one of them.
14	MS. BARRIE: Thank you very much.
15	Yes, and I will keep this as brief as
16	possible. The petitioner himself had some
17	health issues to deal with, and he asked me to
18	read his comments.
19	CHAIRMAN MELIUS: Okay. Can you
20	just identify yourself?
21	MS. BARRIE: Oh, I'm sorry.
22	CHAIRMAN MELIUS: We know who you

are, but the people on the phone --

MS. BARRIE: My name is Terrie Barrie, and I'm the co-petitioner for the Rocky Flats workers. I want to thank you for this opportunity and thank everyone who has helped me, you know, Rocky Flats workers and other advocates on this presentation on the SEC.

The interviewee, the EPA interviewee attested that the bioassay samples sat on the shelf. NIOSH said there is no scientific basis for concluding that sample counting performed weeks after collection would compromise the results.

Really? NIOSH said he couldn't find any bioassay procedures for Rocky Flats, which I find very odd, especially since the original author of the Site Profile was the manager of the Health Physics Lab. Fortunately, the LANL petitioners shared a section of his petition, and the NCRP report states, and I quote, "All biological samples

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are subject to deterioration by bacteriological action that may interfere with subsequent analysis."

I would think that the health physicist would know that. And what about tritium assays? If a sample's slated to be checked for tritium exposure, sat on the shelf for a week or two, would the bioassay results be accurate? I don't know the answer to that. That is something that I think NIOSH needs to address.

NIOSH also interviewed a Mound employee, and the Mound employee said that it is high opinion that, and I quote, "It's a valid assumption that Mound procedures would be representative of other DOE sites."

However, I found a 1995 document that states otherwise, and it says, Slide 3, "The information gathered from these questionnaire responses illustrates the diversity of international dosimetry practices at DOE facilities."

So it cannot be assumed that the Mound dosimetry procedures are representative of all other DOE sites. I am thankful that DOL is willing to check with their Legal Department, on whether it would be possible to petition the Court to unseal the records seized during the FBI raid.

ANWAG and other advocates will be happy to submit briefs to the Court supporting this proposed motion. But I still don't understand why DCAS is insistent that there was no problems at the Rocky Flats plant, both with regards to the bioassay program and the worker protection program.

I have discovered a few documents with support worker statements that all was not well at Rocky Flats. I won't be able to go into and explain every one of them, but I'll be happy to talk with you individually if you need some explanations.

These following slides from DNFSB shows a variety of problems here. We have

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concerns here with the Rocky Flats plant in 1993, with worker air monitoring and "air monitoring in the workplace at Rocky Flats is not in compliance," and like I said, there's others on this projection.

I'm just going to skip back to the GAO report. So even the GAO identified worker protection issues pre-FBI raid. This report also notes that there was improper use or placement of air monitors, and a year later, they identified a lack of adequate measurements and documentation on extremity doses for certain workers.

So let's move on to the tritium, since there is just small time limit here. My understanding about tritium is like the element itself. Information seeps into my brain and just as easily it's seeped out. Some knowledge remains, though. So a good portion of this presentation is just going to be observations or questions to the Board, NIOSH and SC&A.

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In Slide No. 9, which is Okay. before this one, we had asked DCAS to explain to find other information about tritium stripping on Building 444 in 1987. This has not been addressed as far as I can tell in the tritium White Paper, and I think that's important, that we need to get to the bottom of that.

Is there more documentation? What is tritium stripping on or in Building 444? I think the workers need to understand that, and the Board also obviously.

You should also note on this slide, this is the document seized during the raid, which refers to a tritium release from Building 776 in April of 1989. I just located this, and that's why it hasn't gotten to, very far before today.

I'm not sure if DCAS was aware of release, of this release. If they were, did they determine that this release was less than the levels in 1973 and '74? But this is one

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thing I would appreciate an answer on.

During last week's teleconference,
Dr. Makhijani asked NIOSH if metal tritides
were present at Rocky Flats. NIOSH replied
no. There are many Rocky Flats workers who
are helping us, and one such worker informed
me that tritium metals and tritium oxides were
indeed on site, and experiments were done in
Building 559, Glove Box C-1 by [Identifying
information redacted] using tritides.

Many of the workers interviewed during the focus group and other interviews mentioned the fact that tritium alarms went off frequently. I know some of those accounts occurred after 1974, including one from the petitioner. Did DCAS find any information on tritium alarm incidences, and if so, did they rule out those alarms were caused by releases that were less than the 1974 exposure?

Special tritium compounds, where does that fit in, or does it fit in when it comes to reconstructing dose for tritium

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exposure? Considering the documents that I located on the monitoring insufficiencies at Rocky Flats, there are serious doubts in my mind that the records NIOSH is using to reconstruct dose are true and accurate.

One more slide here. Which brings us up to this document. I have that document right here, a copy of it, and it's a 1996 memo from Silverman that Mr. Mark says destroying Rocky Flats records." So it's easy to assume that even though the documents that say we're not going to destroy records and we have everything, they were still doing it in 1996.

leave you with one I'd like to last thought from before Ι read me [Identifying information redacted] quick response, and while this SEC petition is two years old, that's kind of short in the time in SEC petitions, these Rocky Flats issues have been around for eight years.

We deserve an answer and quickly.

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It's not like these are brand new issues to anyone. They need to be investigated sure, but not another eight years down the road do we need an answer. I've been working on sick worker issues for 18 years. I want to retire in two. So I would like to have this resolved by then.

for the quick response from [Identifying information redacted]. "While will DCAS believe the workers or other experts that fit into their predetermined position? of people telling DCAS have tons records are missing, that they have a zero for badge reading, that instruments а were recalibrated to show a background reading that was higher than what the workers' badges read.

"Could the reason all of these zero readings be that the lab was remiss in following scientific protocols? No one has believed these workers. We supplied the EPA interview to bolster the workers' testimony. It came from someone who had direct knowledge

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of the lab's practices, and worked in the lab for a number of years.

"It is unforgivable that NIOSH would dismiss this important information. Yes, the raid happened because of environmental crimes. But the Tiger Teams looked at the whole plant.

NIOSH all four "Does have assessment team reports, just the orenvironmental one? If they do have the four reports, did they review them all and determine that the Tiger Teams found similar problems with personal bioassay lab procedures, or didn't they?

"If they had the reports but didn't read them, why didn't they? Ιf all four reports are not available to DCAS, not? Who has them? If DCAS cannot obtain all four reports, how can they emphatically assert that there were no problems with the worker bioassay program? Will we ever learn the truth?

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1	"I too want to thank all the Rocky
2	Flats workers, and the other advocates who
3	have helped me with this petition. I hope in
4	October, when the Board comes to Denver, we'll
5	have as much time as need to explain our
6	position." Thank you very much, and we'll
7	take any questions.
8	CHAIRMAN MELIUS: Okay, thank you
9	very much, Terrie. LaVon, responses or
10	MR. RUTHERFORD: We got Terrie's
11	presentation with her, and we're going to
12	follow up on every one of the things she put
13	in there, and we'll make sure that we provide
14	a response to the Work Group as we work
15	through these issues.
16	CHAIRMAN MELIUS: What my
17	recollection is from when the Evaluation
18	Report was first presented, that you were
19	following up on the other tritium scripting
20	time frames, weren't you, or is that
21	MR. RUTHERFORD: No, not the
22	tritium stripping issue.

CHAIRMAN MELIUS: Okay, okay.

MR. HINNEFELD: Yeah, the specific phrase of "tritium stripping" I don't think was in front of us at that point.

CHAIRMAN MELIUS: Okay.

MR. RUTHERFORD: I would like to say that some of the issues, you know, recognize that we went back and we looked at the issues that one, we didn't feel had been fully vetted in the previous evaluation, and new issues that came up during our classified interviews or other interviews and our document reviews.

of the things So some that occurred in the previous evaluation haven't come back on our radar as well, I'll say. So you know, I think we've caught all the issues that I know of, and I certainly have been talking with, or following Terrie's presentations in her documents that she sends over when we look through issues, and we'll follow these up as well.

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1	CHAIRMAN MELIUS: Okay. Thank
2	you. Mark?
3	MEMBER GRIFFON: I mean I just
4	going to ask while you're there, LaVon. One
5	question, I don't know if someone can answer
6	or maybe answer before the meeting's over.
7	Terrie mentioned, I guess it's really an
8	assessment report, right? It's not a Tiger
9	Team. But this four volume report that
LO	exists, does NIOSH have this?
L1	MR. RUTHERFORD: Yeah. I'm going
L2	to say I believe we do, because I believe
L3	those were part of those that were looked at
L4	during the previous evaluation under SEC 30.
L5	I'll verify that we have them, and get you an
L6	answer before the end of the meeting.
L7	MEMBER GRIFFON: Okay, and maybe
L8	if they can be posted or somewhere where
L9	others can look at them, you know, that would
20	be useful, I think.
21	CHAIRMAN MELIUS: Any other Board
22	Members have questions or comments? I want to

indicate just in response to one of your comments, Terrie. I think everyone's trying to, doing their best to keep this moving along, and we agree.

It's a long time, and although it's -- I don't think we even are comfortable saying it's only two years, because it's something you try to resolve, but try to resolve thoroughly. But every attempt to move it along.

I think it's a part of NIOSH and everybody else involved. But we appreciate your input, and input of the petitioners. Thank you. Okay. Let's see where we are, a little bit ahead of time. Hear from somebody new now. LaVon. Is LaVon here?

SEC Petitions Update

MR. RUTHERFORD: Okay. I'm going to give the status of the upcoming SEC petitions. This is a report that we routinely do at the Board meetings, give you an idea. Gives the Board Members an idea of current

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status of existing SEC petition evaluations, what SECs do we have that have just recently qualified and 83.14s that we're working on.

It also gives them an idea that on, so they can prepare for future Work Group meetings and Advisory Board meetings.

Petitions. We are, as of July 5th, we're up to 213 petitions. We have two petitions right now that are in the qualification process, and one petition that's actually in the evaluation process. remember, the last couple of meetings, haven't had any really new petitions. have received petitions some new here recently, and I'll talk about those shortly.

Currently, there are a number of petitions that are with the Advisory Board. They have had some action taken since they were initially presented to the Advisory Board, but they have not been completely closed out. Some of these I'm thinking or hoping are going to come off the table during

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this meeting.

We've got the Fernald Feed Materials Production Center, Hanford, Pantex plant, Los Alamos National Lab, Savannah River Site, Brookhaven National Lab. Brookhaven, I believe, is actually going completely away. Baker Brothers, Joslyn Manufacturing and Supply Company.

Some of these are in various stages of final closeout and should be coming off -- this should be a much shorter list soon. We have one petition that is waiting for its initial action, and that is the Rocky Flats petition plant evaluation. Again, we did have an evaluation performed under SEC 30. But under this petition, there has been no action taken by the Board as of yet.

