UNITED STATES OF AMERICA

CENTERS FOR DISEASE CONTROL

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NATIONAL INSTITUTE FOR OCCUPATIONAL SAFETY AND HEALTH

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ADVISORY BOARD ON RADIATION AND WORKER HEALTH

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102nd MEETING

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THURSDAY, NOVEMBER 6, 2014

The meeting convened at 8:30 a.m., Pacific Standard Time, in Hilton Garden Inn Los Angeles/Redondo Beach 2410 Marine Avenue, Redondo Beach, CA, James M. Melius, Chairman, presiding.

PRESENT:

JAMES M. MELIUS, Chairman
HENRY ANDERSON, Member
BRADLEY P. CLAWSON, Member
R. WILLIAM FIELD, Member*
DAVID KOTELCHUCK, Member
RICHARD LEMEN, Member
WANDA I. MUNN, 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

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AL-NABULSI, ISAF, DOE ADAMS, NANCY, NIOSH Contractor BARRIE, TERRIE* BLAZE, D=LANIE BURGOS, ZAIDA, NIOSH CRAWFORD, CHRIS A FRANK, DOL FITZGERALD, JOE, SC&A HARTSFIELD, DEKEELY, HHS HINNEFELD, STU, DCAS HUGHES, LARA, DCAS KINMAN, JOSH, DCAS KLEA, BONNIE KUROWSKY, LORRAINE LEWIS, GREG, DOE LIN, JENNY, HHS MCKEEL, DAN* MCFEE, MATT, ORAU Team NETON, JIM, DCAS PACE, JOHN* ROESCH, CHARLEEN ROESCH, DANIEL RUTHERFORD, LAVON, DCAS SCHULTZ, JEFF* STIVER, JOHN, SC&A WORTHINGTON, PATRICIA, DOE ZEITOUN, ABE, SC&A

$\texttt{T-A-B-L-E} \ \texttt{O-F} \ \texttt{C-O-N-T-E-N-T-S}$

Welcome and Introduction Dr. James Melius4
NIOSH Program Update Stuart Hinnefeld6
DOL Program Update Frank Crawford
DOE Program Update Greg Lewis
Procedures Subcommittee report on Completed Review of Program Evaluation Report 14: Construction Trade Workers Wanda Munn
Sufficient Accuracy/Coworker Dose Modeling Dr. James Melius
SEC Petitions Status Update by LaVon Rutherford
Board Work Session by James Melius
Subcommittee and Work Group Reports by James Melius
Santa Susana Special Exposure Cohort Site Profile Update by Lara Hughes
SC&A Site Profile and SEC Updates by John Stiver
Public Comment

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

1 (8:29 a.m.)2 CHAIRMAN MELIUS: -- Radiation and 3 Worker Health, call to order. And I'll turn it 4 5 over to Ted. MR. KATZ: So welcome everyone to the 6 meeting. For everyone listening on the phone 7 too, the materials for the meeting today, for 8 9 people in the room -- I don't see any public 10 members yet, the materials are outside on the table. 11 12 For people on the line, the materials 13 are on the Internet, on the NIOSH website, under 14 the Board section for Meetings, today's date. And you'll find -- you should find all of the 15 presentations that are being given today there 16 17 for your perusal. 18 And there is also Live Meeting. 19 can -- so you can follow along while people are

giving presentations on Live Meeting. that link for Live Meeting is on the agenda, which is on the NIOSH website.

> Roll call. We have no topics for

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which there are conflicts, so I'm not going to address conflicts for Board Members. So we're just going to run through roll call. And let's just go alphabetically from the top. There are a few Members that -- we're having feedback here, I think -- who I know are going to be absent.

(Roll call.)

MR. KATZ: Very good. Okay. And a few other things just to note, there's a public comment session that begins at 4:30 today, 4:30 to 5:30.

So people on the line, we don't have members of the public in the room right now, but people on the line, we will be taking people in the room first, addressing Santa Susana, but then we'll get to you folks on the line.

And we have a number of people in the public who have already signed up, by send -- sent me emails, but you don't -- the rest of you don't need to send emails or what have you. We'll get to you after we finish with people in the room.

And then just let me note, for everyone on the line, please mute your phone, except when you're addressing the group. And if you don't have a mute button, press *6 to mute your phone. Press *6 again to take your phone off of mute. But please keep your phones muted while you're just listening. Thank you. Dr. Melius, this is your meeting.

CHAIRMAN MELIUS: Okay. Thank you,
Ted. And we'll start, as usual, with a NIOSH
update. Stu Hinnefeld.

MR. HINNEFELD: Okay. Now we'll try it. Okay. Thank you, Dr. Melius. And I'm here for my normal program update. Okay, good.

I always -- I try to start with program news on these updates, and sometimes I don't think of very much news, probably because it's not news to me, because it happens to me every day. But since our last meeting, we have mainly outreach activities to talk about. We've done a number of them, either -- well, really in conjunction, mostly in conjunction with the other agencies.

The first one, though, was one that we did with our outreach contractor, ATL, and that is the Dose Reconstruction and Special Exposure Cohort workshop that we offer once a year -it's been in September the last couple of years, where we invite interested parties, a few advocates, mainly lot of labor representatives, some retiree organizations, representatives of those folks Cincinnati, and we conduct a two-day workshop about the program, dose reconstruction, various aspects of it.

So certain members of our staff present certain sections or certain topics. Staff from ATL present other topics, and then there's some working sessions for people to become more familiar with our website and where to find information.

The idea behind this is that these people then can be resources for their constituencies back at home, and can be sources of information for those folks back at the facilities.

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So it went pretty well. ATL does a nice job of setting that up, and we've -- typically get very positive comments. They do an attendee assessment, essentially a course assessment thing at the end.

Pretty consistently, we get pretty high marks about the usefulness, people, you know, really happy to be there and they thought the information was presented well and they enjoyed it.

So we did go through that in September. And then there were several Joint Outreach Task Group meetings in the months since the last Board meeting, in Richland and Spokane -- that was one trip for the two locations, Rochester and in Paducah, just a -- I guess it's about three weeks ago now.

So those are most -- that's most of the news that I thought of to put on the slide. Getting into the claim, or the statistics, I won't get into -- spend a whole lot of time on these. They're on the handouts, and I think they were probably in information you received.

The claims are, continue to go up at 1 about -- we continue to get about 200 a month, 2 new -- maybe slightly less than 200 a month of 3 new claims. We have a kind of a constant flow 4 5 of reworked claims. Typically when people 6 get an 7 additional cancer, they're sent back So in combination of the new and rework. 8 9 reworks, we're probably around 250 a month that we get. 10 And we continue to send cases back. 11 12 These are the various categories. The 1,200 13 cases still with us, I believe, a number of those are cases where the initial draft is in 14 15 the hands of the claimants, and we're waiting for the close, either doing a close-out 16 interview or going to get the OCAS-1 form back. 17 18 So it's really, the number we have is 19 somewhere around, well 9 -- it looks like about 20 970 based on this accounting here. 21 So those are -- that's how the game's

-- you know, we've -- that's kind of been our

sort of inbox for a while now. We're not -- we

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have certainly timeliness objectives, in terms of getting cases done, I think 90 percent of the cases within five months of getting all the information we need to do it.

And so we're not making -- we don't really have an objective to reduce the number in the inbox. We just want to make sure we're timely in the response to the claims, as timely as we can be.

Here is the breakdown of Probability of Causation, how the -- these are ones done from DRs. This is just the DR statistic. It doesn't take into account cases that were compensated through the SEC process.

As I recall, that's either 28 or 29 percent. I did the arithmetic and -- but I didn't write it down. Or I don't -- I wrote it down, but not with something I have with me. I think it's like 28 or 29 percent have been successful through dose reconstruction.

Again, a lot of the -- a number of the cancers that tend to be successful with, through dose reconstruction, like lung cancer

in particular is an SEC cancer, and so as you add additional SEC Classes, you don't have those, you know, those don't come to dose reconstruction so you don't get those successful dose reconstructions when they're paid through the SEC.

This is our chart, our long-term chart of submittals versus production. It doesn't change much. The last data point really shouldn't be on there. That's a partial quarter.

These are quarterly data, and there is

-- there was no precipitous drop in claims

recently. That's just a partial quarter, and

we were close enough to the meeting and getting

the meeting materials ready that it was a little

hard to re-run.

So we said, well what the heck, I'll just explain it. That's a, that's only a partial quarter. There's no, there was no precipitous drop in claims recently.

Here's status of the first 5,000 claims and how they break out. Nothing's much

different there. This slide caught my attention because -- and if you look at the claims at NIOSH, in the first 10,000, there are 20 DRs in process, and three are initials.

And that bothered me a little bit, because I knew that we had, a while ago, we had a number of claims that were initial even though they had low numbers because they were CLL claims, that CLL originally referred to us. And so we gave them a claim number, even though DOL shouldn't have referred them to us, because CLL wasn't -- you know, chronic lymphocytic leukemia wasn't covered.

So, and then when the regulation changed, we -- the DOL returned those to us. And so they came back to us, and they still had their low numbers. So I was pretty confident, for a while, that we, when we'd have these low numbered initial cases, it was because they were a CLL case.

And I saw this slide and I said, you know, that was quite a while ago. That seems like we shouldn't have those. So I looked

those up. Excuse me a minute.

Two of the cases were pulled before being completed. Either the claimant opted out of the process or died, unfortunately, before the claim was submitted. And it was closed for years, seven, eight, nine years, until a survivor either was identified or decided to pursue the claim.

And so it was reinitiated then, with the survivor's action, and that happened just very recently. And so those claims are now active. They never had a final dose reconstruction done. And so it's still in the initial category, but they were just recently renewed and reinstated because the survivor picked up the claim.

The other -- the third one was a CLL case that was activated with the rest of the CLL cases, and then it, because it was at a site where -- well it was the Fernald site. It was a claim from the Fernald site.

The Fernald site, we had SEC decision debate. We -- typically we don't pend claims

when we have an SEC claim, you know, petition in front of us. But when we get close to the -- when we think we're close to the end, we'll pend the cases, and so that we're just going to finish up this SEC, we're going to finish up all the DR, or the TBD issues, and so we'll pend the claim so we just do them once.

Well, with Fernald, there's still some Site Profile issues still hanging on, and it was getting longer and longer, and we said, the heck with it. Let's do them the way we would do them now. When we finish the DR issues, if we have to, we'll do a Program Evaluation Report and we'll do them, rather than just have them sitting there waiting then.

So that's what the third one is, an initial one. So, I am -- I did -- at least for this meeting, I did pay attention to that slide.

Okay. These are our DOE response numbers. I do not have the comparison to last month but I could probably find them if I need to. These really look pretty good to me. We don't have any sites, I don't think, that are

particularly problematic. And the 256 is, I think, a really pretty low number, considering how many claims we get all the time.

And our SEC summary table, I won't move too far into this, because LaVon has a presentation all about SECs at some point today or tonight. So we'll -- I'll let him go through that, but as -- while at one time, let's see, yes, while at one time the breakdown between 83s and -- 13s and 83.14s was pretty even.

That was the time when we were finishing our research on the various AWE facilities where we didn't have very many claims. We went through that and we ended up getting, adding a lot of SECs for that. And so we kind of caught up with .13s.

Well, we're pretty much through that process now. So the petitions from now on will probably be, for the most part, 83.13 petitions. So, I think that was it. Yes, I guess it's not going to take me out of it, but I can get out of it over there.

So, are there any questions?