There are a number of petitions that we have dialed up for 83.14s. Sandia National Lab Livermore. We're actually working this 83.14 in preparation for the October Board meeting. We'll be recommending

a Class up through 1994.

Sandia National Lab Albuquerque.

This is the early years that used to be the Z

Division at LANL. We're still waiting for a

claimant for this one, to support an 83.14. We

have no claims for this period as of yet.

General Atomics, this is -- we're modifying an existing Class Definition. This was one of our old Class Definitions that identified, was very building-specific, would not have been defined this way under current practices today. However, at this time, DOL is implementing this Class, such that it would be just like it was all employees. So we have not received any claims to support modifying the Class.

Dayton Project Monsanto. There's a couple of things going on here. We need to modify the Class based on the facility designation change to a DOE facility, and then we'll also add a nine month period where operations shifted from the Dayton project to

the Mound.

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Again, we have no claim to support this at this time. We've put this on our -routinely checked for claims to support getting these moved forward, so are looking. Okay, new petitions. We actually had two petitions in the qualification process, K-25, 1993 to '97.

If you remember, the statutory SEC goes up to February 1992, and this period is just post that statutory period. We've very close to a finding on this one. The LANL petition is actually for -- this happens every once in a while. We had a petition for a period that's already covered under the SEC. It happens.

Sometimes we get individuals that have non-presumptive cancers that would not be a part of the SEC, that petitioned to get their in under the SEC, which that can't happen. So this petition will not qualify.

Then we have one petition that is

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1	in the evaluation process, and that is the
2	Kansas City plant qualified for the period
3	1949 to 1993. We had some gaps in monitoring
4	for some activities in Kansas City plant that
5	supported qualification, and we are in the
6	evaluation phase on that one.
7	That petition evaluation will not
8	be ready for the October meeting. I think
9	based on our current project schedule, it
10	would be the following meeting after that. I
11	believe that's it. Questions.
12	I tried to race through it.
13	CHAIRMAN MELIUS: I was talking to
14	the Secretary. I had to hang up.
15	MR. RUTHERFORD: You know, Josh
16	just pointed out that I failed miserably. I
17	forgot we have one other petition that is the
18	qualification phase that we just recently got
19	for Argonne National Lab East, and we are
20	reviewing that petition as well. It is in the
21	qualification phase.

CHAIRMAN MELIUS:

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You may have

1	said this, but what puzzled me when I looked
2	through the slides is why have you
3	predetermined that LANL will not qualify?
4	MR. RUTHERFORD: Okay. I did
5	answer that. It was pretty the person
6	petitioned for a period that's already covered
7	under the SEC.
8	CHAIRMAN MELIUS: Okay. I assumed
9	that, but you had accepted the petition?
10	MR. RUTHERFORD: Well, any
11	petition comes in, we go through the petition
12	process.
13	CHAIRMAN MELIUS: Alright, okay. I
14	thought you had an administrative way of
15	dealing with those also.
16	MR. RUTHERFORD: Yeah.
17	CHAIRMAN MELIUS: No, that's good
18	of you. Any other Board Members have
19	questions for LaVon? Good thing we found you
20	other work to do then.
21	MR. RUTHERFORD: Yes.
22	CHAIRMAN MELIUS: Yeah. Thank

you. Okay. It's a little bit early for a break, and we can do a break now. We can go into the Board work session. We can do that for a while. We can do that to say 3:30 or 4:00, and then take a break until 4:30, and then that will give us a break and then come back at 4:30 for the INL, sort of a longer break.

I'm not sure. We'll see how we do through the Board Work session. But in terms of Board work session, we're going to need to save some for tomorrow, because the SEC petitions are all, actionable ones are all tomorrow and Fernald, both Pantex and Fernald will be, good chance they will be letters and so forth. We'll need to talk about a little bit.

We don't have any scheduled Board work time tomorrow until late. So why don't we go through and let's see how we do. If we're doing well, try to break and, a break before the 4:30 time frame and so forth, or

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1	have a break and then come back. Is that
2	reasonable for everybody?
3	(No response.)
4	Okay. Hearing no objections,
5	we'll move forward. Why don't we start with
6	what's usually the, and I don't have it on my
7	set of this copy of the agenda, but you had
8	some dates you wanted to throw out?
9	MR. KATZ: So this just to
10	schedule further out. So presently, we're
11	scheduled the latest meetings. We have the
12	October meetings, October meeting in Denver.
13	MEMBER ROESSLER: For what, two or
14	three days?
15	MR. KATZ: I'm pretty certain it
16	will be two days, 16th and the 17th. Two
17	days.
18	MEMBER ROESSLER: Two days.
19	MR. KATZ: 16th and 17th of
20	October.
21	MEMBER ROESSLER: Okay.
22	MR. KATZ: So the 16th and 17th of

1	October, we are planning to go to Denver, and
2	then we have scheduled December 9th at 11:00
3	a.m. for a teleconference, and then the next
4	in-person meeting is January 28th through 30th
5	we have blocked out for that, January 28th
6	through 30th.
7	CHAIRMAN MELIUS: I think Brad's
8	invited us back.
9	MR. KATZ: For the January
10	meeting. That would be lovely. Okay, 28th
11	through 30th.
12	CHAIRMAN MELIUS: Yeah. Most
13	likely the Tuesday and Wednesday.
14	MR. KATZ: Yeah, most likely 28th
15	and 29th.
16	CHAIRMAN MELIUS: Or we could do
17	the 29th, whatever people's preferences are.
18	MEMBER ROESSLER: Okay, I got it.
19	MR. KATZ: No. We can talk about
20	that if we have ideas, but
21	CHAIRMAN MELIUS: Yeah. I think
22	we're looking for suggestions. But at this

1	point in terms of what
2	MEMBER ROESSLER: Kansas City.
3	CHAIRMAN MELIUS: Kansas City.
4	MEMBER ROESSLER: I was just going
5	to suggest that.
6	CHAIRMAN MELIUS: I was reading
7	your mind. Many of you have never been there,
8	and I didn't listen to all of the, LaVon's
9	thing, but my understanding is that the I
LO	think I heard that the SEC report probably
11	would not be ready in October?
L2	MR. KATZ: Oh yeah. Not in
L3	October, but
L4	CHAIRMAN MELIUS: Yeah. So it
L5	would be ready for the January meeting.
L6	MR. KATZ: Correct. So that's one
L7	option.
L8	CHAIRMAN MELIUS: Okay.
L9	MR. KATZ: Augusta is another
20	option, if SRS is ready by then. I couldn't
21	hear.
22	CHAIRMAN MELIUS: Rio. Yeah, I'm

1	already working on, since Joyce Lipsztein's so
2	involved, that we should do at least do a Work
3	Group meeting. I told him I would displace
4	Mark from the SRS Work Group.
5	MR. KATZ: Not yet.
6	CHAIRMAN MELIUS: Somehow, I think
7	we would be reading about ourselves in USA
8	Today if that occurred.
9	MR. KATZ: I don't know. There's
10	Pinellas. Do we expect progress in this time
11	frame for Pinellas?
12	Okay. So that's a no from the
13	program for Pinellas in that time frame.
14	Sandia is another arena.
1 -	
15	DR. NETON: I think Pinellas might
16	DR. NETON: I think Pinellas might be ready.
16	be ready.
16 17	be ready. MR. KATZ: Oh.
16 17 18	be ready. MR. KATZ: Oh. DR. NETON: It's very close to
16 17 18	be ready. MR. KATZ: Oh. DR. NETON: It's very close to closure.
16 17 18 19 20	be ready. MR. KATZ: Oh. DR. NETON: It's very close to closure. MR. KATZ: Okay. So Pinellas

1	action we're working on is Sandia Livermore.
2	So that one, I think, will be done before.
3	MR. KATZ: Before then.
4	CHAIRMAN MELIUS: I'd rather have
5	Rio. Kansas City, I mean we've never been
6	there, and I think that would be first
7	priority. I'll say weather's a factor, but
8	weather's a factor getting anywhere there.
9	Yeah. We'll probably hit the blizzard of
10	yeah. So let's tentatively do Kansas City,
11	and then
12	MR. KATZ: Okay. It's Kansas City
13	in January. Okay. So then now we're
14	scheduling out beyond that, and the right
15	weeks for a teleconference beyond that would
16	be the week of March 19th or March 26th. I
17	don't know if I chose the Wednesdays as date
18	marks for those weeks or something else.
19	MEMBER ZIEMER: What were the
20	January dates again?
21	
	MR. KATZ: The January dates are,

1	29th.
2	MEMBER ZIEMER: Thank you.
3	MR. KATZ: So is the 19th a Monday
4	or a Wednesday? Okay. So how does March 19th
5	work for people? For a teleconference, that
6	would be 11:00 a.m. Eastern Time normally.
7	Does that work for everyone, and on the line
8	too? Paul, David?
9	MEMBER ZIEMER: That's okay with
10	me, Ziemer.
11	MR. KATZ: Okay, great. Okay. So
12	the 19th it is, and then the next face to face
13	meeting, approximately the week of April 28th,
14	May 5th, May 12 th . I said April 28th or May
15	5th, or as far out as May 12th, and then we're
16	getting pretty far after that.
17	Anybody have any trouble with the
18	week of April 28th, or with one part of the
19	week or the other?
20	CHAIRMAN MELIUS: What are the
21	holidays? They're not in my calendar year, so
22	Microsoft is falling down again.

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1	MR. KATZ: Yeah. Perhaps none. So
2	do you want to go the 29th and 30th?
3	MEMBER ROESSLER: Yeah.
4	CHAIRMAN MELIUS: Yeah.
5	MR. KATZ: Okay. Let's do that.
6	29th and 30th of April.
7	CHAIRMAN MELIUS: Place to be
8	MR. KATZ: Place to be determined.
9	No, that's a Tuesday and Wednesday.
10	MEMBER ANDERSON: April 29th?
11	MR. KATZ: April 29th and 30th.
12	MEMBER ANDERSON: Oh, 29th, okay.
13	MR. KATZ: And 30th. That way,
14	people aren't traveling on Sunday.
15	MEMBER ANDERSON: Oh, that's good.
16	I just put 28th in my mind.
17	MR. KATZ: Week of, yeah.
18	CHAIRMAN MELIUS: And let's keep
19	in mind, I guess we have Sandia, Pinellas and
20	Livermore as possible locations. For those.
21	So let's do, start with Work Groups and
22	Subcommittees, and recently, Dave Kotelchuck

2	Subcommittee, and you're on.
3	Dose Reconstruction Subcommittee
4	MEMBER KOTELCHUCK: We have a
5	conference call meeting, actually what do they
6	call it Live Performance on August 7th, and
7	we're basically going through the Sets 10
8	through 13. We have LANL, we have SRS and I
9	forget, one more, and then we're going to be
10	selecting for, selecting cases for a new set,
11	which will be either 17 or 18.
12	There's some question. I think
13	17, something else coming up that we're going
14	to call 17, yes.
15	MR. KATZ: It's just the blind
16	reviews for 17, so we've moved to 18.
17	MEMBER KOTELCHUCK: That's right,
18	that's right.
19	MR. KATZ: To accommodate that.
20	MEMBER KOTELCHUCK: So it will be
21	18. Okay. That's it. Moving right along.
22	CHAIRMAN MELIUS: Okay. Any
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took over as chair of the Dose Reconstruction

questions, comments? Okay. So where are we with the blind reviews, in terms of --

MR. HINNEFELD: It's been a bit of a struggle. The blind reviews intend to use the existing ORAU dose reconstruction tools, which allow the dose, you know, the dose reconstruction makes a certain number of choices when they use those tools, and provide those tools to the SC&A reviewer to do the cases essentially so the arithmetic all comes out. Its choices are consistent.

CHAIRMAN MELIUS: Yeah.

MR. HINNEFELD: And getting that going has been a bit of an ordeal, but the last I saw is they are not accessible to Doug, the reviewer, and the data input files, you know, the pre, the already coded, you know, spreadsheets essentially of a person's dose record have been now found, and should have been made available to Doug, will be made available this week.

We've notified Doug. I think we

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have those files now. What folder, where you want us to put them, so you can get them and use them on the tools. This has been a little bit of a complicated computer security issue from our point.

That was the complication, getting the tools in a place where Doug can get to them through our system, because normally they run on the ORAU system, and getting them to where they would run in what believe it or not is called the demilitarized zone, in IT tech speak, and I don't know exactly what that means.

So that's the last report. The last message I saw on this was Grady sent a message to Doug that says I have those data input files, so Doug doesn't have to key in the data, and where would you like them, and that's the last message I saw. But once he has those, I think things should relatively quick.

CHAIRMAN MELIUS: So we've

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demilitarized the -- will we allow an incursion into the -- there's got to be some jargon there.

MR. HINNEFELD: It is a virtual area between our servers and ORAU's server, and I can't do any better than that.

CHAIRMAN MELIUS: Anybody have questions on that? If you do, too bad. Okay. Procedures, Work Group, or excuse me, Procedures Subcommittee, which is going to have some presentations tomorrow. But in addition to those.

Procedures Subcommittee

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MEMBER MUNN: Yeah. We met in April, April 25th, one of the first Live Meeting calls, which I know the order finds to useful, this be very and particular Subcommittee chair finds to be disastrous. But we are making a few adjustments to the Board Review System, which when I say "we," actually NIOSH and the folks who actually do the hard work with it, are tweaking it a little bit,

making it we hope a little bit more user friendly than it already is.

It's working very well for the Subcommittee. We had a large number of items on our agenda, especially now that we are really into the PERs. We have probably seven or eight of them actively involved right now, either being resolved in NIOSH or with SC&A.

We had reports on PER-11, 30, 14, 17, 44. I don't know that the numbers mean anything to you. I have to go back and look at each one of them to identify where they are. But they've covered the wide range of sites and a wide range of issues.

We have been adding, for the first time talking about how best to handle the overarching issues, which we've opted to track through the Subcommittee. But they're going to have to be handled, because of their nature, and not being site-specific, they're going to be handled differently than most of our findings.

1	We had quite a bit of a discussion
2	about that, and I think we've come to some
3	resolution on some of the simple, basic
4	elementary things. But we'll be working on
5	that a great deal, I think, from time to time.
6	We had some responses from OTIB-
7	55. We had two PERs, I believe, from the
8	Hanford site, and those I believe have been
9	transferred to the Work Group for resolution.
10	Status reports on several of the PERs, and we
11	had a status report on the revisions to OTIB-
12	54.
13	I believe our next meeting is
14	going to occur day after tomorrow, if we're
15	still present and functioning there, and
16	that's, I believe, yep. Unless someone has
17	some questions.
18	CHAIRMAN MELIUS: So this will be
19	live Live Meeting.
20	MEMBER MUNN: This will be a real
21	live meeting, not a digital live meeting.
22	CHAIRMAN MELIUS: A virtual.

1	MEMBER MUNN: I much prefer real
2	live meetings to virtual live meetings.
3	CHAIRMAN MELIUS: Yeah, in a
4	demilitarized zone.
5	MEMBER MUNN: Yes, uh-huh, with
6	both computers that are necessary to
7	accomplish this, yeah.
8	CHAIRMAN MELIUS: Any questions
9	for Wanda? Okay. Now we'll turn to the Work
10	Groups. Anybody want to volunteer? Yeah, go
11	ahead. Now we'll just start going through
12	alphabetically, but I figure, yeah.
13	MEMBER ROESSLER: Have a little
14	variety.
15	CHAIRMAN MELIUS: Variety, yeah.
16	We've been doing the alphabet for how many
17	years, right?
18	ORNL Work Group
19	MEMBER ROESSLER: Right. Okay.
20	I'll report on ORNL Work Group, and I do have
21	progress to report. Tim Taulbee, who's the
22	DCAS lead on this site, is in Alaska right

now. But he sent me an update this week. I tried to send it out to the rest of the Work Group Members, and I didn't get your email address right, but I'll give you a copy of it.

But recall, Tim as you may presented the DCAS Petition Evaluation Report to us at a meeting in Denver last September. You might also recall that this is a complex site. The petition covers the period from June 17th, 1943 to July 31st, 1955. So historical information there's lot of involved.

This is the X-10, site which involved the historic graphite reactors, some other very unique reactors, and many research labs. In September, DCAS reserved the exotic radionuclide portion for further follow-up, and Tim reports that since them they have made significant progress, but it is slower than anticipated due to difficulty in obtaining and assessing data from DOE.

He says they now have a good

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handle on the radionuclide production, that ORNL. They've identified 254 radionuclides that were produced there over the years. I think just that number itself should tell people that that's a pretty interesting site to evaluate.

He said they will do a triage on these radionuclides. I'm not sure if that's quite the right word, but to assess their exposure potential. Some of them were very small in quantity. Some had very short half - lives. Some were encapsulated when they were used, and some have low dose conversion factors.

However, some have significant exposure in internal dose potential. So they're looking at that. They're also doing validation and verification of the bioassay database that they reported on in September. So they're targeting the October Board meeting for an Evaluation Report addendum, but are not 100 percent sure they can do that. So stay

tuned. Any questions?

CHAIRMAN MELIUS: Questions for Gen? Okay. Okay, very good. Any other volunteers? Then I'll start calling on people. Yeah. We'll see who's been slacking and who's been into that. Brookhaven is reporting, Fernald is reporting.

Hanford, Hanford has been working but has not been meeting. Arjun, correct me if I'm wrong here, but we're actually updating the matrix, catching up on -- there have been some more recent interviews and so forth, pulling information together into the matrix on where we need to go from here.

So we hopefully will have a Work Group meeting between now and the next meeting, and see where we would need to focus in in terms of further SEC evaluation at that site.

DR. MAKHIJANI: Yeah, that's more or less right. Dr. Melius, we sent you a report with a number of findings in April, and

1	at that time, there were some loose threads
2	regarding neptunium especially that needed
3	interviews, and we have been struggling to get
4	them organized.
5	Finally, I think, they will be
6	done in the week of July 29th, and we will
7	have completed our work. So we may issue a
8	supplement to the report. So we're very, very
9	close to done.
10	CHAIRMAN MELIUS: So any questions
11	on Hanford? Okay. Idaho?
12	Idaho Work Group
13	MEMBER SCHOFIELD: Idaho. There
14	are some revisions to be done to the TBD and
15	the recent White Papers.
16	CHAIRMAN MELIUS: Mic, mic. Use
17	your microphone.
18	MEMBER SCHOFIELD: Sorry. There's
19	some revisions to be looked at in the TBD,
20	both by SC&A and NIOSH. There's also some
21	White Papers hopefully will be out supposedly
22	the latter part of October. So without a

1	current SEC qualified petition, we're still
2	kind of setting in the water, not moving much.
3	CHAIRMAN MELIUS: Dr. Ziemer, if
4	you're on the line, Lawrence Berkeley.
5	Lawrence Berkeley Work Group
6	MEMBER ZIEMER: Yes, I'm here.
7	Lawrence Berkeley?
8	CHAIRMAN MELIUS: Yes.
9	MEMBER ZIEMER: Right. Lawrence
10	Berkeley, currently NIOSH is reviewing four
11	White Papers from SC&A that were generated
12	following the initial meeting of that Work
13	Group, which goes back a little over a year,
14	and those responses are still under
15	preparation. We have received in late May, I
16	think May 31st actually, one White Paper from
17	NIOSH on thorium.
18	So the Work Group has that in
19	hand. We're awaiting the responses for the
20	other four White Papers. Most of those are in
21	draft status. I understand from Dr. Lara

Hughes that pending resolution of some of the

1	questions raised on their internal review, and
2	some additional data captures, they will be
3	passing this information on to the Work Group,
4	at which point we can schedule a meeting. But
5	we're still awaiting those. So that's our
6	current status.
7	CHAIRMAN MELIUS: Thank you, Paul.
8	NIOSH, any
9	MEMBER ZIEMER: Incidentally, if
10	we're talking about possibly a meeting in the
11	Lawrence Livermore area in April, hopefully we
12	would have something more substantial on
13	Lawrence Berkeley at that time as well.
14	CHAIRMAN MELIUS: That makes
15	sense. Stu or I guess I'm reading your
16	report here on Lawrence Berkeley and do you
17	have some
18	DR. NETON: I'm the hold up on
19	Lawrence Berkeley, so I guess I should take
20	responsibility. The internal review process,
21	I looked at some of the comments that were
22	made, and it's my opinion Lawrence Berkeley

1	has similar issues to what other cyclotron-
2	type facilities have, that is a slew of
3	exotic-type radionuclides that were handled in
4	various degrees and quantities.
5	So I just wanted to make sure that
6	we button up those issues before we proceed.
7	That's where we are, and that's where the idea
8	of possibly additional data captures are
9	necessary. I'm not sure. I raised the
10	question, and people are looking at the data
11	that we currently have.
12	We have a lot of data we captured
13	there, and I just asked folks to go back and
14	look through what we have, and make sure that
15	we can put some brackets around some of these
16	exotics. I'd feel a little better moving
17	forward then.
18	CHAIRMAN MELIUS: Good. Thanks,
19	Jim on that, and Paul. Kansas City, I think we
20	Josie, do you want to add anything to what

MEMBER BEACH:

has been said?

21

22

No, I don't have

anything to add at this time.

CHAIRMAN MELIUS: LANL, Mark.

LANL Work Group

MEMBER GRIFFON: Yeah, I'm going to -- it's sort of like Hanford. We've been -- there's work going on. We haven't had a meeting in a while. But I'm going to ask if Joe, if you have anything significant to add. I mean we need to at some point schedule a meeting. But there's been, the Work Group hasn't met in quite some time, and nothing on the horizon as far as I know.