1	CHAIRMAN MELIUS: Any questions for
2	Stu? We'll let LaVon explain some of the
3	MR. HINNEFELD: Yes, the
4	CHAIRMAN MELIUS: SEC issues.
5	Dave?
6	MEMBER KOTELCHUCK: I was just
7	curious. I mean, you for administrative
8	reasons you separate out the DR and SEC cases.
9	MR. KATZ: David, would you speak
10	right into your mic, so that
11	MEMBER KOTELCHUCK: Right. Is the
12	mic on? Okay. For administrative reasons,
13	you separate SEC and DR cases, but I've tried
14	to look at the numbers and see, of all the cases
15	that are submitted, of all the individuals
16	submitting claims, whether they end up in DR or
17	SEC, if they are accepted by DOL as having been
18	employed over 250 days, that is that they are
19	potential, what fraction of them are, in fact,
20	compensated?
21	MR. HINNEFELD: Well, there might be
22	some statistics on the DOL website that might
23	answer that. I'm not familiar with them. The

information that we have, I think I have a slide 1 back here, you can see in the second major side 2 where you have all the sub-bullets, there are 3 4 3,351 cases for SEC Cohort, Special Exposure Cohort. 5 Those are claims that were referred to 6 And while the claims were with us, 7 us by DOL. an SEC was added that included, probably, those 8 9 I say probably because when DOL first 10 refers the case to us we'll have a particular cancer diagnosis on it. And we will pull it 11 12 based on the cancer diagnosis that they had when 13 they sent it to us. It has happened that when they go to 14 final adjudication they take another look at 15 the case and they have a different diagnosis. 16 17 And so it may, essentially, fall off the SEC. 18 So that is a pretty good approximation. 19 you would add that number, the 3,351 to -- oh, 20 I'm sorry, to the 10,073 --21 MEMBER KOTELCHUCK: Oh, okay. 22 MR. HINNEFELD: -- you would have an 23 approximation of -- but then you've got to add

1	the same got to add it to the 35,667 also.
2	MEMBER KOTELCHUCK: Right.
3	MR. HINNEFELD: So you'd have an
4	approximation of the fraction, the ones that
5	got to us. Now, once an SEC Class is added, we
6	never see any more claims from that site because
7	the Department and it's an SEC cancer, the
8	Department of Labor just pays them.
9	MEMBER KOTELCHUCK: I see.
10	MR. HINNEFELD: So the best
11	information about how they're I think would
12	be on the DOL website. I think they have some
13	statistics about site-specific outcomes there.
14	MEMBER KOTELCHUCK: Oh, very good.
15	I will do that. I will check that.
16	MR. HINNEFELD: Okay, my handy
17	assistant Kato has just sent that was
18	probably inappropriate, wasn't it? In Part B,
19	according to the DOL let me see what I'm
20	looking at, there have been 92,609 cases filed.
21	And it looks like there are 42,000 cases that
22	were paid. So that's a little less than 50

percent.

1	MEMBER KOTELCHUCK: Well, it's close
2	to 50 percent, just
3	MR. HINNEFELD: Yes. And, now that
4	well Part B, though includes silicosis and
5	
6	MEMBER KOTELCHUCK: Beryllium.
7	MR. HINNEFELD: berylliosis.
8	MEMBER KOTELCHUCK: Right.
9	MR. HINNEFELD: Beryllium disease.
10	So
11	MEMBER KOTELCHUCK: Nevertheless,
12	it's much larger than the number you had, than
13	the DR number.
14	MR. HINNEFELD: It's much larger than
15	the 28 percent that is done through dose
16	reconstruction, yes.
17	MEMBER KOTELCHUCK: Okay, thank you.
18	CHAIRMAN MELIUS: Yes, Paul?
19	MEMBER ZIEMER: Stu, my question is
20	on the workshop that you held, and you described
21	the types of individuals who participated.
22	Can you give us an idea of the actual numbers
23	of people who participated in the

1	MR. HINNEFELD: It was about 30.
2	MEMBER ZIEMER: About 30?
3	MR. HINNEFELD: Yes. Slightly less,
4	I think.
5	MEMBER ZIEMER: And is that covered
6	by your budget, in terms of their travel and so
7	on, or do
8	MR. HINNEFELD: Yes, yes.
9	MEMBER ZIEMER: Okay.
10	MR. HINNEFELD: Yes. We pay their
11	travel, and for certain attendees, we provide
12	wage replacement.
13	CHAIRMAN MELIUS: Any other
14	questions for Stu? Any of the Board Members on
15	the phone wish to ask questions?
16	MEMBER FIELD: No, not at this time.
17	Bill.
18	CHAIRMAN MELIUS: Okay. Yes, Henry
19	has a question though, here.
20	MEMBER ANDERSON: Yes. Stu, you
21	have here that you've completed 131 of the SEC,
22	and ten of them are with the Board. Are there
23	any that you've completed that aren't with the

1	Board?
2	MR. HINNEFELD: No. Anything that
3	we've completed has either had action taken
4	MEMBER ANDERSON: Yes, okay.
5	MR. HINNEFELD: or is with the
6	Board. There are two that we are working on
7	MEMBER ANDERSON: Yes, right. Okay.
8	MR. HINNEFELD: that we have not
9	presented to the Board yet, I think. Bomber
10	will give the numbers later on
11	MEMBER ANDERSON: Okay, yes.
12	MR. HINNEFELD: but there are some
13	we are working on now that we've not presented
14	to the Board, but
15	MEMBER ANDERSON: Right.
16	MR. HINNEFELD: everything where
17	we've finished an Evaluation Report, it's
18	either in the Board or it's been it's with
19	the Board or it's been dispositioned by the
20	Board.
21	MEMBER ANDERSON: Okay. Thank you.
22	CHAIRMAN MELIUS: We'll reveal that
23	in greater detail some time between now and

midnight tonight. Okay. 1 Nofurther questions? Okay, thank you, Stu. We'll work 2 on coming up with some more to that. 3 Okay. We'll now get an update from 4 5 the Department of Labor. Frank Crawford. MR. CRAWFORD: Good morning. I'm 6 Frank Crawford, a health physicist with the 7 Department of Labor. And let's see if we can 8 get -- yes. 9 Sorry. Just parenthetically, it's hard to 10 tease out the exact effect of SECs, in answer 11 I have some slides here that 12 to your question. 13 give you a slightly different view of the thing, but between the CBD cases and the silicosis 14 15 cases, there are so many factors floating around -- I'm sorry, that it's difficult to say 16 exactly how many cases have been paid because 17 18 of the existence of an SEC alone. 19 MEMBER KOTELCHUCK: Yes, I know. Ι 20 And often people ask, well what fraction know. 21 of the cases submitted are compensated? And I never have been able to say, other than the DR, 22

which is about 30 percent, right?

23

I think it's

1 about 30 percent. MR. CRAWFORD: 2 Yes. 3 MEMBER KOTELCHUCK: Yes. Also, before we MR. CRAWFORD: 4 Yes. started, you'll notice there's 5 get discrepancy between the numbers that I 6 presenting and the numbers that Stu presents. 7 There are various reasons for that. Part, it 8 9 is different reporting periods, and part is definitional, in that we have a different view 10 because Stu never sees SEC claims that are 11 12 simply disposed of by DOL alone. All right. 13 Okay, we'll go to the first slide. Ted mentioned, this is all on the website, so 14 I'm going to hurry through some of the slides, 15 and there are also slides that we will not see 16 17 in the presentation but are generally 18 informational, about what kinds of claims are permitted under Parts B and E and so forth, who 19 20 are the survivors under both cases. But 21 there's no point in presenting that here. 22 By our count, then, that is DOL, we have about 175,000 cases filed, and almost 11 23

billion in total compensation paid to date.

This is where the numbers start differing a little.

We show that 43,000 cases, approximately, have been referred to NIOSH for dose reconstruction, and approximately 41,000 have come back, leaving NIOSH with about 2,000 cases. As Stu has told you, really, it's probably 1200 cases, but we'll have to live with those discrepancies.

We show a slightly higher fraction approved with DRs, just a little bit over a third, and I don't know how to account for that except perhaps, again, reporting periods. And also, we're only talking about final decisions, which is a distinction, perhaps, that may not be in Stu's statistics. I don't know.

So we have 35,000 cases returned with a DR, and 28,000 now have a final decision as well, and that's where we get, under that, our 35 percent approval rate. This pie chart is maybe a little hard to read, but I'm sure on the website it'll be much clearer.

The only thing I'd like to point out 1 here is that the other category probably does 2 include silicosis and CBD, and it's -- there may 3 be other kinds of failed claims, where they 4 5 weren't federal workers as it turned out, or didn't qualify for some other reason, I'm not 6 And that's a little bit opaque for us. 7 In terms of SEC cases, we see that it's 8 9 almost 50/50 between cases not referred to NIOSH and cases referred to NIOSH. 10 It's 12 percent versus 14. So with that, we have some 11 12 evidence of what's going on, but again, it's 13 very hard to tease out exact numbers. 14 go. In this slide, we show slightly over 15 50 percent, what you might call a total approval 16 17 rate, that is, with SEC cases taken into account 18 with simply dose reconstruction cases. 19 Now, in this next slide, we see that 20 the accepted SEC cases far outnumber the 21 accepted DR cases by more than 2 to 1. 22 21,700 versus about 9,200 DR cases.

There's also one other interesting

stat here, which is the third bullet point, cases accepted based on SEC status, and with a PoC greater than 50 percent. That's a relatively small number, 700 cases.

And remember, on a previous slide, we saw that 12 percent of cases were referred to NIOSH and had an SEC. So of those cases, which must be a substantial number, only 700 returned with a positive result, you might say.

Hard to interpret, but my best guess is that SECs sweep up everybody at a site, of course, I mean typically -- not all sites, but typically, and many of those people would not have had significant exposure.

So if they're accepted first by an SEC and then ask for medical benefits, and therefore a dose reconstruction, they're less likely to be approved than somebody who has gone the other way, through the dose reconstruction process and has had significant exposure. So I think that's what's going on there, but -
MEMBER RICHARDSON: I'm sorry. I'm

a little confused.

Right. 1 MR. CRAWFORD: MEMBER RICHARDSON: 2 You're saying that half the claimants who are covered under 3 4 SEC nonetheless having an are dose 5 reconstruction by NIOSH? MR. CRAWFORD: Yes, but that could be 6 7 because they had the dose reconstruction first and then the SEC was approved second, or it 8 9 could be because they're SEC Class members and 10 they requested a dose reconstruction in order 11 to get medical benefits. Stu, you can help me 12 out. Just one other point, 13 MR. HINNEFELD: the medical benefits is for a cancer that's not 14 15 an SEC cancer. So in other words, the claimant would have more than one cancer. One would be 16 17 an SEC cancer, and that would put them into the 18 SEC and they'd be compensated through SEC. 19 With another non-SEC cancer, their 20 successful claim for the SEC cancer will not pay 21 for medical benefits for that other cancer. successful 22 they have to have а

reconstruction to have medical benefits paid

1	for the non-SEC cancer.
2	MEMBER RICHARDSON: So in this case,
3	that number refers to the Probability of
4	Causation under a calculation where the target
5	organ may not have been the primary may not
6	have been an SEC cancer?
7	MR. HINNEFELD: Correct. It would
8	not have been an SEC cancer.
9	MR. CRAWFORD: But both are evaluated
10	for the PoC. In other words, if a person had
11	a lung cancer and a prostate cancer, they
12	qualify for the SEC solely on the lung cancer,
13	of course. But if they ask for medical
14	benefits for the prostate cancer, then when the
15	case goes back to NIOSH for a DR, both cancers
16	are evaluated de novo, as if there had been no
17	SEC.
18	MEMBER RICHARDSON: I'm still
19	confused why you would do that. Isn't that
20	just a lot of time spent by NIOSH to do a dose
21	reconstruction for something which is going to
22	be compensated regardless?

Well

CRAWFORD:

MR.