MR. RUTHERFORD: Well, I can a little bit to that. We've provided a questionnaire to the site on some, trying to get a better feel for the end date of the existing SEC from the end date in 1994, up beyond when, you know, does the site have a good handle on the program. Do we know that they were looking at the exotic radionuclides, if they had good methods.

We provided a questionnaire to the

1	site some time back, to answer some questions.
2	They responded generally to those questions
3	back to us, but it just brought on additional
4	questions or additional clarification that we
5	needed.
6	We sent that back to the site. The
7	sequestration kind of held that up a little
8	bit. I think we expect that response back
9	from them very soon though, within the next
10	week or two.
11	CHAIRMAN MELIUS: According to
12	your schedule, you were expecting it back last
13	week.
14	MR. RUTHERFORD: Yeah.
15	CHAIRMAN MELIUS: Yeah. So we're
16	close enough for that. Okay. Mound.
17	Mound Work Group
18	MEMBER BEACH: So far Mound, we're
19	complete. We have completed all of our SEC
20	issues. We're currently awaiting some
21	response from NIOSH on some Site Profile
22	issues, and just heard from Jim earlier today.

1	Don't really have a time frame on those, but
2	they're aware that they're due.
3	CHAIRMAN MELIUS: Questions for
4	Josie? I will add somebody, some Board
5	Members asked me at the break about the Mound
6	emails and where they're found. If you go to
7	the ANWAG block, A-N-W-A-G. If you Google
8	that, you will find on that blog a series of
9	postings on Mound, from what, about two months
10	ago, something like that.
11	MEMBER BEACH: Yeah. I can also
12	send a link to anybody that wants it, because
13	there's actually two spots that they're
14	available.
15	CHAIRMAN MELIUS: Yeah, you can do
16	that. So they're there, if you're interested
17	in reading those. Okay. Nevada Test Site,
18	Brad.
19	Nevada Test Site Work Group
20	MEMBER CLAWSON: We haven't met
21	right yet. We've got, we've got SC&A has
22	gone through and reevaluated the Site Matrix

1	and we're just in the process. We should be
2	setting up a Work Group to be able to go
3	through those in the next little while.
4	MR. HINNEFELD: I'm trying to get
5	my computer going.
6	CHAIRMAN MELIUS: Yeah. You have
7	something revealing to say to us, but Pantex
8	we're going to hear about tomorrow. Pinellas.
9	Pinellas Work Group
10	MEMBER SCHOFIELD: The main
11	outstanding issue of Pinellas is still dealing
12	with the tritium issue, and if we get all
13	those settled with Mound on how to deal with
14	that, that will probably help us close out
15	Pinellas.
16	DR. NETON: Just a slight
17	correction. It's really a tritide issue at
18	Pinellas that's holding it up, and we are
19	interviewing or have recently interviewed, I
20	hope, a health physicist at Pinellas that can
21	inform us, maybe in some more detail, how they

actually monitor for tritides, because there

1	were some there, and there were spills.
2	There's references to a
3	Bremsstrahlung counter, which I know what
4	Bremsstrahlung is, but I've never seen a field
5	instrument called a Bremsstrahlung counter. So
6	we want to see what's going on there. We
7	approached it. We had listed in the Site
8	Profile had some flaws in it, because they had
9	filtered the tritium solutions prior to
10	counting them. So clearly the tritides
11	weren't in the solution.
12	So we're trying to shore that up a
13	bit more, and Phil's right. That's the only
14	outstanding issue that I'm aware of at
15	Pinellas.
16	CHAIRMAN MELIUS: Thanks for the
17	update, Jim. You're on, Portsmouth-Paducah-
18	K25, Phil.
19	Portsmouth-Paducah-K25 Work Group
20	MEMBER SCHOFIELD: Okay.
21	Basically, Paducah is closed on the they
22	are looking at currently the neutron/photon

1	ratios, because all three facilities had
2	cylinders where they stored highly enriched
3	uranium. So they're trying to get a handle on
4	those ratios, so that we can close out that
5	issue, and then we're just I think that
6	will pretty much close us out on the gaseous
7	diffusion plant.
8	CHAIRMAN MELIUS: Rocky. We've
9	done Sandia. Dr. Lemen isn't here. I don't
10	believe there's been I think there's
11	ongoing activity in the NIOSH end on this one.
12	LaVon, you want to
13	MR. RUTHERFORD: You want
14	Albuquerque or do you want Livermore?
15	CHAIRMAN MELIUS: Both.
16	MR. RUTHERFORD: Okay, Livermore -
17	_
18	CHAIRMAN MELIUS: I said Sandia.
19	Albuquerque and Livermore Work Groups
20	MR. RUTHERFORD: There you go.
21	Livermore, as I mentioned, that we are working
22	on an 83.14 for Livermore. It's the same

issues that were at Albuquerque. Livermore, the radiological control program kind of mirrored Albuquerque's, and so ultimately we'll be adding Class, very similar to what was -- or recommending a Class very similar to what was done at Albuquerque.

That is on schedule for a presentation at the October Board meeting. We also have some additional, it's still -- the open period at Albuquerque and Livermore will be the post-'94 period. We had some onsite visits scheduled at this time for August, to do some additional interviews, to see if we can close out some of the questions.

Similar questions that we have at LANL that we'll have at Albuquerque-Livermore, basically updating, understanding their program at that time. Also understanding their availability of records. Sandia is also dealing with at this time a backlog of claims.

So we can't, we've been kind of pushed off our going to the site to work on

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these issues, and so they can use their resources to get these backlog of claims taken care of. So that's pretty much where we stand at this time with this.

CHAIRMAN MELIUS: Okay, thank you.

Questions for LaVon? Yeah, okay. Okay. Santa
Susana.

Santa Susana Work Group

MEMBER SCHOFIELD: I'll have to give DCAS a compliment here. They've been doing a tremendous amount of work on revisions of the TBD, and also they've received, I don't remember what it was, a boatload of documents, mostly exposure records and stuff. They've been having it entered by hand.

Presently, there's a coworker study, hopefully will be done on the internal. That's not due out until the middle of February next year, and the coworker study will be due out end of November, quote-unquote.

CHAIRMAN MELIUS: Good. Thank

1	you, Phil. Questions? Okay, you're on.
2	Savannah River Work Group
3	MEMBER GRIFFON: Savannah River
4	is, there's some significant progress that's
5	gone on. SC&A, I just talked to Arjun earlier
6	today. They've completed a review of the
7	neptunium model, and have several findings
8	ready to bring that back to the Work Group.
9	Also making quite a bit of
10	progress on the thorium issue, and we're just
11	trying to figure out the timing, best timing
12	for a Work Group meeting. But if we have
13	those two significant issues, it's probably,
14	it will probably good to schedule something in
15	the near future on those. So that's sort of
16	an update.
17	CHAIRMAN MELIUS: Any questions or
18	that? Just going back to the issue with the
19	DOE and the site and so forth, is that holding
20	up the Work Group at this point?
21	MEMBER GRIFFON: Arjun, you want

to speak to the site access issues?

DR. MAKHIJANI: Well you know, as Greg Lewis explained in the morning, we've had some difficulties, and DOE has been working hard with us to make some progress. Joe is going to get access to review the documents from his government computer, without going to Savannah River.

So we did some interviews. did some interviews recommended by CPWR, Center for Protection of Worker Rights, and we have a number of other interviews scheduled. The schedule for review, hope the we classified document searches will be done But it's difficult, you know. There's soon. a fair amount of sorting out.

However, since NIOSH already has put a very amount of analytical information, coworker models, data, compiled a lot of data, we're able to do quite a lot of work, as you know. Besides the neptunium and thorium, and the thorium is done, the trivalent actinides will also be largely done. So you'll be able

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to review those at the same time.

CHAIRMAN MELIUS: Thanks. I do want to bring up the issue, since the sort of site and data access issue is ongoing, and may be getting resolved, may not. But should it be necessary or helpful for the Board to write a letter to appropriate parties involved at DOE in this, that we not have to wait until a Board meeting.

So what I would plan on doing would be to draft up a letter and I will circulate it to the Board for comment or input, obviously work with the Work Group. But it would be something I'd rather not have to wait until the next Board call or Board meeting to do that. If no one objects to that, I think that would be the procedure. Yes, Wanda.

MEMBER MUNN: Have we in fact decided that we're going to send such a letter?

CHAIRMAN MELIUS: No, we have not

1	decided that we will send, should it be
2	necessary if things are not making significant
3	
4	MEMBER MUNN: Oh. I missed the
5	"should it be necessary" phrase. Thank you.
6	CHAIRMAN MELIUS: Yeah. Science
7	Issues, David, are you on the line? David
8	Richardson? David Richardson are you on line
9	and off mute? Okay. I have been informed
10	that Jim Neton is the controller of what's
11	happening on the Science Issues Work Group.
12	Science Issues Work Group
13	DR. NETON: Yes, Dr. Melius. I
14	did send a report to Dr. Richardson. If he's
15	on, I'd be more than happy to have him present
16	it. But it's very short. The dose rate,
17	effectiveness factor, so-called DDREF tome,
18	the document written by us, by Senes a while
19	back, has been sent out for external and
20	internal review.
21	We solicited reviews from five
22	external experts and two experts from within

NIOSH. As of this morning, we have all six reviews back. We were missing one review for the complete package. Once we get that final review, we will pass that on to Senes for their consideration and response to the comments we've received.

But it's been a while. But it's a large document and we had to cajole some people to get it to us in a timely manner. But they're there, and I think the one remaining review will surface fairly soon.

CHAIRMAN MELIUS: Maybe the Board chair needs to write a letter to the recalcitrant party.

DR. NETON: They do this not for much money. We can only offer a very meager honorarium. So but I'm happy with the reviews that we got and the panel that we selected, and that's public knowledge. It should be public knowledge. We're going to de-identify the actual reviews themselves, but we will publish the qualifications of the individual

1	reviewers.
2	CHAIRMAN MELIUS: Yeah, I found
3	that those of us redoing journal reviews get,
4	especially some of the electronic ones, where
5	you get varying levels of email reminders and
6	then threats, you know. Then I think at some
7	level it's a public shaming or something that
8	goes on if your review isn't in, and how you
9	personally are holding up the progress of
10	science and failing to save the world and so
11	forth.
12	MR. HINNEFELD: That's just my
13	life at a Board meeting, Jim.
14	MEMBER ROESSLER: Jim, can you
15	tell us who the reviewers are?
16	DR. NETON: Yes, I can.
17	CHAIRMAN MELIUS: Just say the
18	name of the recalcitrant reviewer really
19	slowly.
20	DR. NETON: I'll just mention the
21	reviewers. Dale Preston from Hirosoft; Rick
22	Hornung, University of Cincinnati; Bill

1	Morgan, PNNL; John Boice, representing NCRP in
2	this particular instance; and the fifth one,
3	this is terrible. I'll think of it in a
4	second. Oh yeah, Jerry Puskin from the
5	Environmental Protection Agency.
6	Yeah. I thought it was a fairly
7	good list, and to get all of them to respond
8	was great. Internally, Doug Daniels and Mary
9	Schubauer-Berigan of the DSHEFS Division are
10	also reviewing, because they're fairly well
11	familiar and interested in this area.
12	CHAIRMAN MELIUS: Just let us
13	know. We'll write the letter. Paul Ziemer,
14	if you're on the line, for TBD-6000.
15	TBD-6000 Work Group
16	MEMBER ZIEMER: Right. Can you
17	hear me?
18	CHAIRMAN MELIUS: Yes, we can.
19	MEMBER ZIEMER: Okay, yeah. We're
20	dealing with actually four different sites, so
21	let me report briefly on each of those.
22	General Steel Industries first.