23

compensated

1	monetarily, but if the prostate cancer, for
2	instance, was treated at great expense
3	MEMBER RICHARDSON: No, but the lung
4	cancer.
5	MR. CRAWFORD: Oh, the lung cancer is
6	covered by the SEC, and that's
7	MEMBER RICHARDSON: Yes.
8	MR. CRAWFORD: Yes.
9	MEMBER RICHARDSON: But you're
10	saying that the calculation is done,
11	nonetheless?
12	MR. CRAWFORD: Using both cancers to
13	be I mean, it's only fair, essentially. If
14	there were no SEC, that person would be
15	evaluated on the basis of both cancers.
16	MR. KATZ: The reason for that is
17	because multiple cancers, there's a
18	statistical treatment that's given it's
19	advantageous, if you have multiple cancers, to
20	have all those considered when they do the PoC.
21	So that's why they reconstruct the
22	doses for all the cancers, including the one
23	that's covered by the SEC. Because for each

1	primary cancer, there's a statistical
2	treatment that combines those in giving you a
3	total PoC.
4	MEMBER RICHARDSON: And that's going
5	to help them with medical compensation?
6	MR. KATZ: That'll help them with the
7	cancer that's not covered.
8	MEMBER RICHARDSON: For the medical
9	compensation of the
10	MR. KATZ: Exactly. Exactly.
11	MEMBER RICHARDSON: non-covered
12	cancer.
13	MR. KATZ: Exactly. So that's just
14	beneficial for them.
15	MR. CRAWFORD: Then I think we can
16	leave this slide. No surprises here. Our top
17	four work sites still generating new Part B
18	cases are Savannah River, Hanford, Y-12 and
19	K-25.
20	This slide, which is perhaps a little
21	difficult to read, but shows basically two
22	things of interest, I think. One is that DOE
23	site cases are slowly declining, in general,

and that AWE cases are slowly increasing.

Part of that is an artifact, I think, because we started with the DOE sites, and it took a long time to gather in all the AWE site information and act upon it.

Stu has already mentioned outreach events, which DOL, of course, participates in and sponsors, through the Joint Outreach Task Group. This slide is rather small, but -- has small text, but these are some of our recent meetings. I don't think there's any point in going through them unless a Board Member cares to ask. They're all on the website. And as you can see, there are quite a few meetings.

And we're now into FY15 of course, and here's the combined slide for both 14 and 15. We had three meetings in October alone. And our next outreach meetings, we see one in Los Alamos. Well, one, but it's on three dates in November. And the IBEW Union Hall meeting in Albuquerque on November 13th and 20th.

Now on the Santa Susana site, which is being discussed later today, just as some

1	background information, we already have
2	combined B and E claims or cases, rather,
3	1,000, approximately, of which NIOSH has
4	already completed a DR of 240 of them.
5	We have final decisions on 500 such
6	cases, which is an interesting thing, but I'm
7	not sure of the discrepancy between those two
8	numbers. But as it okay, earlier SECs,
9	right. And then we have 200 Part B approvals,
10	and 200 Part E approvals.
11	And that's the last slide, I think,
12	that needs to presented. The rest is, as I
13	said, general information about claim-filing
14	and qualifications. Any questions?
15	CHAIRMAN MELIUS: Any questions for
16	Frank? Anybody on the phone have questions?
17	Okay. Thank you. We'll now get an update from
18	Department of Energy. I'm not sure just
19	you, Greg, or how are we doing this?
20	MR. LEWIS: It's got to be me.
21	CHAIRMAN MELIUS: Oh, okay.
22	MR. LEWIS: You're stuck with me.
23	CHAIRMAN MELIUS: Put him on the

spot, huh?

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MR. LEWIS: Yes. All right, good morning, everyone. I'm Greg Lewis with the Department of Energy. And of course, Pat Worthington and Isaf Al-Nabulsi are also here with me. I guess while Stu is getting the slides up -- I was going to -- oh, I guess there again. Thank you.

All right. So the DOE mission is to work on behalf of program claimants to ensure that all available worker and facility records and data are provided to DOL, NIOSH and the Board. So basically we provide records. That's our role in the program.

We do that in primarily three ways. The first is with respect to individual records request. So we respond to employment verification requests from the Department of Labor, we -- and then I quess the second is to respond to large-scale records requests, like the Special Exposure Cohort projects. And third then the is to research covered facilities, particularly with respect to AWEs,

are primarily where we do the research.

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So the most important, you know, front lines that we have in terms of responding to these records requests are, you know, with the POCs. You know, some of you all who took the tour at the ETEC facility yesterday yet Phil Rutherford who was our primary POC.

still involved, He's but the contract, of course, has transferred over to North Wind, so those folks are going to be stepping up and doing the new remediation responses, where Phil and his team are still going to be handling, you know, the legacy Rocketdyne, the workers, the Atomics so International, those folks, they're going to still be handling.

So as you saw yesterday, Phil has been on-site for 25 years. So, you know, although the site goes back to the 50s, you know, he's been there for, not the whole the time but, you know, 25 years. He's been there quite a while. He understands the site. He knows the history, you know, as well as anyone does.

He knows what was done, where it was done, what these folks might have been exposed to, and where the records are, most importantly. So he's -- him and his team are the ones that pull those responses together.

So with the individual records requests, we do about 16,000 per year. As Chris Crawford alluded to, it has gone down slightly over recent years, but it's still roughly about 16,000. And it's split between the employment verifications, the NIOSH requests and the Department of Labor document acquisition requests, or DARs.

And as you know, you know, claimants often worked at multiple sites. They worked in multiple departments, divisions, held multiple job titles, particularly if they were career employees over 30 years. They might have been in a number of different locations around the site, job titles, things like that.

So often we have to go to a number of different locations to pull together those records, both -- you know, particularly for

historical workers, we might have microfilm, microfiche. We might have to go to multiple databases, you know, because on some of these sites as different contractors came in, they brought their own systems, their own databases, their own ways of doing things.

So, you know, at any given site, we might have to go to 20 to 30 different locations to pull together an employee's records. You know, and of course, these would be for the long-term career employees.

You know, again, we might have to go to 20 to 30 places, of many different types of records. You know, and unfortunately, this is not always true for the subcontractors. For the subcontractors, we check everywhere that we can but, you know, for a short-term construction contractor, you know, we might not have anything. So we try to be as creative as possible finding those records.

So we will look at gate logs when we have them, sign-in sheets, badging records.

If they were on-site and went to medical, or

happened to be badged for dosimetry, we'll check those type of locations. But often times we won't have a formal employee, you know, human resources employment record for those folks, so we have to be as creative as we can.

large-scale So for the records projects, know, research you those incredibly time consuming, as you know, and can be very difficult. They can take years. can cost us quite a bit of money. You know, and we work with you all and with NIOSH and SC&A to try to make sure that our resources are in place to be able to support these records research efforts.

Currently, we're -- these are some of the sites that we're working on, although we're getting requests for many more, whether it be for an SEC project or for a Site Profile update, you know.

And those listed there, some are, you know, just starting. Most are kind of in the tail end, or we're kind of doing the final stage of the research but we are supporting NIOSH and

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the contractors on those sites.

Then of course, I already mentioned that we provide site tours when requested. So yesterday we took Members of the Board, NIOSH and the contractors over to the Santa Susana field lab up by the Simi Valley and drove them around the site, took them into quite a few buildings, and for more of the buildings, showed where they used to be and what used to happen.

I think the numbers, there used to be 200 and something buildings there, and I think they mentioned there's somewhere around 20 buildings there that used to do DOE work.

So there's very little left compared to what they used to do, but we were able to show the group the layout, how the site is configured, the difference between Area IV, which is the facility that's covered by this program, and then Areas I, II and III, which were NASA and Air Force sites, which are not covered.

So we were kind of able to show the

layout, some of the buildings, what different processes were done and what's there now. So hopefully it was helpful to you all.

And then of course, document reviews, due to, you know, security and classification concerns, there are, you know, certain sets of documents that we have to review.

We've committed to do this. We have a security plan that kind of lays out what we do and how we do it. The average turnaround time for documents reviewed by headquarters is about eight working days.

It's not always that quick out at the sites because they're not always able to juggle their resources and their staff to accommodate the reviews, and also because the headquarters reviews are typically NIOSH-drafted reports or Board-drafted reports and they're shorter, whereas out at the sites we could be talking about hundreds of documents that are thousands of pages so it can be quite a workload for the sites.

We -- you know, when we get a

large-scale request, we will try to work with, you know, the requester or NIOSH, the Board, whoever it is, to determine at least a path forward to completion, what the priorities are, what we should work on first.

And we'll provide a time frame. And if that time frame is not workable for the Board or NIOSH, we can bring in assistance from headquarters sometimes, or try to work out alternate pathways.

And in fact, with -- recently with Savannah River, there was a very large request for declassification, and given the staff and resources at Savannah River, they were going to struggle to do that in the time frame that was needed on your end, so we've sent a large group of those documents up to DOE headquarters in Germantown.

And so folks can come in and work on them there. So as they're being reviewed and declassified, there will be a collection at headquarters that can be used to help draft the reports. So things like that, we do try to come

up with solutions that meet your all's needs.

And then facility research, you know, and that's kind of the smallest of our responsibilities in terms of workload, but it is extremely important, particularly with respect to the AWEs.

When questions are raised, and often those questions can come from NIOSH research into either a Site Profile or SEC, they might, you know, come across documents that suggest facility the should be covered for additional time period, or they can't find evidence that work was done during some parts the covered time period. We'll work of together with DOL to pull together documents and do some research into those facilities.

And then I just want to mention the SERT. I think I've mentioned it before, but again, this was our big accomplishment last year, bringing this on line. SERT is the Secure Electronic Records Transfer System.

It's an electronic system that DOL, NIOSH and DOE are all a part of. It allows us

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to send requests for documents, and then, you know, upload the responses and send them back instantaneously.

So it's two-factor authentication, which is basically a complicated way of saying it's the highest standard in terms of the protection of private information. I know that, you know, there's been many high-profile hacking incidents and data releases and things like that, so two-factor authentication is the, you know, is sort of the gold standard for protecting information.

So it allows us to quickly and efficiently send and receive information with these groups. It has cut down the time to respond by probably about ten days or so for each claim. And it also allows all of the different agencies to track and manage their responses and requests, you know, cohesively, so everyone has the same numbers.

And then both Chris and Stu have mentioned outreach. Again, we do participate in the Joint Outreach Task Group meetings. And

we've had quite a few meetings this fall and we'll be having a few more in the winter and spring.

And then I also want to mention the other program that we work on at DOE that isn't directly tied to EEOICPA, but it, you know, I kind of consider it a sister program. It's our Former Worker Medical Screening Program.

We provide free screenings to all federal contractor and subcontractor workers at DOE sites. Again, that's not the AWEs, but at all DOE sites, you're eligible for a free screening. We work to provide you with that screening close to your house.

And have specially trained we that occupational medical physicians familiar with the DOE sites, that know the hazards that you might have been exposed to and are able to custom tailor a screening to your particular situation. And I've got website up on there for anyone who wants more information.

And then, of course, for this area,

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1	the two programs that would cover the Los
2	Angeles area are our Supplemental Screening
3	Program for all production workers and our
4	Building Trades National Medical Screening
5	Program for the construction workers.
6	And with that, questions?
7	CHAIRMAN MELIUS: Paul, go ahead.
8	MEMBER ZIEMER: This is not really a
9	question, but I just wanted to highlight the
10	tour that DOE hosted of the Santa Susana Field
11	Lab yesterday. Excellent tour, and I think we
12	owe Dr. Worthington and Greg and the staff a
13	word of thanks for really an excellent tour, one
14	of the best ones we've had.
15	MR. LEWIS: Thank you. And we know
16	these tours are important, so any time you
17	know, we're always happy to work with our sites
18	to facilitate these tours for you.
19	MEMBER ZIEMER: And Isaf, too, is
20	here. So include all of the DOE folks that are
21	here.
22	CHAIRMAN MELIUS: Any other Brad,
23	you don't have a usually you have something

you want to pin Greg down for. 1 MR. LEWIS: Taking it easy on us. 2 3 CHAIRMAN MELIUS: We're doing fine We do appreciate, both the tour -- I 4 now. 5 heard, as Paul said, rave reviews this morning while eating breakfast, from people that were 6 on the tour, but also I think everything's been 7 going well with the sites. 8 9 I think the -- at least my impression of it seems to be that we have a number of sort 10 11 of special focus now. That list you put up, 12 somehow I feel that I'm not sure we're really 13 fading away in terms of the records requests from some of those sites, Savannah River and 14 Idaho and --15 MR. LEWIS: No, well they seem to be 16 17 coming fast and furious. 18 CHAIRMAN MELIUS: -- Hanford to do 19 things, get turned over. But I think, again, 20 that's good. We need to get caught up on 21 everything, so good. But we appreciate the 22 assistance and cooperation and so forth with

Anybody else -- any of the

the, just that.