NIOSH is preparing the final details on how dose is demodified for all the components. So it's both the source operational and the residual periods, and then following that, SC&A will have a chance for final review prior to the next Work Group meeting. We haven't scheduled that meeting yet. But that should occur fairly soon.

Also, we have yet also the complete closure of all the findings matrix and that is the other item on the agenda for General Steel.

For Joslyn Manufacturing, I'11 just remind you. Currently, there is an SEC through 1947, and we're reviewing the remainder of the operational years. DCAS responses to the SC&A are expected I believe by the end of July. At least that was the last date I heard, and then will be reviewed by the Work Group at its next meeting.

Simonds Saw and Steel, and a reminder again. There is an SC&A, not an

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SC&A, an SEC for Simonds Saw and Steel already. I'm trying to remember. I think it was '48 to '55 or something like that. I don't have that right before me.

But in any event, we're focusing on the TBD itself and the response to some of the SC&A findings. NIOSH has agreed to some revisions in the TBD, and those are currently underway. Then NIOSH is also reevaluating the urinalysis data for internal doses, and also the methods for dose reconstruction in the residual period. So that is going on to supplement what we already have for the existing SEC.

Then Baker Brothers, the Work Group has voted to recommend that the SEC Class not be granted for the residual period, and we'll be reporting on that in detail tomorrow. So those are our four areas that we're looking at.

CHAIRMAN MELIUS: Thank you, and Paul, you put all the other Work Groups to

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1	shame with your productivity, going through
2	all these sites. But no, it's been a lot of
3	work and it is appreciated by that Work Group
4	for that.
5	MEMBER ZIEMER: Thank you.
6	CHAIRMAN MELIUS: Questions for
7	Paul? If not, Henry.
8	DuPont Deepwater Work Group
9	MEMBER ANDERSON: We got the SC&A
10	response to the NIOSH commentary findings on
11	DuPont Deepwater the first part of June. So
12	the committee's going to be looking at that,
13	and then we'll hopefully have a call to close
14	out or at least discuss DuPont again, and
15	hopefully we'll have something by the October
16	meeting.
17	CHAIRMAN MELIUS: Questions on
18	okay. Weldon Springs. Dick Lemen is yes,
19	John.
20	MR. STIVER: Yes. This is John
21	Stiver. I just wanted to kind of expand on
22	what Andy said. We do have a couple of sites.

1	I think there's about three of them where we
2	have recent work products that would fall
3	under the URAWE Work Group. Whether it be
4	NUMAC or General Atomics. So we might want to
5	consider bringing those in.
6	CHAIRMAN MELIUS: We'll take that
7	
8	MEMBER ANDERSON: If they're
9	assigned to us, we'll take it on. If they're
10	assigned to us, we'll take it on. Yeah,
11	right. Yeah, I know. We could add that to
12	our teleconference. I don't think the DuPont
13	will take too long.
14	CHAIRMAN MELIUS: Weldon Springs?
15	I don't think there's much. Dr. Lemen isn't
16	here. Okay, and last but not least, Worker
17	Outreach.
18	Worker Outreach Work Group
19	MEMBER BEACH: Okay. Not too much
20	new to report. SC&A did deliver the
21	evaluation for LANL to NIOSH, and we did make
22	DCAS' work list, but there's no date

associated when they are going to have that review completed at this time.

MR. HINNEFELD: Yeah. I will say here that that is a bit of a victim of sequestration and the resources available. So it's going to take us some while to get some people free to do work on that.

MEMBER BEACH: Yeah. I kind of expected that. Once we do get that back from NIOSH, then SC&A will go back and finish up the report and send out the finished version. So to be continued.

CHAIRMAN MELIUS: Thank you, and I again, I think it is fair to remind that sequestration has taken some toll on what we do overall in this program. So we will have to decide, and it's also one of the reasons we've hesitated. I know we've talked about some other Work Groups and we have products and so forth out there that could be reviewed.

But at the same time, we have to

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1	keep in mind some of the resource limitations,
2	in terms of making these assignments as we go
3	along here, and we need to consult with NIOSH
4	about making sure that those resources are
5	appropriate.
6	Okay. So that completes our Work
7	Group update, and unless I'm mistaken, we have
8	one more quick thing to do, and I just have to
9	find the right file here.
10	Yeah. Okay. All of you have
11	received, and you've probably all memorized
12	the comments from the the file that had the
13	comments from the last Board meeting.
14	There's two files. One's a
15	spreadsheet that summarizes the comments and
16	the response, and has that categorization code
17	that none of us can ever remember. But
18	usually someone brings it to mind if it's
19	important.
20	Then there's a second, much longer
21	file that actually has the transcripts
22	pertinent to those comments, should we have

questions or recollections of -- what went on is a little different from what's being reported here.

So I will go through these briefly. I will try to group them as we go through. There's only a small number, I believe, what is it, 17 from the last comment. So if you want to sort of read along with me and so forth.

Most of the first set are from [Identifying information redacted], who is commenting on the Savannah River Site, and had a number of questions and comments on some of the methods that were being used by NIOSH and addressing those and so forth.

I think everything looks like it's referred properly and so forth in this comments. Yeah, there's two from him. Dr. Ringen also made a number of comments relative to the SRS petition, and again, I think these are all, for the most part referred to the Work Group or the Board, addressing things

that are in progress in terms of the evaluation of that site.

There's a comment from another person there, Comments 9, 10 and 11, again related to the Savannah River Site and SEC. Again, I think these are all straightforward in terms of how they were handled and so forth and who responded.

Comment No. 12 is basically just someone indicating they supported the petition. There's a comment from [Identifying information redacted] regarding General Steel Industries, and had some issues about the process for his, how the petition was being communicated to the -- results of the petition review is being communicated by the NIOSH Director and the Secretary.

I think those have all been clarified in the response. I'm aware of at least one direct response from Ted on that, and others within NIOSH, and set of questions from Terrie Barrie, comments from Terrie

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1	Barrie pertinent to the set of these relate
2	to Hooker Electrochemical in Rocky Flats. Some
3	of them were related to another set of emails
4	and the FOI process.
5	Again, I think pretty
6	straightforward, in terms of their responses.
7	Finally, there were comments from [Identifying
8	information redacted], both some general
9	comments about individual dose reconstruction
10	and then raising several issues about the
11	Pinellas SEC petition and dose reconstruction
12	at Pinellas.
13	Again, I think these were for the
14	most part fairly general and addressed pretty
15	directly. So anybody have comments or
16	questions on those, or the nature of those
17	responses?
18	(No response.)
19	CHAIRMAN MELIUS: If you haven't
20	had a chance to go through in detail, or wish
21	to, we can also talk about this tomorrow

briefly, if you want to raise issues then.

1	not, if everyone feels ready, I think we need
2	a motion to accept these as an order. Is that
3	how we do this? No, we don't do anything.
4	Okay.
5	MEMBER ANDERSON: Just consider it
6	done.
7	CHAIRMAN MELIUS: Consider it
8	done, good.
9	MEMBER ANDERSON: No. It's a
10	useful process and good exercise, and the
11	summary is nice, so we can go through it
12	quickly.
13	CHAIRMAN MELIUS: Yeah, yeah, and
14	I will compliment I'm not sure who does all
15	this work. Yeah, but it's
16	MEMBER ANDERSON: It's a lot of
17	work.
18	MR. HINNEFELD: It's generally
19	done by our Outreach contractor, ATL, and then
20	we they collect them, and then we provide
21	responses.
22	CHAIRMAN MELIUS: Yeah, good.

1	MEMBER MUNN: Well that's really
2	reassuring them check them over. One forgets
3	from one meeting to the next, and then just
4	being able to look at them. That's what we
5	asked for, to be reassured that they were
6	being addressed. It's very well done.
7	CHAIRMAN MELIUS: Yeah, I agree.
8	Good, okay. I think that completes our, any
9	Board Work Session business. Is there
10	anything else? Ted?
11	MR. KATZ: I don't think so, I
12	don't think so. I don't have any
13	correspondence.
14	CHAIRMAN MELIUS: Right. So it is
15	3:15. If we can reconvene at 4:30, and we'll
16	start with an update on INL, and then we'll go
17	into the public comment period. So we'll
18	stand adjourned or we're on break until 4:30.
19	(Whereupon, the above-entitled
20	matter went off the record at 3:18 p.m. and
21	resumed at 4:54 p.m.)
22	CHAIRMAN MELIUS: We'll reconvene

the meeting. Welcome back, LaVon.

MR. RUTHERFORD: Thank you.

CHAIRMAN MELIUS: Glad you're still here, and LaVon will give an update on the -- I know.

INL Site Profile Revision Update

MR. RUTHERFORD: Alright. I'm going to give a little update on the INL status and Site Profile, what issues we're working on, and when we expect to be complete. For background, SC&A conducted a Site Profile Review, and identified roughly 38 issues from the initial Site Profile.

Since that review, some of the documents have been updated. Because they were updated, NIOSH and SC&A both wanted to go back to review the existing or review the issues, to determine if all the issues were still applicable. Of the 38 issues, ten are closed. NIOSH is working on 11 issues. SC&A is reviewing 22 issues, six in conjunction with NIOSH.

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Last year, we actually went out to -- out here, and went did a data capture, data review, April, May and June of last year, and identified а number of documents for We received the last of those capturing. documents in April of this year, and ORAU completed loading those documents into the Site Research Database in June.

Working on a number, a couple of White Papers to address these issues, or some of the issues that were identified. Working on a White Paper investigation of the NTA film dosimeter limits of detection being used for INL dose reconstruction. We expect that delivery to the Work Group later on this month.

In fact, it's under review right now. It's on Tim Taulbee's desk, as soon as he gets back from Alaska, to finish that review. Working on a White Paper on INL Environmental Monitoring. We expect a delivery to the Work Group scheduled for

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October of this year.

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Hot Particle issue, delivery will late, late this year, and be an issue with aircraft associated the nuclear propulsion, and we expect that delivery to the Work Group in late this year. We've also been working on a coworker model. We started working on that in June of last year.

It was a large amount of data that was actually -- it was loaded, and we went back to review that data, did some QA work on that data, and in addition, we've been adding data from the April 2013 data capture as well, and we expect to have the QA analysis done on the data in late 2013, late this year.

Again as I said, we did some additional data entry as well. The schedule for completing the model will be actually lined out after the data entry and the QA efforts are complete. Once we've done the coworker model, we've completed the coworker model. We've addressed SC&A issues.

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2	Site Profiles will be updated, and then the
3	Program Evaluation Report will be completed,
4	as necessary. If the Program Evaluation
5	Report identifies that, you know, from the
6	Program Evaluation Report it may identify that
7	claims need to be returned back to us for
8	rework, and we'll contact DOL about that.
9	If issues come up or if issues
10	cannot be resolved, obviously those issues
11	would move, we would move forward with either
12	an 83.14 or something with the SEC process, if
13	issues can't be resolved with some of the
14	issues that were previously identified from
15	SC&A. And that's about it.
16	CHAIRMAN MELIUS: Okay. Questions
17	for LaVon?
18	(No response.)
19	CHAIRMAN MELIUS: So can you go
20	back a slide?
21	MR. RUTHERFORD: Yes.
22	CHAIRMAN MELIUS: Because this is
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The Site Profile will be up --

1	sort of the speculative part of it. Okay. So
2	the key issue is the development of the
3	coworker model?
4	MR. RUTHERFORD: That is one of
5	the key issues. There are issues that
6	there are a number of issues that were
7	identified previously from the Site Profile
8	Review. The coworker model is being worked as
9	we go through, but there are other issues that
10	are being resolved as well, and plus SC&A is
11	reviewing previously defined or previously
12	identified issues, to see if they're still
13	applicable as well.
14	CHAIRMAN MELIUS: Okay. What is
15	the internal model cover intended?
16	MR. RUTHERFORD: What do you mean?
17	CHAIRMAN MELIUS: What exposures?
18	MR. RUTHERFORD: Well, it would be
19	exposures from internal releases for the
20	reactors and all the differences.
21	CHAIRMAN MELIUS: Okay. I'm just
22	trying to get a handle on what the Work Group

1	should be doing. I get concerned that this
2	site is sort of lagging behind, and we're not
3	
4	MR. RUTHERFORD: I know we have
5	one paper we're close to delivering later this
6	month, and then a few other papers. So I
7	think I don't want to speak for SC&A, but I
8	know they're reviewing issues right now, and
9	would come back with probably a follow-up
10	review of what's still applicable.
11	CHAIRMAN MELIUS: Because if I
12	guess what I'm getting at is if we have stuff
13	ready, I think we should start Work Group
14	reviewing and trying to resolve some of these
15	issues if that's appropriate, given what's not
16	completed.
17	MR. STIVER: This is John Stiver.
18	I may be able to fill in a little bit here. I
19	don't know if Steve's on the phone. He had
20	been, he's our lead for INL.
21	We had gotten started on doing
22	this background review of the existing issues

1	and we decided to go ahead and just kind of
2	put that on hold until the new information
3	comes out, to avoid having to go back and redo
4	all over again at a slightly later date.
5	That said, I mean if the Work
6	Group would like us to resume that, we could
7	certainly may get started on it.
8	CHAIRMAN MELIUS: Well, I guess my
9	question is, are there issues that are
10	appropriate to review, essentially independent
11	enough of what work has to be completed?
12	MR. STIVER: I guess the reason we
13	didn't want to spend a lot of effort on that
14	was because if things are changed
15	dramatically, then we would have just kind of
16	wasted that effort, and looked at pre-existing
17	information.
18	CHAIRMAN MELIUS: No, I understand
19	that. But I guess I'm hearing that some stuff
20	that NIOSH has completed.
21	MR. RUTHERFORD: Yeah, we're about
22	to complete.

1	CHAIRMAN MELIUS: About to
2	complete.
3	MR. RUTHERFORD: But we haven't
4	completed yet.
5	CHAIRMAN MELIUS: Okay. So when
6	they complete
7	MR. STIVER: As soon as LaVon
8	delivers the goods
9	(Laughter.)
10	CHAIRMAN MELIUS: Okay.
11	MEMBER BEACH: Isn't there like
12	three items? One's getting close and the
13	other two are at the end of the year? Is that
14	what I heard?
15	MR. RUTHERFORD: Yes, yes.
16	CHAIRMAN MELIUS: Yes. What
17	struck me was this coworker model. If we wait
18	until that's completed, it's well a year from
19	now, and I think that's and since we're
20	continuing to evaluate some of these coworker
21	issues, I won't say we'll shortcut that, but I
22	think we'll have some better criteria going

1	forward. Maybe we'll understand what needs to
2	be done there.
3	But I just hate to put off I
4	don't think we should be putting off
5	everything until the coworker model is done,
6	because that's, as I said, a full year. If we
7	have stuff done before then. Uh-huh. I mean,
8	yeah.
9	MR. STIVER: We'll go ahead and
10	start doing as the products are ready?
11	CHAIRMAN MELIUS: Yeah.
12	MEMBER GRIFFON: The question I
13	have was with the coworker models, it seems
14	like it's only for the reactor exposures. Are
15	there any considerations of coworkers for CPP,
16	and that's the
17	MR. RUTHERFORD: Yeah. I think
18	we're still evaluating that.
19	MEMBER GRIFFON: Okay, okay,
20	alright.
21	MEMBER BEACH: Jim, I don't have a
22	Site Profile question, and I'm not sure if

1	this is totally appropriate, but I'm going to
2	ask it anyway, since LaVon's standing at the
3	mic. Can you give us a little bit of history
4	or background on the site, on the petitions?
5	I know there's been three, and all
6	three of them have not qualified.
7	MR. RUTHERFORD: Right, well and
8	I'll go by memory on those three petitions.
9	One of the petitions was a very broad petition
10	over a large time period, identifying a lack
11	of monitoring data. That petition did not
12	qualify because they could not narrow it down
13	because we had monitoring data and we didn't
14	see it.
15	At that time, we did not see the
16	gaps or anything that would have supported
17	qualification. We had another petition that
18	was that did not qualify, because it I
19	don't know if they, and I'm trying to remember
20	the exact part of it.
21	But it was, they had
22	administrative problems. I remember what it

1	was. They weren't an eligible person to even
2	petition for it. They were an outside entity
3	trying to petition. I can't remember what the
4	third one. I know there was a third one, but
5	I can't remember what one was for.
6	I guess we have not had, you know,
7	and you know, when we work through these
8	issues, there definitely could be situations
9	that will come up that we identify in
10	feasibilities when we move forward with an
11	83.14.
12	CHAIRMAN MELIUS: Phil, maybe we
13	could I thought at one point we had talked
14	about doing a scheduling Work Group call, and
15	then we sort of dropped it, because I think
16	the schedule wasn't certain and so forth?
17	MEMBER SCHOFIELD: Yeah. I mean I
18	would be more than happy if you think there
19	would be some stuff, maybe to have a Work
20	Group in early November.
21	CHAIRMAN MELIUS: Yeah. But what
22	I think first is maybe a Work Group call, just

1	a short one, just to make sure we're all
2	understanding where we are and what the plan
3	is for well, as best we can. I mean you
4	know, things change and there's contingencies
5	and so forth.
6	But I think it would be helpful,
7	so that we're not it seems to me that we
8	start falling into a trap. Every time we're
9	saying well, it's going to take waiting for
10	the Site Profile to be complete, and it's just
11	taking a long time. I think we should be
12	making, trying to make some progress
13	MEMBER SCHOFIELD: I agree with
14	you.
15	CHAIRMAN MELIUS: Yeah, yeah. To
16	do that, good.
17	MR. HINNEFELD: If I could just
18	offer something here, something I think for
19	all the Work Groups to keep in mind is if we
20	meet on Live Meeting, you know, rather than
21	meeting in person, there's no need to have

eight hours' worth of things to discuss. You

1	know, we could have a Live Meeting for one
2	topic. You get one paper, you could have
3	that.
4	CHAIRMAN MELIUS: Yeah.
5	MR. HINNEFELD: So if you could
6	have more frequent shorter meetings online,
7	then you don't feel obliged to meet for a full
8	day when people travel.
9	CHAIRMAN MELIUS: Yeah.
10	MEMBER SCHOFIELD: This is a
11	question for Ted on those Live Meetings. If
12	we try and schedule those, do you guys have to
13	go on the Federal Register notice?
14	MR. KATZ: Work Group meetings
15	don't go in the Federal Register anyway.
16	MEMBER SCHOFIELD: Okay.
17	CHAIRMAN MELIUS: Yeah, I mean
18	yeah. So and certainly with Hanford and
19	others, we've done even short of live
20	meetings, you know. Phone conversations of
21	one or two hours can cover a lot of territory
22	and are easier to do, and I mean I'm a Member

1	of the Work Group with you, Phil, so it's
2	MEMBER SCHOFIELD: Well, it's also
3	
4	CHAIRMAN MELIUS: Up until this
5	point, I was waiting also, and then I see the
6	schedule and I think we're, you know.
7	MEMBER SCHOFIELD: No.
8	Unfortunately, I mean you're correct. We
9	haven't done much on Idaho in a long time, and
10	then we've got some others like Pinellas and
11	stuff. We might be able to just finish that
12	one up. But that actually happened to have
13	face to face. I mean it's just some of these,
14	that's not that much work left.
15	CHAIRMAN MELIUS: Any other
16	questions for LaVon? Again, a reminder for
17	people on the phone, please mute your phones,
18	*6. Yeah, yeah. I think so too, they can do
19	that.
20	<u>Public Comment</u>
21	CHAIRMAN MELIUS: Okay. We will
22	now open for a public comment period, and we

1	have a number of people signed up, some here
2	in person, and some on the phone. I'm going
3	to sort of go in a little order, in terms of
4	people that are here. I will start with
5	those, and then go on the phone and go from
6	there and do that.
7	The first person I have signed up
8	who's here is Joan Stewart. I knew I saw you
9	here some place.
10	MS. STEWART: My name is Joan
11	Stewart.
12	CHAIRMAN MELIUS: Do you need to
13	do your intro?
14	MS. STEWART: Good evening.
15	CHAIRMAN MELIUS: Hang on a
16	second. Ted has
17	MR. KATZ: Just a quick note I
18	should have said before we started this. But
19	you probably realize it, because I think
20	you've been here before. But public
21	commenters, everything that you say gets
22	transcribed, ends up in the transcript for the

1	public. If you say anything personal, that
2	too will be there.
3	But if you speak about personal
4	matters of other individuals, those things,
5	their privacy will be protected. So their
6	information will be redacted sufficiently to
7	protect their identity. So if you talk about
8	other people.
9	MS. STEWART: Okay, because I do
10	have one name to mention.
11	MR. KATZ: Yeah. I'm not
12	preventing you from mentioning the name. I'm
13	just saying that when we publish the
14	transcript for this, their name will be
15	redacted, for example.
16	So just to be aware of that, and
17	that's to protect their privacy. Even though
18	you may, they may tell you it's fine to talk
19	about me, so it's still required. So I mean
20	that's the short of the whole policy.
21	There's lots of details to it, and
22	it should be on the back table there for

people that are here in the room, and it's also on -- for people who are listening by phone, it's also on the website under the Board section at the top part. It talks about a Redaction Policy, and that's what I'm speaking to here. Okay, thank you.