1	Board Members on the phone have questions?
2	Okay.
3	MR. LEWIS: Thank you.
4	CHAIRMAN MELIUS: Yes. We have a
5	mellow Board today. Yes. It's the barbed
6	wire surrounding us. Right. Oh I see, we're
7	saving it all up for the last presentation here.
8	Okay. We'll now get an update from
9	our Procedures Subcommittee, and their review
10	activities, and particularly one Program
11	Evaluation Report, Construction Trades
12	Workers. So I'm not quite sure how this is
13	going to work, but
14	MEMBER MUNN: I'm not sure either.
15	CHAIRMAN MELIUS: Okay.
16	MEMBER MUNN: But all right. Thank
17	you. You're all very familiar, I think, with
18	this material. We've certainly talked at
19	length, both here and in Procedures
20	Subcommittee, with respect to the construction
21	trade workers and how we view their dose
22	reconstructions.

We start with the first slide, which

is one you're quite familiar with. 1 think we need to go over this in great detail. 2 This is just your update with respect to what 3 the audit process for the DERs actually is. 4 Ιt consists of five different 5 subtasks, and the first of which is having our 6 contractor evaluate NIOSH's assessment of what 7 They take a look at specific the issues are. 8 9 methods that NIOSH has used, and they take a 10 look at the stated approach. Generally we see those three subtasks 11 12 in the first initial report, and Subtasks 4 and 5, which are 13 fairly time-consuming, unexpectedly, sometimes come afterward. 14 15 We started this process for the construction trade workers back in 2004. 16 that time, NIOSH brought to the fore the fact 17 18 that there might be some construction trade 19 workers at various DOE sites that hadn't been 20 monitored but had been exposed. 21 At that time there was a considerable discussion with respect to who would be covered 22

by the designation construction trade workers.

We specifically included laborers and mechanics, masons, carpenters, electricians, painters, pipefitters, boiler-makers, millwrights, sheetmetal workers, iron workers, insulators. And that's just a partial list. That's not all-inclusive.

In order to make sure that we had the appropriate process in addressing these issues, we were very pleased to have NIOSH provide for us OTIB-52, their first OTIB with respect to the parameters that needed to be handled when we were going to look at construction trade worker issues.

That was issued on 2006, and that was the founding document which we have used since then in order to address these issues surrounding this particular group of workers.

I think I went too far. No, I didn't. All right. At the time that this model was provided for us, PER-14 was issued to reevaluate the claims that had already been done at ten sites, where external coworker models had already been looked at.

Those are the ten sites; they are Hanford, PNNL, Kansas City, LANL, Pantex, Portsmouth, Savannah River, Weldon Spring and Oak Ridge Labs, Y-12.

Those were not the only sites, of course. There were four other sites for whom coworker studies had been published prior to the issuance of OTIB-52 in 2006. Those claims had already been reevaluated, however, under other PERs.

You see the sites: Paducah, PER-13, Rocky Flats, PER-21, Mallinckrodt, PER-15 and K-12, PER-11. So because those had already been evaluated by PERs, those were not included in the ten that were covered by PER-14.

The timeline is fairly straightforward. I think we covered that reasonably well. In 2007 was when PER-14 was issued first, and it's been under evaluation of one sort or another since then. In 2012, SC&A had its draft review of the entire PER, and in July of that year their six findings were discussed at length, and eventually, just

recently this year, all were resolved.

Our Subtask 1 issues, assessing the circumstances that necessitated the need for the PER to begin with. You'll recall that's the basis for Subtask 1 from our first slide.

During facility modifications, we might have had exposed construction trade workers who hadn't been monitored. The exposures might be different from other radiation workers, and the assignment of coworkers to unmonitored construction trade workers needed to be claimant-favorable.

The empirical data ratios were developed from both external and internal doses, by monitored workers to all monitored workers, using data from seven major DOE sites, which are listed there. The results were the basis for the OTIB-52 guidance.

Subtask 1 focused on both external dose, using -- deriving deep-dose coworker adjustment factors of 1.4 from the empirical data that was available. A shallow dose was bounded by the workers' doses, by the AM

workers' doses, and the 95th percentile of coworker data could be used without having to apply an adjustment factor, for the shallow dose only.

In the internal dose, only the Hanford coworker intakes need be multiplied by a factor of 2. For all the other sites, the internal dose was going to be assessed using the same method that's applied to all other workers.

I apologize for these slides where we give you the finding information. You know, we always have the same process when we are presenting PER information to you.

We want you to be able to see the finding and the approximate solution -- and resolution to that finding at the same time, but if we put each one of these on a different slide, then I end up having 75 slides here, and I don't think you want that any more than I do.

So I'm going to have to ask you to bear with us with respect to the way we present these findings. It seems to have worked well in the past. If you don't find that amenable to your

needs now, please let us know. 1 We're hoping that providing these 2 slides gives 3 you in advance you 4 opportunity to read through this at your 5 leisure, and making them available to the public makes it possible, we hope, for them to 6 be able to address that in the same way. 7 But if you find this onerous for any 8 9 reason, please let me know what your desires are 10 with respect to presentation to you. That being said, Subtask 1 findings, 11 12 of which we have four shown here. One, two and three were conditional. 13 Number 4 is a true finding. These have to do with the deep-dose 14 15 adjustment factor of 1.4. The inclusion of the construction 16 trade workers with all monitored workers might 17 18 obscure the dose differences, and a shallow 19 dose adjustment factor may be required if NIOSH 20 failed to adjust for the construction trade workers' shallow doses. 21

guidance

internal dose says the internal dose should be

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OTIB-52

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determined using the same method as is used for all the other workers in the absence of internal monitoring data and coworker data. It's unclear what was meant by that recommendation.

So these are the types of things that we debate at considerable length in the Subcommittee proceedings. If you have any question about these that we can't answer for you, please do -- I refer you to the Subcommittee's transcripts, which give you the detailed information about the discussions that were had with regard to them.

A conditional finding, as the footnote indicates, it's just indicating that SC&A didn't have access to the original dose data, and would -- didn't know if the data that they had was all completely accurate. And it was established for efficiency purposes, so that we could have the issue in front of us, even though there was some question about whether it was really an issue or not.

CHAIRMAN MELIUS: Excuse me, Wanda.

MEMBER MUNN: Yes.

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1	CHAIRMAN MELIUS: I think David
2	Richardson had a question.
3	MEMBER MUNN: Yes, David?
4	MEMBER RICHARDSON: Wanda, if just
5	help me understand the first Subtask 1, the
6	problem was laid out that the construction
7	trade workers might have been exposed but not
8	monitored, and their exposures might have been
9	different from monitored radiation workers.
10	MEMBER MUNN: Yes.
11	MEMBER RICHARDSON: And I thought it
12	was useful, what you took one step towards
13	that which was to say that there was some
14	monitored construction trade workers, and you
15	compared them to the monitored radiation
16	workers.
17	MEMBER MUNN: Yes.
18	MEMBER RICHARDSON: But then it opens
19	the question, are the unmonitored construction
20	trade workers like the monitored construction
21	trade workers? How did you answer that
22	question?
23	MEMBER MUNN: Do we have one of our DR

experts who had addressed this issue for us? 1 DR. NETON: We have the expert. 2 There he is. 3 MEMBER MUNN: DR. NETON: I'm sorry. That wasn't 4 5 done. That comparison was taken at the face If construction trade workers had a 6 value. 7 higher, on average, dose than the regular workers, the adjustment was applied without any 8 9 correction at all, or any evaluation of what the 10 status of the unmonitored workers actually was. Because that --11 MEMBER RICHARDSON: 12 it's -- I mean, I -- to me, I mean, I've felt 13 some kind of struggle with the same problem, and I think what you've done is extremely valuable. 14 15 It's -- the counter-argument that 16 I've made in my head is that there are people 17 I guess part of this gets to the who --18 definition of a construction worker, there are 19 people at many of these sites who hold jobs with 20 titles like painter or pipefitter, who were 21 monitored. And yet they may have 22 employees of the prime contractor, and they

were monitored for that reason.

And then there were a lot of people who we often conceptualize as construction workers, who were employed as subcontractors doing construction work, and they were not monitored.

And whether somebody who has a job title of painter or pipefitter or so on, who worked for these subcontractors and falls into this large group of unmonitored workers who were doing the same sort of tasks and facing the same sort of occupational hazards as the monitored workers has been an open question in my mind.

DR. NETON: Yes. That's a really tough question. I think we're maybe going to get into that a little bit later when we deal with the coworker model and sufficient accuracy and how that all plays out. There's an implementation guide that we have in draft form that we're going to discuss at a later session this morning.

This TIB-52 was our very, very early attempt -- I think this document was issued in

1	2006, of trying to address these types of issues
2	that you're bringing up. And I admit that it's
3	somewhat crude's probably not the right word
4	but, you know
5	MEMBER RICHARDSON: But it's
6	DR. NETON: used the data that we
7	had available.
8	MEMBER RICHARDSON: So it's
9	leveraging the assumption that the
10	construction workers who were monitored are a
11	simple random sample of all construction
12	workers
13	DR. NETON: Right.
14	MEMBER RICHARDSON: and that you
15	can make the extrapolation from these data to
16	the unmonitored
17	DR. NETON: Yes.
18	MEMBER RICHARDSON: I guess that's
19	the only thing. It's just to be explicit on
20	that.
21	DR. NETON: Yes, exactly. Yes.
22	CHAIRMAN MELIUS: Can I just add I
23	think we're going to be coming back to this

issue, as Jim described in his presentation, later today, because I think we're sort of re-looking at the whole coworker issue.

And plus, I think we better understand, maybe, the -- how the variability between sites and these issues and also the lack of good information on which construction worker fell into which category, because they all come out of the same union, they often could have been an employee of the prime contractor and then also, before or after that worked for other subcontractors.

I mean, it's just a very complicated picture to do, and I think we're going to have to look at what information's available and what's the best way to do that.

And I suspect it's going to be on a site by site basis, but I think we're going to have to sort of -- I think if we can come to grips with the, sort of the coworker model issue and what the criteria ought to be for evaluating that, I think we'll make progress on this.

Wanda and I went back and forth a

little bit on what would -- what to present today, in terms of this, and I think the focus is going to be, you know, what I'm presenting is, there were a number of other specific findings as part of this review, that I think are relevant, aside from the coworker issue, which we'll be coming back to.

Subtask 2 was assessing MEMBER MUNN: methods and corrective actions. The OTTB requires multiplying external coworker dose by 1.4 for the construction trade workers and at Hanford, multiplying by 2. But the claims at other sites where the coworker studies externally were issued prior to 2006 had to be reevaluated.

Identifying all the workers that had been exposed as a member of the construction trades, NIOSH went through NOCTS and all the original DR reports. They used a 31 word key search, and they identified 977 claims that might be potentially affected.

They reviewed the list of sites when they established the coworker models, and used

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the key word list, determining that their screening methods were going to be adequately inclusive and complete.

SC&A didn't have any findings under our Subtask 2 review, so we'll take a look at Subtask 3, evaluating the approach for identifying the number of DRs that were going to require reevaluation.

To check to see if the 977 potentially affected claimants were going to need reworked DR, NIOSH had applied the screening criteria of confirming that a claim was a construction trade worker, because the key word search doesn't always do that appropriately, to verify that the external coworker dose, or in the Hanford case, the internal dose, assigned in the original dose reconstruction. To screen the claims based on the ability to raise a PoC that was equal or greater than 45 30 percent, to ensure that IREPs were performed, and that would automatically be triggered by an original PoC of 36.8 or 29.0 from Hanford, any claims with PoCs less than the

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trigger value, to determine whether any other PERs might increase that dose.

So under Subtask 3, we had two findings, one of which was conditional. The first of those findings, which was Number 5 for these purposes -- remember, we had four under Subtask 1, Number 5 was the assertion that the PER was incomplete and that the extent of the screening and evaluation of the universe of those 977 claims was not fully discussed in the PER.

So they asked -- indicated they didn't feel NIOSH had identified the actual number that were eligible for PER dose adjustment factors. We did resolve that. We looked at it closely, and the criteria that would -- they'd used to request, that NIOSH would use to request, was shown there with the 977 totals, and how those were broken down.

It was acceptable with the explanation that was given. And the conditional, which we had indicated was Number 6, would -- may be highly restrictive in

addressing the problems of these unmonitored workers and the uncertainty of the fate of the claims that had been adjudicated before the issuance of a coworker model.

I read that badly, but you recognize,

I trust, what the conditional finding was

concerned with. It's one of the issues that

we, again, discussed at considerable length but

were able to resolve satisfactorily.

Subtask 4 gets to be the sticky wicket where you get into real audits. Referencing a finding that we discussed earlier, Finding 5, in respect to the possible, potential failure to identify that adequate number of claims out of the 977, it was determined that regardless of what the number was, SC&A felt that they should have one case from each one of the ten sites that had been impacted by the PER.

And again, those sites are listed there. They're the same, I believe, as the ones you saw listed in Subtask 1.

Timeline for this crucial Subtask 4 began in 2012 when SC&A was asked to review one

case from each of those 10. In addition to that, we also asked them to evaluate the site TBDs and workbooks that were applicable to what we were doing here, to make sure that they had been properly updated and that they fit the recommendations for construction trade workers.

So during their review, the contractor determined that there were no reworked cases at four of the ten sites, and so for those ten -- for those four sites, Kansas City, Pantex, PNNL and Weldon Spring, they were going to need to be limited to just verification that the TBDs and workbooks had been updated. They didn't actually have cases to look at.

In 2013 we received SC&A's draft of their subtask for review, and later that year, we had, at Subcommittee meetings, discussed their findings and were able to resolve all of the findings from them.

Here is Subtask 4's review of the sample sets, the DRs that were looked at by the PER. Out of the 977, as you can see in the first

table there, the selection criteria that were applied, the first of those items was NIOSH requesting that the case not be returned for a new DR. There were 620 of those.

The cases that were requested to be returned based on some other PER other than 14 that we were working with, there were 221 such cases out of those 977. The cases that were requested to be returned for a new reconstruction were 52 in number, and there were 84 cases that had been returned to NIOSH prior to completing the PER evaluations.

Underneath, the selection criteria for the site are listed for you for each of those ten sites, so that you get the breakdown two different directions, if you read the criteria and the breakdown shown by site.

The findings that we had, Findings 7, 8, all the way to 21, but because of the number of sites that were involved, many of these findings were applicable to more than one site, and so some of them were grouped together.

We've done that grouping as we go through here

in this presentation, and you'll see them.

Finding Number 7 -- and remember, we're under Subtask 4 here, this is the actual audit process that we're looking at, SC&A found many of the cases identified as requiring rework didn't meet all the requirements of the selection criteria.

NIOSH indicated they'd reviewed all the potential cases of less than 50 percent to determine if they were affected by another PER. SC&A, in Finding 8, noted that some of the cases had been returned as a result of this PER, but those cases were not revised.

And NIOSH indicated that not all cases had been returned by Labor, and since some of those are contained in an SEC, or the claimant has died and there's no survivor. But each case had to be individually verified in order to be included in the cases that were going to be -- come forward for review for the PER.

Subtask 4, case reviews, involved an audit of six reworked cases from -- one each from Savannah River, X-10, Portsmouth, LANL,

Hanford and Y-12.

That review focused just on the doses that were impacted by this PER, and they were concerned solely with assessing the accuracy and correction -- correctness of the coworker external doses. For Hanford, both coworker external and internal doses had been evaluated.

Continuing with the case review findings, this next finding, as you see, is a group of actually four findings lumped together. As I indicated to you earlier, many of these findings were repeated because the same finding was applicable to more than one case, but we lumped them all together when we were looking at them.

This one has to do with the 1.4 adjustment factor being applied to the measured coworker data at each site. And after discussing them and looking at them at considerable length, the contractor and NIOSH agreed that we had adequate empirical evidence to indicate that the 1.4 adjustment factor had been appropriately applied, so this finding

resolved.

Finding Number 12 had to do with whether NIOSH is planning to revisit the one returned case for a construction trade worker coworker dose at LANL. And as it turned out, DOL did not return that case to NIOSH, so -- because the worker had qualified under an SEC.

Finding Number 13 had to do with the correction factor not being applied to a LANL coworker dose, and it turned out that in this case, the worker had a job title on the list, but, after looking at the CATI looking pretty closely, it was shown that the claimant actually was an in-house employee and not a construction trade worker.

Finding Number 14 had to do with application of the construction dose factor for -- dosimeter correction factor in coworker dose, and there was agreement that the dose correction factor of 1.244 hadn't been applied to an unmonitored photon dose.

The correction factor was one during this period and wouldn't impact the dose. And

we had a file indicating that the impact of the finding on the case outcome didn't change the level of compensation, so that was resolved.

And Finding Number 15 was another dosimetry uncertainty, it hadn't been applied to a Y-12 coworker. And NIOSH didn't do that. Their explanation was, the value would be entered into IREP as a mean of the normal distribution with a 30 percent uncertainty. So the TBD was re-evaluated by the contractor, and they recommended the finding be closed. The Subcommittee agreed.

Finding Number 17, again, а construction trade worker correction adjustment factor, failure to do that to an unmonitored internal dose at Hanford. And adequately explained that was with the employment in 1944, intakes being based on air monitoring.

And when the case was reassessed by the contractor, the technical documentation, and recommended that -- they recommended the case be closed. We agreed it was appropriately

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resolved.

Finding Number 18, the contractor felt that there didn't appear to be any Hanford-specific technical guidance documents that required implementing OTIB-52 for internal coworker doses. But NIOSH indicated that the OTIB-52 requirements were built into the reconstruction tool, which was used by all the dose reconstructors. And that was found to be the case, so we closed the finding.

Again, one of those group findings from three different sites, Kansas City, Pantex and Weldon Spring, Findings 19, 20 and 21, was a concern about whether there were guidance documents or workbooks for implementing the dose adjustments that had been listed in OTIB-52.

And the response indicated that the requirements had been built in through the tool which was used by the dose reconstructors, and that was found to be the case and it was closed.

That's a full review of what we've done with PER-14 and where we are right now.

Questions? 1 CHAIRMAN MELIUS: Gen? 2 MEMBER ROESSLER: This was a lot to 3 absorb, but I do have one question on Slide 8, 4 5 and it appears on other slides too. It seems that Hanford internal doses are different than 6 some of the other sites. 7 On Slide 8, you talk about -- you say 8 9 only Hanford coworker intake rates needed to be 10 multiplied by a factor of 2. What's the difference at Hanford? What makes that site 11 12 different? Is it because they started 13 employment earlier and --14 MEMBER MUNN: No, not entirely. 15 It's the scope of the activities at Hanford more than anything else. The level of -- I should 16 17 say the scope, also, of the type of materials 18 that were handled. If anything was going to be 19 handled at all, it was undoubtedly passed

And because of the types of materials, because of the level of activity, as well as the periods of activity, this decision was made

through the workforce at Hanford.

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relatively early that a dose correction factor 1 of 2 was going to be implemented at the Hanford 2 I believe that has been done across the 3 site. 4 board. 5 CHAIRMAN MELIUS: There's some Yes, Jim, do you want to -- yes. 6 others here. I could just elaborate a 7 DR. NETON: These were, you know, just little bit on that. 8 9 empirical data sets that we had, you know, 10 construction trades versus regular workers, and using the available data, it just came out 11 12 that way. We really made no judgment as to why 13 that was particularly true. Hanford was the only one of the sites 14 15 we evaluated, though, that did have that difference. I'm not sure I really understand 16 17 why it existed there, but that's the way we 18 treated it. It was just purely based on 19 empirical evaluation. 20 other CHAIRMAN MELIUS: Any 21 questions? If not, I have a -- it's more of a 22 comment than a question. Findings 9, 10, 11

and 16, and then later on in -- towards the end

in Findings Number 19, 20 and 21, the -- it seems that we're referring findings back to the other site, individual site documents, or to individual Site Reviews to be addressed.

And I think -- again, it's not a fault of the Procedures Committee but, you know, we do these reviews on multiple levels, and we've always had problems with when things get referred, or assuming that, you know, another Work Group or that NIOSH will address it, or someone will address the site-specific finding.

And I think we -- in this case, we have a number of these sites that are currently under active review. We have some that are under inactive review. I mean, Brad's doing, I think, still working on some stuff at Pantex. We have Hanford, which we're actually mostly focusing on SEC issues still, and I won't go through the whole list, Kansas City, and so on. And I think we just need to make sure that this all gets communicated and that NIOSH also be aware of these issues and SC&A also.

It's not clear to me that these, 1 either in the case where there's been a problem 2 found or where it's been the construction 3 4 worker adjustment is subsumed under 5 instructions for doing individual dose reconstruction, that these get, you know, 6 properly evaluated and reviewed. 7 They may be, they may not, and I think 8 9 we just sort of need to formalize that process and make sure there is some follow-up on these 10 11 issues. Dave? 12 MEMBER KOTELCHUCK: Okay. If I can 13 get through, I can do this. 14 So, I'm not quite sure where this 15 leaves us. Have -- would you say that the PER audit process has been completed for the 16 17 construction workers, or are there next steps, 18 and what are they? 19 MEMBER MUNN: You would not have me 20 here giving this review for you if we had not 21 fully completed our review of the PER. Subcommittee is done with PER-14. And what we 22

have is now history, and you have it.

1	MEMBER KOTELCHUCK: Thank you.
2	MEMBER MUNN: You bet.
3	CHAIRMAN MELIUS: And I think my
4	comment was saying that, you know, the PER has
5	been review has been completed but we
6	there are findings that still need to be
7	addressed in other venues or other groups and
8	so forth. And so we should not lose track of
9	that.
10	MEMBER MUNN: Yes, our Chair says yes
11	but.
12	CHAIRMAN MELIUS: I mean, we have a
13	similar problem with individual, you know, dose
14	reconstruction reviews where we find a Site
15	Profile issue or something, and we just need to
16	make sure we connect back rather than coming,
17	you know, ten years later finding
18	MEMBER LEMEN: Right.
19	CHAIRMAN MELIUS: discovering the
20	problem again.
21	MEMBER LEMEN: So in addressing your
22	question, who has that responsibility?
23	CHAIRMAN MELIUS: We do.

1	MEMBER MUNN: Yes.
2	MEMBER LEMEN: I mean, how are we
3	going to keep track of it is what I'm asking.
4	I mean
5	CHAIRMAN MELIUS: Well, we
6	MEMBER LEMEN: is somebody going
7	to follow this?
8	CHAIRMAN MELIUS: I think all the
9	Board Members have some responsibility for all
10	the sites here. And I certainly took note of
11	Hanford.
12	MEMBER LEMEN: Should it be a
13	continuing agenda item or?
14	CHAIRMAN MELIUS: And I would hope
15	that SC&A and does that, and NIOSH also.
16	MEMBER LEMEN: Yes.
17	MR. KATZ: Let me just add to what Dr.
18	Melius is saying. So I generally, when one of
19	these findings comes up, that we're going to
20	refer, from whatever Subcommittee or Work Group
21	to somewhere else, generally either I or the
22	Chair will send an email to the Chair of the
23	group it's being referred to, saying this

finding, and provide -- I'll provide a transcript for the discussion as well.

This finding is being transferred to your Work Group to resolve. So that's part of the process that goes on here. I think one thing that'll help, though, it doesn't -- that doesn't necessarily make it easy to track. So, because then that Chair has it, but that doesn't mean it's easy to track.

So what we have in -- we're trying to put in place, but it's difficult because a number of these Work Groups are longstanding, and we're putting into place a system, you know, midstream for those, but as Stu has discussed at a number of these Board meetings, we have this tracking system that we started off using just for Procedures.

Now we've expanded it and we're using it for Dose Reconstruction Subcommittee, and we want to use it -- and for new Work Groups, and we want to use it, ultimately, for everyone.

That tracking system, actually, is great for just this problem, because that

finding sits there unresolved for all to see until it is resolved. And that way we won't lose anything.

So to the extent we can move towards using that system, I think we'll be better off for just this problem. Because it is tough. It is tough for everyone to keep track of these items. A number of these chairs are chairs of a number of Work Groups and Subcommittees, and it's a lot to mind.