CHAIRMAN MELIUS: Okay. Sorry for the interruption. Go ahead.

MS. STEWART: No problem. Hi, good evening. My name's Joan Stewart. Prior to being the senior-most radiological control technologist, technical supervisor at Rocky Flats, I was a union steward. I was the union steward that filed the 1987 grievance and safety concern over dosimetry at Rocky Flats, changing doses that were high into "no data available."

The aggrieved was, may I say it, [Identifying information redacted]. We went through three steps in our grievance process. During those three steps, it was noted by dosimetry that they had been doing this for

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years, changing doses to "no data available."

Generally, these were high doses. They were written down in pencil, and they were adjusted.

This came to the union's attention when [Identifying information redacted] had injured her knee at Rocky Flats, and was assigned up to dosimetry for a period of time, because she couldn't work the metallurgical operator. She was willing and able to provide us with proof that this was going on.

During the third step, second and third step, they admitted to not only doing this for years, but they said if they changed it, they would have to start pulling people out of the area. Hence the creation of the 100 Millirem Club.

DOE should have copies of, I don't know who the DOE person would be here. DOE should have copies of all safety concerns and grievances that were filed and their adjunct answers. As far as I know, all data was

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1	turned over to them. The union did destroy
2	their copies. I have a copy at home some
3	place I can possibly come by.
4	Now we have another question that
5	has arisen on the laboratories, to include
6	dosimetry. In the 90's, there was a DOE
7	accreditation program called laboratory
8	accreditation program. I don't know if you're
9	it was through DOE. Rocky Flats did not
10	qualify.
11	It caused quite the uproar at
12	Rocky Flats. They had to readjust a lot of
13	their procedures, because they were so far out
14	of qualification. They couldn't even be
15	certified. So you might want to look that up.
16	That was, I believe, a Tiger Team, part of the
17	Tiger Team report. So are there any
18	questions?
19	CHAIRMAN MELIUS: Any questions

MS. STEWART: I apologize.

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Yes, Dave. Dave, please use your

for Joan?

microphone.

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]	MEMBER	KC	OTELCHUCK:	What	do	you
mean	by	100	Millire	em	Club?			

MS. STEWART: There was a thing called an 100 Millirem Club. It was whether or not you were pulled out. They pulled the people out of the area if they had achieved 100 millirem in any given quarter.

That was started slightly about 1989, I would say. It took a little while to resolve the grievance and the safety concern.

Now I have heard that there's some talk on tritium in '76. We had a gettering system in '76. We had tritium bubblers in '76, and there were times when monitors would change -- of course monitors or RCTs, as you well know, would change out the tritium bubblers.

Sometimes you would have bubblers that were dry, and they were not sampling anything. So you may have times in your data that you will find that they couldn't get a reading on anything, because it wasn't pertinent, because they didn't have any

1 distilled water into the sampler. So if you 2 come by that. 3 overall, on the dosimetry But 4 practices, it appears as if you're working 5 with some very skewed statistics on this, 6 because your data set is off. Thank you. 7 CHAIRMAN MELIUS: Okay, thank you. 8 The next person I have signed up is Mark Nelson. 9 10 MR. NELSON: Hi. I'm Mark Nelson. 11 I spent my career at the INL here in Idaho, and I don't have all the details that Joan 12 13 had. However, I do have some questions and an I'll give you the observation observation. 14 15 first so I don't forget it, is it looks to me 16 like this whole mess is just going to stall until all of us die and it's no, never mind. 17 I started at the INL in November 18 19 1977 with a subcontractor called Chem 20 I did my orientation the Monday before Thanksgiving. By the 15th of December, 21

I was exceeding 2,800 mR. So I couldn't enter

an area until the 1st of the year. Then I hired on with -- I'm getting all -- anyway, with the contractor.

I worked at the ICPP, and every year for the next five years working for them, by October I couldn't enter any hot areas because of the amount of radiation factor I got. Now I really didn't pay a whole lot of attention to it, because I'm not that kind of guy.

But I got to thinking about it after I heard about this meeting just yesterday. I got to thinking about it, and my lifetime dose at the INL shows up at 10,000. I'm kind of wondering where the extra, because I was right at three for six years, 3,000 a year. I'm kind of wondering where the others went, kind of like Joan.

I'm here not primarily for myself, but for those who really ended up being in poor health because of their dedication to their country and their job, and I'm kind of

1	concerned that in my application I had
2	prostate cancer, and the reason that I was
3	turned down is because cadmium, exposure to
4	cadmium is not carcinogenic.
5	Yet every safety training I've
6	ever been to says stay of cadmium because it's
7	carcinogenic. I'm a little concerned about
8	the discrepancy there. I didn't quite catch
9	everything about that 100 mR Club. But when I
10	was not old at the site, we didn't consider it
11	worth our time to go in a hot area if we
12	didn't pick up 100.
13	Since I'm one who doesn't really
14	know what to do if it's not really hot. I am
15	not the only one who exceeded 2,500 to 3,000
16	every year for six to seven years, and yet it
17	does not show up on any of our records gross.
18	That's pretty much it.
19	MEMBER SCHOFIELD: Can I ask you
20	one question?
21	MR. NELSON: Yes.
22	MEMBER SCHOFIELD: As you

1	approached the administrative limit, which I
2	assume was 5R for the year.
3	MR. NELSON: Yeah.
4	MEMBER SCHOFIELD: Did they pull
5	you out of the area, and when they pulled you
6	out of the area, was it documented, or was
7	that just you were moved to another area?
8	MR. NELSON: Actually, they didn't
9	pull me out of an area. I just couldn't go
10	into hot areas. I could still go in and
11	operate in the operating corridor at 601 at
12	ICPP, but I couldn't go in the corridor and
13	pull samples, because the samples run 50 to
14	60R.
15	So I couldn't get those because I
16	would get more. I couldn't go in and decon
17	the cells. So I was not really pulled out of
18	my area. I was still able to operate, but I
19	was not able to do any decon or sampling or
20	those types of activities. So there was no
21	need for documentation. I was still working.
21	need for documentation. I was still working.

MEMBER SCHOFIELD: Okay, thanks.

1	CHAIRMAN MELIUS: Thank you.
2	MR. NELSON: You bet.
3	CHAIRMAN MELIUS: I'm going to go
4	to the phone now, and the first person I have
5	signed up on the phone is Chris Barker. Is
6	Chris Barker on the phone?
7	(No response.)
8	CHAIRMAN MELIUS: Okay.
9	[Identifying information redacted]?
10	MR. KATZ: You have to remind them
11	to press *6.
12	CHAIRMAN MELIUS: If you may have
13	it muted. Is [Identifying information
14	redacted] or Chris Barker on the phone? If
15	you have your line muted, hit *6 to unmute.
16	(No response.)
17	CHAIRMAN MELIUS: How about
18	Stephanie Carroll. Yeah. They have you
19	signed down as phone. I couldn't figure out
20	
21	(Off mic comments.)
22	CHAIRMAN MELIUS: Again, Chris

Barker, [Identifying information redacted] on the phone?

(No response.)

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CHAIRMAN MELIUS: Okay. We'll come back in a few minutes then. Stephanie Carroll can go.

MS. CARROLL: Okay. First, I'd like to thank the Board for allowing me to speak on some issues and concerns that I have on the ability of NIOSH to reconstruct dose. I'm advocate for Rocky Flats workers an regarding application for compensation under EEOICPA.

I therefore have access to many DOE documents from the site and from personal archives of the Rocky Flats workers. First, I would like to address the destruction and falsification of records. A document being presented by the petitioner, Terrie, refers to a DOE memo dated April 25th, 1996, issuing a moratorium on the destruction of records at the site, and including the Denver Federal

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This memo reinforces a sworn affidavit presented at the last meeting in Denver by a worker that admitted to being ordered to destroy records herself. She doesn't have a SEC claim and I believe that this memo should solidify her sworn affidavit.

[Identifying information redacted], a well-respected operations manager at Rocky Flats, Building 771, swore affidavit that he changed, my words "falsified." incident report the an at direction of his supervisor, to ensure that the cost was set to a level that would not have to be reported to DOE. If you'd like, you should maybe review that affidavit that [Identifying information redacted] swore to.

He's also used by NIOSH as somewhat of a site expert. He does have an SEC claim either, so he has no financial gain from his passing.

OPERATOR: The conference is now

in talk mode.

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MS. CARROLL: Next, I would also like to address the thorium issue. I spoke to former RCT this week that relayed information to me concerning thorium. He said that he surveyed the upstairs area of 444 by keeping down the center aisle while thorium ingots were displayed on what he referred to as "wine racks."

Ι spoke to another RCT actually, who said that they remembered the term "wine racks" being used in 444. I also have a question about thorium. After the thorium was removed from U-233 during thorium strikes, what happened it? it to Was processed and reclaimed? Was it treated as Where did it go? waste?

Another question. Rocky Flats did not record dose to the lens of the eye until the mid-90's. How is NIOSH assigning dose to the lens of the eye now for claimants with brain tumors?

Another question I have or concern is about radon, and I've brought this up before. But why was radon considered in Mound's petition and not ours? Buildings that were underground were located underground at Rocky Flats as a requirement of the processes performed on the site.

I do not believe that this is a natural environmental exposure. Workers were required to work underground. interviewing many workers, I learned that all short-lived isotopes were assumed to be radon and discounted. In my research, I learned most harmful effects uranium that the to workers are the effects of the short-lived isotopes, which were ignored on site.

I have documentation on short-lived, on an incident where a short-lived isotope was found on the worker's hands and a broom in Room 996, Building 991, and not found on the walls or the floor, mind you.

It was ignored, and put into the

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category of radon. This incident was not taken into account in the dose reconstruction, because it was assumed to be radon. So this man never got any incidence recorded in his dose reconstruction.

This assumption is not backed up by scientific evidence. Were all short-lived isotopes ignored because PU and uranium were the elements of concern? Background in all the documents I've seen, background continuously changes, depending on what room you're in, and it seems to always be slightly higher than the actual count taken on the worker.

What is the definition of background, and I also wanted to ask about if the workers were not paying attention to short-lived isotopes and actually any other isotopes besides uranium and plutonium, what about the exotic radionuclides? Who was paying attention to those? That's what I need to know.