MEMBER MUNN: And it was an extremely painful process for us to get that database up and running. We spent a disproportionate amount of the Board's time giving you reports on our blow-by-blow, step-by-step process to do that. But we -- our IT folks have been very diligent in helping with this.

And in the cases that we have in front of us right now, in almost all cases -- I do believe I can safely state in all cases, these dose reconstruction factors, the correction factors that we were talking about, have been very carefully applied in all of the tools that

1	are used by the dose reconstructors.
2	So as long as the claim is identified
3	as a CTW, it's I personally have, am well
4	reassured that the tools that are available to
5	the dose reconstructor will be adequate for
6	that purpose, will catch that. Any other
7	questions?
8	CHAIRMAN MELIUS: Any Board Members
9	on the phone have questions? I don't want you
10	to be forgotten. Okay, if not, it's almost 10
11	o'clock. Why don't we take a break and we'll
12	reconvene at 10:30.
13	(Whereupon, the above-entitled
14	matter went off the record at 9:59 a.m. and
15	resumed at 10:36 a.m.)
16	CHAIRMAN MELIUS: Okay.
17	Before this deteriorates any more we'd better
18	reconvene. So between now and lunch we're
19	going to be talking about sufficient accuracy
20	and coworker modeling, and do that.
21	I think, as we've talked at the last
22	Board meeting and a little bit on the last Board
23	meeting call, we've been working the SEC

Review Group's been working with Jim Neton,
NIOSH, and SC&A on addressing both sufficient
accuracy issues but with more of a focus,
recently, on coworker modeling issues.

And we've gone through, well one meeting just before the, our Idaho meeting, you know, the day before, and we had another conference call a few weeks -- a couple of weeks ago on this, to do that.

And Jim has -- Jim Neton's been working through a document describing sort of an approach to developing coworker models, and a sort of set of guidelines, I guess we will call them, similar to sort of the guidelines we have for reviewing surrogate data, reviewing SEC Evaluation Reports.

So that's all. It's not totally prescriptive, but the idea is to try to get what are the key factors that will be used in developing and therefore the key factors that would be used in evaluating the coworker models.

So as I said, we had a meeting, what

two weeks ago, something like that, and Jim's done some updating then. The plan is that Jim will do a presentation, go through sort of the key points in the document. We'd like to get your, all the Board Members' comments here today. And then we'll ask everyone on the Board to also, you know, when you have time to review the document, some time in the next couple of weeks get comments in to Jim. And we'll do another revision, and hopefully some time in the near future we'll get this finalized.

I'm hoping, certainly by the March meeting, I think we'll be able to do that. So that's sort of our target, but there'll be some other iterations as we go along. So let me turn it over to Jim Neton.

DR. NETON: Thank you Dr. Melius. As Dr. Melius said, this is something that's been going on for quite some time now. It started shortly after the release of ORAU Report 53, which was a report that described how to analyze stratified data sets.

And SC&A had a number of findings on the statistics and such, and going through that it became apparent that, well let's first get an idea of how we evaluate data to see if it's stratified in the first place, and sort of, not bypass the statistics but talk about in general, how you look at, how you approach coworker data sets.

And so we -- I volunteered, and we've been working hard on putting together an implementation guide, we're calling it -- right now it's a draft, on exactly that. What criteria are used to evaluate these data sets that go into coworker models. And we're up to Rev 3 now, and I will go over that in a second.

But before I get into that, I thought it might be useful for me to go over sort of a case study, if you will, of what -- how a coworker model is developed. We use them a lot.

We -- almost -- many, many sites have them, and it became sort of obvious to me, or apparent to me, at our Working Group meeting in

Idaho that there wasn't a general, you know, 1 general understanding of what we mean, how we 2 go about establishing a coworker. 3 So I'm going to briefly go over -- I 4 5 have like six or seven slides, and just sort of go over, you know, what -- how is a coworker 6 7 model developed. I'm going to use the internal dose example because they tend to be the most 8 9 complicated. External is a little troublesome. 10 11 So here goes. So you've seen this 12 before, and this is right out of Report 53. 13 This is the -- these are the, this summary of the steps that go into an internal coworker 14 model calculation. 15 You start with Box 1 on the upper left, 16 17 which is the urine data, just a database you get 18 from the site that has all the urine data over, 19 let's say, the history of the plant, from '57 20 to 2007 in this case, is what we're going over 21 today. 22 But we need to do something with that,

to apply to workers that didn't have any

monitoring data. And we start that by doing this OPOS analysis, this one person, one statistic analysis.

We're going to talk about that a little later, but what that means is if you have multiple data points in one time interval, in one year, you need to somehow account for that, group them together so you have one value in that one monitoring period. We'll talk about it a little later.

There was some disagreement between SC&A and NIOSH and the Working Group on that. I think we've come to some agreement.

The third step is to take those OPOS data, the one person, one statistics data, and generate these distributions. You take the log of the data and you do a cumulative probability plot, and then you look at how they fall on a normal distribution.

And from that you can generate the 50th and 84th percentile of the data. So that can characterize the data for that one particular time interval, one year, three

months, whatever data set you have.

The fourth step is where you really start getting more detailed. You take the 50th and 84th percentile excretion rates, what the person was excreting, and you convert that into what they were actually inhaling. That's where the IMBA program comes in. That's Step 4.

IMBA, Integrated Modules for Bioassay is a very sophisticated computer program that can take urinary excretion data and say, what were these people actually breathing in over this time period if the exposure was chronic. A key to these coworker models for internal is that, for all the models we've developed so far have been chronic intake models.

You use that to generate the 50th and 84th percentile intakes, what these people were breathing in, and then you can use that for workers, an unmonitored worker, to calculate what their internal dose was over a certain period of time. I've got some examples.

And then, of course, the sixth step is

you'd use the dose calculated to the organ to generate some Probability of Causation result. So this would all be -- we take monitored worker data and try to apply it to unmonitored workers.

This is what I was talking about. Let's see, Step 3 here is generate the 50th and 84th percentile urine data. I just have an example here of what one of those distributions might look like, where you see the geometric mean and the 84th percentile. You've got 196 individual workers represented here that left 332 samples in a single year.

So you'll have one of these for every single year in this particular case. It could be three months. If we had -- if urine samples were taken every three months, we would have, every three month, a graph like this. But typically, a year seems to be about the most common monitoring period.

So you have a graph for each monitoring period, and you generate graphs, in this case for Savannah River, from 1955 through 2007. These are the distributions of urine --

urinary excretion over the entire monitoring history that we have.

And this is Table A-3 right out of the Savannah River Site. I think it's in the Site Profile. I don't think they have a separate -- I think it's a TIB. This is a TIB.

Anyway, these are the data that were generated. This is the real data. So you can see, from the middle of the year, July 1st, '50 -- actually '55. I've only presented here through 1965, or '84, but it continues on through 2007. I didn't give you the entire sheet.

But you can see that you can generate the 50th percentile, the 84th percentile for every particular year. And now these are the data that are going to go into IMBA, the Integrated Modules for Bioassay Analysis, to estimate what the people actually breathed in during these periods with these data sets.

And this is where it gets a little tricky to explain, but the blue dots, in this particular case, are the 50th percentile data

points from that graph. So that second column there from the left, these are the 50th percentile data points presented over the monitoring period that we have, for which we have data.

Now you can see that there's two separate type, two separate sort of collections of data, the blue dots and the red dots. The idea here is that as long as you have a monitoring period where the data appear to be similar, where the exposure -- the excretion patterns were similar, that represents what we call one intake regime.

And then, so you would have Intake Regime 1, the blue dots, and Intake Regime 2, the red dots. So we will fit a chronic exposure model through the blue dots up through where the red dots start.

And so you can see that increasing black solid line. It starts down by zero because when you first start work, you start -- it starts going up because you start inhaling material. And then it stops at the

intersection of the blue and red dots, and starts to decrease.

That's because once the intake regime stops, we're saying there's no more exposure, but the person will still continue to excrete uranium that they had -- or plutonium, in this case, that they had inhaled in the earlier period.

So that's Intake Regime 1. Now you go to the bottom one where you say Intake Regime 2, and there's another chronic model fit to that. So here is an example of where we fit data, those 50th percentile data points, to the data. There is some subjectivity involved in this, but I'm not sure there's any way around that. Okay.

So this is the complete data set, so you can see all the data including the first and second regimes. And what's interesting, you really can't see it very well, but there's a green line there, the solid line.

That's what the excretion would be if the person inhaled plutonium from day one of the

plant operations all the way through 2007.

That's the combination of an intake from Period

1 and Period 2.

And here is, taken right out of the TIB, the values for those intake amounts. So between '55 and '90, the person would receive -- would get about 1.8 picocuries per day intake. The 84th percentile is 5.1. You fit the same curve at the 84th percentile to get that value. And the GSD on this particular fit is 2.88.

You see the value that says adjusted GSD? That is the minimum geometric standard deviation that we will allow for an internal exposure, because there is a lot of uncertainty involved in internal dosimetry, and based on some references that we've obtained from the literature, it seemed to be 3 is about as low as you can go.

So we won't assign any GSD less than 3 for internal exposures. So everyone will have that uncertainty associated with it.

And then you see the second period,

1991 to 2007, it's a little lower, 0.9 picocuries per day. Interesting, what you see here is the change in the out -- in the urinary output really is more of a function of the detection limit of the measurement system rather than changes in work -- necessarily changes in working conditions.

I think they probably went to alpha spectrometry back then, in 1991. Detection limits went down, and so you have a much lower median value, because many of the 50th percentile values that we calculate are usually right around the detection limit of the measurement system. And that's pretty much true in this case.

So, that's just a sort of a quick run-through of how a coworker model is established, just so everybody has a feel for what we've been doing for a long time now, at least in the internal dosimetry world. Is there any questions on that before I move on to the implementation guide?

CHAIRMAN MELIUS: Yes, Brad?

1	MEMBER CLAWSON: I just all the
2	dots, they represent the whole spectrum of
3	workers, or
4	DR. NETON: In this particular case,
5	this would be all the workers. Now it could be,
6	if you decide to have some sort of a strata, it
7	would represent the 50th percentile of all the
8	urine values for that particular strata.
9	This is a general model for all the
10	workers, you're right. And we're going to talk
11	about how we might make some decisions about how
12	to partition or stratify these in the different
13	data sets. That's the, sort of the point.
14	MEMBER RICHARDSON: Can I ask you
15	just a follow-up question.
16	DR. NETON: Yes.
17	MEMBER RICHARDSON: A dot is not a
18	worker but is
19	DR. NETON: It's the 50th percentile
20	of the urinary excretion of the workers in that
21	year.
22	MEMBER RICHARDSON: A dot represents
23	a year?

1	DR. NETON: Right.
2	MEMBER RICHARDSON: And it's
3	DR. NETON: In this case
4	MEMBER RICHARDSON: median value
5	of excretion in a year.
6	DR. NETON: Right.
7	MEMBER RICHARDSON: And you had
8	one slide back you had two colored dots.
9	DR. NETON: Right.
10	MEMBER RICHARDSON: And could you
11	tell me once more the transition?
12	DR. NETON: Well, when you fit
13	chronic exposure models, you like to fit intake
14	regimes, as we call them, with that have
15	similar excretion patterns. So the blue dots
16	is Excretion Pattern 1, the red dots are
17	Excretion Pattern 2. Those were fit as
18	separate chronic intake exposure scenarios.
19	So the blue dots were fit all the way
20	through whatever that number is, 12,000 days
21	post start of the site, through 1990 I can't
22	read it from here, 1990 it looks like. So the
23	blue

1	MEMBER RICHARDSON: Yes. So the
2	origin was 1955?
3	DR. NETON: Right.
4	MEMBER RICHARDSON: And you've got, I
5	think, 60 years of data, 20,000
6	DR. NETON: Right.
7	MEMBER RICHARDSON: days or
8	something like that.
9	DR. NETON: Right.
10	MEMBER RICHARDSON: So you're going
11	up. And so that the red, the transition from
12	blue to red is not related to the two rows of
13	your summary table, 1955 to 1990 and 1991 to
14	2007? That's something different?
15	DR. NETON: No. Well, the
16	transition in 1990 was because you can see there
17	was a there's a distinct, an abrupt decrease
18	in the urinary output in 1990. So to fit a
19	continuous chronic exposure model through all
20	of those dots just doesn't seem reasonable.
21	MEMBER RICHARDSON: So, okay. So
22	the again, the transition from blue dots to
23	red dots is because of a change in monitoring

1	practice. It's not because I thought you
2	were describing it as an assumption about a
3	particular exposure scenario.
4	DR. NETON: Well, it's not a change in
5	monitoring. It's a change in the qualitative
6	look of the plots. I mean, you can see there
7	that there's a fairly abrupt change in the
8	output of the urine. And so
9	MEMBER RICHARDSON: Well, I mean, you
10	might, or one might look at it and see that
11	there's two dots, perhaps, that look like
12	outliers and everything else looks like
13	DR. NETON: Wait, wait. I'm now,
14	I
15	MEMBER RICHARDSON: Or two or three,
16	yes, but I mean, they're but, you know, you
17	see that in a lot of
18	DR. NETON: Yes.
19	MEMBER RICHARDSON: quirky things
20	in
21	DR. NETON: But I'm saying, if you
22	look at the red dots themselves, they are about
23	an order of magnitude lower or about a factor

1	of five, I can't tell from here, than the dot,
2	the blue dots. So something happened there.
3	Something is inherently different about the
4	urinary excretion pattern in that second
5	period.
6	This was a qualitative judgment here.
7	This is not quantitative.
8	MEMBER RICHARDSON: No, I know. I'm
9	just trying I was just trying to understand
10	the interpretation.
11	DR. NETON: Right.
12	MEMBER RICHARDSON: That was a
13	post-hoc color coding. And then you assume
14	that there's two different chronic intake
15	patterns
16	DR. NETON: Correct.
17	MEMBER RICHARDSON: in, among, on
18	average, among the workers at the site.
19	DR. NETON: Right.
20	MEMBER RICHARDSON: Commencing
21	one commencing in 1955 and the other commencing
22	in 19
23	DR. NETON: 91.

1	MEMBER RICHARDSON: 91.
2	DR. NETON: Correct.
3	MEMBER RICHARDSON: Okay. How does
4	that happen? I guess, you know, I think about
5	an exposure pattern as happening for a worker
6	but not for the median population.
7	DR. NETON: Yes, well this raises
8	some issues with the model. Now remember, the
9	50th percentile is not the same person in every
10	particular year.
11	MEMBER RICHARDSON: Right.
12	DR. NETON: So in some ways, I think
13	it's fairly it's somewhat favorable, in a
14	way, to pick the 50th percentile for every year.
15	It's probably not the same person. You're
16	picking the median value for every particular
17	year.
18	Yes, it's this is the way we've been
19	doing it. I'm not saying it's perfect. But
20	again, you are applying this to unmonitored
21	workers, not this is the experience of the
22	monitored population. Now you're trying to
	II

apply this to people who had no monitoring data

at all, and what is their exposure experience. 1 And if you think of the excretion dots 2 3 as sort of representative of t.he air concentrations in the plant, I think you can see 4 5 that -- in this case, it's a little different because I do believe that the red dots are lower 6 because of a change in the technology. 7 They had a lower -- ability to measure 8 9 lower levels of plutonium, therefore you're 10 seeing lower values. It's very possible that some of these chronic exposure models, the 50th 11 12 percentile is actually equal to the MDA or the detection limit. 13 So some things, I 14 MEMBER RICHARDSON: 15 mean, sometimes it's easier to see changes in monitoring by following an individual. 16 17 you would have workers who maybe were -- you 18 would see the transition easier an 19 individual basis. 20 DR. NETON: That's -- but then, Yes. 21 over this long period of time I'm not sure we

know,

you're

You

thousands and thousands of samples here.

can

do

that.

22

23

talking

1	find that individual thread that you can and
2	then you have one individual for the whole site.
3	Yes, I'm not sure that's possible.
4	MEMBER RICHARDSON: Okay.
5	MEMBER ZIEMER: Well, just a couple
6	of comments. One, there is some subjectivity
7	in looking at plots and saying these are two
8	different ones.
9	DR. NETON: Right.
10	MEMBER ZIEMER: That's one comment.
11	Number two, I think this could reflect either
12	a change in work practices or change in
13	detection abilities. There's several
14	possibilities.
15	DR. NETON: Correct.
16	MEMBER ZIEMER: But
17	DR. NETON: Yes.
18	MEMBER RICHARDSON: I've seen, at
19	Savannah River, changes in recording just
20	because recording practices of or, you know,
21	data issues as well. I mean, so you could see
22	the median dropped to zero because
23	DR. NETON: Right.

1	MEMBER RICHARDSON: zeros are
2	recorded for detection limits.
3	DR. NETON: Well, yes. And that's
4	another issue.
5	MEMBER ZIEMER: There=s some
6	statistical analysis issues that we'll be
7	getting into.
8	DR. NETON: Yes, and that's another
9	issue.
LO	MEMBER ZIEMER: I think it's
L1	important. You can't just take one individual
L2	and talk about the work practice change for
L3	DR. NETON: Right.
L4	MEMBER ZIEMER: I think you're taking
L5	a large amount of data in these things, and
L6	looking at an overall effect.
L7	DR. NETON: Yes.
L8	CHAIRMAN MELIUS: But doesn't that
L9	sort of that sort of begs the question of
20	well, should there be stratification within
21	that large population
22	DR. NETON: Well, and that's
23	CHAIRMAN MELIUS: which I think is

what, sort of what David was getting at, maybe 1 not on an individual --2 3 DR. NETON: Yes. CHAIRMAN MELIUS: -- but that's one 4 5 way you'd want to, you know, sort of examine But by group, there could be, and again, 6 that. 7 you got to sort of define what the plot is plus what you see in, you know, as the mix of workers 8 9 or what -- I mean, there's lots of possibility. 10 And so I don't think you can look at 11 any of this without sort of a pretty thorough 12 knowledge of the site and --13 DR. NETON: Right. 14 CHAIRMAN MELIUS: -- you know. 15 DR. NETON: I'm not trying to imply 16 that this is the right way to go, or one size fits all here. 17 I was just trying to say, if 18 this were a stratified subset, and we knew, for 19 instance, that these were maintenance workers, 20 and they were stratified out, this is how we 21 would go about trying to establish what their 22 unmonitored colleagues' exposure experience

I don't want to belabor this too much.

1	MEMBER KOTELCHUCK: Dave Kotelchuck.
2	Let me ask you, this these data points are,
3	of course, coming out from the IMBA program,
4	right? These are
5	DR. NETON: No. These are actually
6	excretion data points that we received from the
7	DOE, or AWE in some cases.
8	MEMBER KOTELCHUCK: Oh okay. So
9	these are the excretion points?
10	DR. NETON: These are urinary
11	excretion values we have in a database.
12	MEMBER KOTELCHUCK: Okay.
13	DR. NETON: Of the actual exposed
14	workers, sorry.
15	MEMBER KOTELCHUCK: Suppose but
16	suppose you were to follow, not one worker as
17	a representative, but
18	DR. NETON: All worker?
19	MEMBER KOTELCHUCK: Well, it could be
20	a but just a few dozen, you should be able
21	to, if you will, visually see that there is
22	really a transition going on. There should be
23	some kind of tailing off.

1	DR. NETON: It may be, but those would
2	be more demonstrable in the higher exposure
3	levels. And we're trying to get the 50th
4	percentile established here, and those tend to
5	be down into the weeds.
6	They'll monitor 300, 400 workers in a
7	year, and you'll see that the 50th percentile
8	worker, who was monitored, is already bouncing
9	around the detection limit.
10	Yes, there's going to be workers up in
11	the 95th percentile that were more heavily
12	exposed, and you could do individual dose
13	reconstructions, but in this particular case
14	are going to be dose reconstructions using
15	missed dose, for the most part.
16	We've been down this path before about
17	
18	MEMBER KOTELCHUCK: Yes.
19	DR. NETON: can we really use
20	individual workers, and I'm pretty convinced,
21	at least, that it's it would be almost
22	impossible.
23	MEMBER KOTELCHUCK: Individual

1 workers OPOS.

DR. NETON: Well to take OPOS data as an individual worker and reconstruct everybody's dose, individually, would be really, really difficult.

MEMBER KOTELCHUCK: Okay.

DR. NETON: Oftentimes you only have one sample a year on these people, and now you're saying well, I can do more with that than I really can.

You know, there's some substance here. Chronic exposure, the annual values that were taken can be used to bound those chronic exposure scenarios. I mean, you know, those are inherent assumptions but I think that we've sort of gotten through that in the past, and that part, I think, is okay.

What I'm going to talk about next is really, you know, how we determine which data points are used for which sets of workers. That's sort of the thrust of the talk. Maybe I complicated things a little more here than I expected to. But I just wanted people to be

1	aware of, this is what we're talking about, how
2	so. Okay. Any other questions? Henry?
3	MEMBER ANDERSON: Are you using the
4	median value or are you using the mean value?
5	DR. NETON: Median.
6	MEMBER ANDERSON: Median?
7	DR. NETON: Median value.
8	MEMBER ANDERSON: Got it. Because
9	that would help you adjust for the change of the
10	limit of detection. Because although I
11	mean, what I was going to say is if you have a
12	change in the limit of detection, you could
13	apply the same limit of detection from the
14	earlier years to the later years.
15	You'd lose some data, but you'd see,
16	does that change? Is this level that's going
17	down that, starting in 1991, is that an artifact
18	of detection? But if you're using medians
19	DR. NETON: Yes. Well
20	MEMBER ANDERSON: It as long as
21	that median is above the limit of detection, I
22	mean, if in a given
23	DR. NETON: If it is, yes. Yes.

1	MEMBER ANDERSON: Is it? Is it
2	typically, in the earlier years, also
3	DR. NETON: In the very, very early
4	years, it's above the detection limit. As you
5	get more contemporary, maybe 1970s, 80s, it's
6	almost very often about the detection limit,
7	sometimes even below the detection limit.
8	MEMBER ANDERSON: So you artificially
9	assign the limit of detection to the value? Or
10	a square root of
11	DR. NETON: Yes. We have techniques
12	for accommodating for what we call a missed
13	dose, right.
14	MEMBER ANDERSON: Because that
15	DR. NETON: But remember, each of
16	these values is going to have a geometric
17	standard deviation of 3 associated with it
18	anyways, so.
19	CHAIRMAN MELIUS: Okay. Any Board
20	Members on the phone have questions, before we
21	move on?
22	MEMBER FIELD: No, this is Bill. I
23	thought this was a very helpful presentation.

1	Appreciate it.
2	DR. NETON: Thanks.
3	CHAIRMAN MELIUS: Okay.
4	DR. NETON: Okay.
5	CHAIRMAN MELIUS: Next.
6	DR. NETON: All right. That being
7	said, now we'll switch gears a little bit and
8	talk about how we ended up with this coworker
9	model draft again, I emphasize draft
10	implementation guide.
11	It doesn't say even implementation
12	guide yet, but the idea is that this will end
13	up becoming NIOSH Implementation Guide, I
14	think, 006. You know, we have one for covered
15	exposure, we have one for surrogate data. So
16	this will be the latest in our collection.
17	So I thought the best way, maybe, to
18	go over
19	MR. HINNEFELD: Excuse me just a
20	minute, Jim. I just wanted to make sure people
21	who are on the phone, we have just left Live
22	Meeting on the presentations. So if you're

following on Live Meeting on the phone, you

won't see what's on the screen here in the room, 1 but it is on the website. 2 called Draft. Criteria for 3 Tt's Evaluation of Coworker Data. I think it's Rev 4 5 3. I think there's a rev -- it's coworker data and Rev 3 is in the title of the file. 6 And that's what's on the screen here. 7 DR. NETON: I'm reasonably certain 8 9 that it's on the website. At least I requested 10 that it be put there, so. So yes, and the idea 11 is, here, we're just going to scroll through, 12 because I find it more comfortable to talk from 13 something rather than speak in generalities. So the idea was, we're up to Rev 3. 14 15 started off saying okay, we didn't have any such 16 quidance in the past. I mean, we've built a lot of coworker models, and our approach, from the 17 18 very beginning, has been, let's just take all 19 the data, rank it and apply it, and not spend 20 a lot of time thinking about where these little 21 subsets may have been. 22 I mean, we've done some of that. We 23 talked earlier about Report 52, or TIB-52.

we went back to the drawing board, said what really do we need to think about when we're doing these coworker models?