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1	And well, that's all I have. But
2	I would like to thank Terrie Barrie, who has
3	tirelessly dedicated herself to the Rocky
4	Flats workers, to their families, to truth and
5	justice, and I just want to say she is our
6	hero. So thank you very much.
7	CHAIRMAN MELIUS: Well, thank you.
8	I think we have the phone issue taken care of,
9	sorted out. So Chris Barker, are you on the
10	line?
11	MR. BARKER: Yeah. Are you asking
12	for me? This is Chris Barker. Can you hear
13	me?
14	CHAIRMAN MELIUS: Yes, we can now.
15	Thanks, good, and we apologize. It does get
16	confusing with these phone systems, so but go
17	ahead. We can hear you now.
18	MR. BARKER: Great, thank you.
19	Chairman Melius and ladies and gentlemen of
20	the Committee, thank you for the opportunity
21	to speak today. Just so you know, I have a
22	little chest cold, so I may pause briefly and

go on mute, because I have to cough. So my apologies.

I'm Chris Barker. I have a Ph.D. in biostatistics from the Graduate School of Public Health. I am a consultant and I have an appointment as an adjunct associate professor of biostatistics. I am providing comments today on behalf of an individual, who I will refer to as the claimant.

The claimant was denied compensation for multiple cancers caused by exposure to plutonium after working at Rocky Flats. The claimant requested that I review the decision and the methodology for the determination of Probability of Causation.

Αt the claimant's request, Ι reviewed 1,000 over pages of NIOSH documentation for dose reconstruction assigned share. I documented and assembled 106 pages of material errors, questions, concerns, objections, false statements, admissions gaps, factual mistakes, circular

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logic, calculation errors, inconsistency and false and misleading claims in the NIOSH documentation, and the use of the NIOSH assigned share, the NIOSH models and processes and procedures, based on the documentation that was publicly available.

Furthermore, I requested that NIOSH provide several additional pieces of information about the claimant's dose reconstruction and other details about the I stated in calculation. the 106 document that upon receipt of the requested materials, I may have additional comments about their processes and procedures and software.

The clamant forwarded 106 pages of my document with the 1,000 pages of document that I had reviewed to the appropriate NIOSH office. Recognizing my time is limited, I will highlight only a few of the errors from the 106 pages of errors and calculation errors and misstatements and false claims that I

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When I discovered, for example, in NIOSH documentation that they're talking about Probability of Causation, I'll discuss that in a little bit. The full, as I say, full 106document and the 1,000 of page pages additional materials have been forwarded to NIOSH earlier for review, and I have colleague at the meeting there who has, make additional copies available Committee as needed.

The issues underlying the calculations used by the NIOSH models involve matters of life and death. I am an expert in the statistical methods that are appropriate for the correct cross-analysis of data arising in these circumstances.

What remains inexplicable is although we submitted these documents, this 106 pages plus the 1,000 pages of documentation months ago to NIOSH, we have never received any reply about the comments

and questions and concerns that I had raised.

So because of the errors in the documentation, Ι have not been able replicate the calculations of the assigned share or the dose reconstruction. all aware, replication of а result, а particularly a calculation, is a fundamental principle of science, because part of consulting work involves pharmaceutical drug development for life-threatening illnesses, I prepare work that is forwarded to the Food and Administration and other regulatory agencies, and I can assure you that no part of the documentation of procedures would ever be accepted by a regulatory agency anywhere in the world.

So I want to refer you to the 106page detailed document, which can be provided,
and I'm only going to highlight a few of the
errors that I found. The first is that the
NIOSH IREP model, which does not incorporate
all the uncertainties in the dose

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reconstruction process, nor the uncertainties in relating dose reconstruction to so-called probabilities of cancer.

I enumerated those uncertainties that are not incorporated in the 106-page document. Just to give you a sense, those uncertainties relate to handling of missing dose information, statistical distributions used and uncertainties about parameters, means and standard deviations that were assumed in the IREP.

The IREP performance were statistical Type 1 and Type 2 errors. In other words, falsely detecting a relationship of the radiation and cancer or erroneously ignoring the relation of radiation dose and cancer when it exists are unknown. These are things that would easily be obtained from an appropriate statistical analysis.

The NIOSH IREP claim of 90 percent uncertainty integrals is misleading because all the uncertainties are not included. Many

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NIOSH documents and letters to the claimant refer to Probability of Causation.

The calculation performed by the program IREP of assigned share is not a probability of any kind whatsoever. It has no probabilistic interpretation. Any use of the term "Probability of Causation" is a false and misleading term.

The NIOSH assigned share is not a probability. They do not indicate causality of any kind. The numerous statistical methods that address causality, I worked with methods for establishing causality for relations such as this every day.

These methods have been available in statistical and other literature since R.A. Fisher considered the father of statistics, developed these procedures in the 1930's, and since Reverend Bayes developed methodologies back in the 1700's for establishing causality.

I documented specific errors in Excel and comma separated files that the

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1	claimant was told were used as inputs to the
2	IREP software.
3	(Interruption.)
4	CHAIRMAN MELIUS: Chris Barker,
5	are you on the line?
6	(No response.)
7	After we got interrupted.
8	[Identifying information redacted], are you on
9	the line? Okay. Terrie? I know. That's what
10	I'm going to do. So Terrie, do you want to
11	read [Identifying information redacted]
12	comments, [Identifying information redacted],
13	yeah.
14	MS. BARRIE: And I do have Chris'
15	comments. He was almost through, and I'd be
16	happy to make copies of whole presentation.
17	CHAIRMAN MELIUS: That would be
18	that would be, yeah.
19	MS. BARRIE: Okay, and this was
20	from
21	CHAIRMAN MELIUS: And we'll make
22	copies for you.

MS. BARRIE: Okay. This is public comments from [Identifying information redacted] from EECAP. She's been involved with the Mound SEC petition, and she says, and this is her quotes, "First, I'd like to thank Dr. Melius and the Board for allowing me at this time to speak. I wasn't able to attend this meeting, but I have tried to listen into it, which given the sound quality has been a real chore at times.

"I appreciate that Stu Hinnefeld discussed some of the problems that turned up in the FOIA request from 2001, which I received a few months ago.

"I couldn't hear clearly, but it sounded like he and Dr. Melius discussed that NIOSH was looking at problems with the conflict of interest policies, problems with undermining the Mound Class Definition for the 1959 to 1980 SEC, and problems with NIOSH employees withholding information from the Board, as well as disparaging remarks made

among NIOSH employees about Board Members who deigned to ask them questions.

"After waiting over two years, NIOSH finally provided me with their emails from their employees on the Mound SEC radon issue. I want to point out that this FOIA was not sent to me until after the Mound SEC was already closed. But whether this was done purposely or not, I do not know.

"I was frankly shocked by these emails that showed NIOSH running roughshod over the Board; rather than being led by the Board; NIOSH employees writing dismissive and disrespectful emails about the Board Members, SC&A and DOL; NIOSH withholding evidence from the Board and DOL for almost a year; NIOSH employees' bias directing the Class Definition, rather than the Board defining the Class; NIOSH employees making assumptions on how things were done at Mound, rather than doing the research in the DOE documents to see what was actually done, and then ignoring the

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research and documentation after EECAP sent it to them.

"NIOSH defining the SEC Class base; the ignoring of the exposures to thoron and actinon and the incorrect assumption that no workers without bioassay had been exposed to radon; blatant disregard to conflict of interest laws; and in many instances the lead NIOSH employee soliciting information from a conflicted NIOSH employee.

"If you wish to see the documents for these claims, you can find it at the EECAP website. These actions raise serious questions that I think need to be investigated for all sites, not just Mound. What allowed this kind of behavior to occur and go on for years? Is part of the problem the culture at NIOSH?

"What allowed NIOSH to blatantly abuse the conflict of interest laws? Why is NIOSH running the Board rather than the Board monitoring NIOSH? The 1959 to 1980 Mound SEC

needs to be reopened, and all employees need to be included. In fact, this is what NIOSH said that would be done on February 11th, 2011.

"The process was begun, because they realized that they had made an incorrect assumption that all workers in the R and SW Buildings had been bioassayed. The NIOSH employee who caused this mess then said he had 'forgotten,'" that's in quotes, "about the cold side of the building, where no one had been monitored.

"A week later, NIOSH reversed its decision after talking to Ted and the OGC. I'd like to tell you more about this discussion, but that email was not provided to me. What did Ted and the OGC say that overrode the science that is supposed to drive the SEC process? I don't know, but I hope the Mound Work Group will find out.

"After seeing how damaging the pages were that were released to me, I am very

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curious to know what was withheld from the FOIA release. A total of 641 pages were withheld, 393 under Exemption 5 and 248 under Exemption 6.

"I would encourage the Board Mound Work Group or SC&A to examine these pages, to make sure they contain no additional illegal or unethical behavior. I have no power to do this, but you do. [Identifying information redacted] sent ANWAG a quote from the law, 18 U.S.C. 1001(a)(1), (2), (3), that indicates NIOSH's actions, as documented by these emails, could lead to criminal charges being filed.

"That statute states 'Except as otherwise provided in this section, whoever in any matter within the jurisdiction of the executive, legislative or judicial branch of the government of the United States knowingly and willfully (1) falsifies, conceals or covers up by any trick, scheme or device, a material fact; (2) makes any materially false,

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fictitious or fraudulent statement or representation; or (3) makes the use of false writing or document, knowing the same to contain any materially false, fictitious or fraudulent statement or entry, shall be fined under this title and imprisoned to not more than five years.'

"While the workers were not harmed only ones by NIOSH employees' misbehavior, they were the ones who paid the cost of loss of benefits and medical care because of it. The damage done to the workers is the reason for this program. They were already betrayed once by their government. second governmental betrayal is really beyond the pale.

" T thank the Board for their attention to this serious matter, and would appreciate if Ι could be informed, as appropriate, on what is being done. let me know if I can help in any way. you, " from [Identifying information redacted].

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1	CHAIRMAN MELIUS: And we've
2	already but Terrie, did you have public
3	comments to make separately? You signed up. I
4	didn't
5	MS. BARRIE: Yes, I did, and it's
6	just really a minor one.
7	CHAIRMAN MELIUS: Go ahead.
8	MS. BARRIE: I would like to ask
9	the Board's reconsideration of the time limit
10	for SEC petitioners' presentations. I fully
11	agree with the time limit. I have been at
12	meetings where, you know, people tend to go on
13	and on.
14	But having a ten minute limit for
15	petitioners, especially ones who are preparing
16	PowerPoint presentations, it's really tough to
17	convey the information we want to convey to
18	the Board. So I was thinking that perhaps we,
19	the Board could go on an individual basis, to
20	see, check with the petitioner, how much time
21	do you think you need?

That's too much, try to, you know,

1	cut it down to X amount, and that way we can
2	prepare, in a timely, you know, within the
۷	prepare, in a crimery, you know, wrenim ene
3	time line and still get the information
4	across. Thank you.
5	CHAIRMAN MELIUS: Thank you. Does
6	anybody else on the line or in the audience
7	wish to make public comments?
8	(No response.)
9	CHAIRMAN MELIUS: If not, we're
10	adjourned. Thank you everybody. We'll see
11	you, everybody, the Board here tomorrow
12	morning.
13	(Whereupon, at 5:44 p.m., the
14	meeting was recessed, to reconvene on
15	Wednesday, July 17, 2013 at 8:15 a.m.)
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