So there's four sections to this

So there's four sections to this documents. I think it's up to about eight pages now. The first section is the introduction, which sort of gives us the basis of why we have coworker models in the first place.

The second section talks about, if you have a set of data, you need to look at it for data adequacy and completeness, and also the type of program that they were trying to implement with that data. So that's a data adequacy type thing.

The third section talks about if you -- once you decide that you can really use the data to establish coworkers, how do you analyze it. And that kind of gets into this 50th percentile, 84th percentile situation.

Then the fourth set, which is still sort of a little bit soft in my mind, is how you actually do a statistical analysis for

stratification. And it'll become obvious as I talk why that's sort of still out there.

So if we could scroll through just the first section, and maybe stop at that quote there. It talks about individuals. The dose reconstruction regulation is directly out of that.

It says, "If individual monitoring data are not available or adequate, dose reconstructions may use monitoring results for groups of workers with" -- and this is where it gets tricky, for "comparable activities and relationships to the radiation environment."

That's right out of the regulation, so that's what we're obligated to do. Now oftentimes, you know, the level of granularity, though, how much you can get in the weeds about who that monitoring data was taken from and that sort of thing is not always as good as you'd like it to be.

And so we have, in the past, generated coworker models that pretty much are one size fits all, you know, a full distribution of data.

I'm going to go into some descriptions here, though, about what we should be considering up front, before we go, before we make that leap of faith. Okay.

So this next section, criteria for the evaluation adequacy and completeness -- go down a little further, Stu -- yes, just this section here. The data adequacy really speaks to the technical ability of the monitoring methods that were employed.

I mean, we have a lot of data that are taken all the way back from the 1940s to the current time. Clearly the technology has changed. And so you have to take the data set that you have in hand and establish, is this -- can this data really, reliably -- can it be reliably used to determine what the person's value was, excretion-wise or on their dosimeter? Is the technology there?

I always use the example of measurement of neutrons in the early days using film. The film couldn't see neutrons, energies probably below say 500 keV.

So you've got to be aware of that and say well, these data -- we have these data but we need to consider what the heck was going on with the technology, and either adjust it or say it can't be used.

So this sets the stage for what type of data could be used in coworker models. We talk about using a bioassay, which we often, very often use. And bioassay, in our mind, also includes in vivo analyses, although we don't do that particularly often. But it can be a valid technique for a coworker model. We've done this.

And I also included as a footnote in there, it talks about you can use breathing zone air samples. We've never done that yet, I don't think, but it certainly would be a viable option if you could show the breathing zone data were pretty good, I mean, they were really lapel air sampling, that sort of thing.

And then of course, you need to talk about external dosimetry type measurements, that measure beta, gamma, neutron, that sort of

1	thing.
2	So it outlines here sort of the
3	criteria to look at, not all inclusive of what
4	should be evaluated when you have in vitro
5	measurements, in vivo measurements, that sort
6	of thing. I won't read all the criteria, but
7	they're in there. And then the last section
8	talks about external.
9	So here we're just trying to vet the
10	quality of the data. Is the data useful? The
11	next step goes into the completeness of the
12	data.
13	MEMBER LEMEN: Can I ask
14	CHAIRMAN MELIUS: Yes, sure.
15	DR. NETON: Yes, maybe I should stop
16	after each section.
17	MEMBER LEMEN: Is there
18	MEMBER MUNN: Microphone.
19	MR. KATZ: Use the microphone please.
20	MEMBER LEMEN: Sorry. Is there good
21	compatibility between the time frames the
22	samples are taken? In other words, does the

methodology of sampling techniques change from

1	one decade to the next decade?
2	DR. NETON: It can, yes.
3	Definitely.
4	MEMBER LEMEN: And how do you adjust
5	for that?
6	DR. NETON: Yes, well that needs to be
7	taken into consideration, the temporal nature
8	of the quality of the data. And I think it's
9	somewhere in here. Dr. Melius raised that same
10	issue the last time. And you're right, you
11	can't take a 1950s technology or,
12	conversely, take a 1980s technology and say
13	that applies all the way back.
14	MEMBER LEMEN: Well, what I wanted to
15	see was how you adjusted to that, and what you
16	did, and if that's later on, why we
17	DR. NETON: Well, actually, this is a
18	
19	CHAIRMAN MELIUS: I was going to say,
20	it's always going to be applied case by case or
21	site by site. So I think it we're just
22	trying to get the general areas of
23	consideration that need to be taken into

account when developing the model. 1 But it's going to be a -- and certainly 2 there's lots of examples that I can think of 3 where we've taken -- we've either had an SEC 4 5 because of a problem with an older monitoring method, from a lot of the early ones. 6 There weren't monitoring methods available or they 7 were very crude relative to what would be needed 8 for dose reconstruction. 9 So I think we have a fairly -- a fair 10 amount of experience with that. You know, if 11 12 I remember, specifically with Fernald, but I 13 think recently the partial -- the SEC was based on a problem with the --14 DR. NETON: I think that was the in 15 vivo data at Fernald. 16 17 CHAIRMAN MELIUS: Yes, right. In 18 vivo data, right. 19 So yes, this document, I DR. NETON: 20 think, it tends to talk about what you need to 21 look at, and why, not necessarily exactly how 22 to evaluate. It's a sort of a general 23 quidance.

There is a section here at the end of that, that does talk about looking at the detection limit of the system. For example, oftentimes with thorium measurements you — thorium urinalysis is a very poor measure of how much you expose. It has a very poor detection limit. And so you could inhale a lot of thorium and not be excreting much in your urine.

So in those cases, even if you have a lot of data, you might say well, does my coworker model provide me plausible values? I mean, you could say well it's less than X, which is a very high number, but is that really a reasonably accurate value?

So you need to take that into consideration as well. So that's the sort of -- it's sort of scripted out here. The in vivo section does talk about using the progeny, the -- not measuring the radionuclide directly because sometimes, like thorium doesn't have any usable photons, so you start using some of the daughter progenies.

And you have to think about the

implications of that, and are 1 equilibrium or are they not, and if they aren't, 2 how do you adjust it, that sort of thing. 3 4 there's a lot of things that need to 5 considered. And I wouldn't want to begin to cover 6 7 each and every one of those in this document. So this is sort of a general road map. Okay. 8 9 Any more questions on data adequacy? 10 Okay, data completeness, this is where you need to look at to see if the data 11 12 actually measured -- had a reasonable handle on 13 the measure -- on the exposed population. they monitor enough people, and enough people 14 15 by job category, for example, of the people that 16 were exposed? 17 I think I called this a gap analysis, 18 looking for, you know, first temporally by 19 years. Do you have data for every year? 20 not, there are years missing, you need to figure 21 out why they're missing. 22 Maybe there's good reason for it, the

plant was shut down. If not, maybe rethink

about what you can do in those years. But then you need to go back and look and see, are the work categories that were represented in those facilities adequately monitored.

I have an example here that came out at one of the -- you have that table, I think, is down there. Yes. This is a good example I just threw in. I thought it was pretty appropriate. SC&A had mentioned this and I thought, yes, it makes some sense.

This is where an SEC was added at the Nevada Test Site. And we, you know, originally our contention was, well we have a lot of monitoring data. We have 290 samples here, of a lot of workers.

But when you really do an analysis of the job categories that were monitored, in this particular case more than 2/3 of the samples were taken on the rad safety staff, while the other workers were not very well represented.

Now, if one could argue, and make the case that the rad safety staff are the highest exposed, okay. But if that's not the case, now

you got a problem.

And so that's all this section is trying to say, is you need to look at the data, and do so qualitatively and quantitatively and establish that, you know, you really can bound these different categories of exposed workers.

Okay. So that's pretty much that in a nutshell. I think the table does a good job, kind of driving that home. Of course, you know, there's language in there about looking at the magnitudes of the exposures.

Very small exposures, you might not see a lot of monitored workers, or if there were, you know, special considerations you need to consider. But in general, I think this is the way it should go. Okay Stu, you can keep moving down.

Yes, I kind of went over this, so you can read this again, but I just want to -- okay, the next section talks about, now that -- if we believe that we have enough monitored workers, and who were monitored in the different job categories, we also need to look and decide --

1	oh, sorry.
2	CHAIRMAN MELIUS: Can we just stop
3	for
4	DR. NETON: Oh I'm sorry, yes.
5	CHAIRMAN MELIUS: comments or
6	questions? Unfortunately some of us have been
7	so close to this, and sort of, so I could
8	recognize where you had updated it that
9	DR. NETON: Yes.
10	CHAIRMAN MELIUS: so we could keep
11	review. At least in my mind, this is sort of
12	an important section that we need to be able to,
13	you know, as part of our evaluation, sort of
14	need to hear about when we're looking at a
15	coworker model.
16	Because it by the time a coworker
17	model comes out this is often hidden. I mean,
18	this is hidden behind the model. And in terms
19	of the judgment that goes into it, in I mean,
20	and I think many of these factors have been
21	evaluated and looked at.
22	I don't think, again, we've always
23	looked at them ourselves during the process,

and there have been problems. We've seen a 1 number of SECs where, when we've sort of poked 2 3 behind the model we've raised questions. But it's -- you know, I think we need 4 5 to urge the other Board Members to sort of look at this and sort of think about what other 6 7 might have if questions you you evaluating, you know, the presentation of a 8 coworker model about the data. 9 10 I've got a few things I want to add here, and I think -- everybody to look at 11 12 because again, I think it is an important part 13 of this. MEMBER RICHARDSON: Can I ask one 14 15 question? That -- just a -- in that, in the table, the -- what's just an illustration of a 16 17 problem, if you could scroll back up to that 18 Table 1. 19 DR. NETON: Okay. 20 MEMBER RICHARDSON: Where just in the 21 last period there are -- there's the bulk of the 22 bioassay monitoring for people other than rad 23 safety staff. It's the bottom right hand

corner where all of a sudden 73 --1 DR. NETON: Security? 2 3 MEMBER RICHARDSON: -- out of 74, is that all like exit bioassay? 4 Is that -- or was it -- what drove that to happen, do you know? 5 DR. Ι honestly don't 6 NETON: 7 remember. I don't think it was exit bioassay, though. I think security just were added in 8 9 the '81 to '92 period. I don't recall why, to 10 be honest with you. 11 MEMBER RICHARDSON: Okav. 12 CHAIRMAN MELIUS: Yes. If you go 13 back to the NTS report, there's a -- at least the SC&A review is a fairly extensive analysis 14 of this. Because this took us a while to sort 15 of get on top of and sort of understand, and 16 17 there was a lot of back and forth as to, about 18 the data set and was it appropriate. 19 I recall, a fairly good But as 20 analysis that those, that the rad safety 21 workers really didn't, really were 22 representative of the other people doing the

And it was more than just sort of a

qualitative assessment, it was also looking at the data from a --

DR. NETON: Yes, there were other issues with the Nevada Test Site. They tended to be episodic samples versus routine, and we had a -- you know, when you develop a chronic exposure coworker model and these are episodic, incident-driven samples, how does that chronic model really fit the picture?

And we're going to talk about that. That's a good segue into this next section. You know, so now that you have the technical adequacy of the data, and you feel like you've got a fairly complete picture of who was monitored and why, and it seems okay, you still need to consider the type of monitoring programs that were applied to these workers.

We outline the three types that we can think of, which are the routine representative sampling of the workers, routine measurements of the workers with the highest exposure, or incident samples.

Those are the three major ways that

monitoring programs sort of come about, and you need to look at each of those populations and say, were they all -- if you want to combine it into one coworker model, first of all, were they all routine samples, yes or no?

If, for example -- and this comes up very frequently, and right now we're discussing this at the Savannah River Site, where building trades workers only monitored on an incident basis whereas everyone else in the plant who were doing routine ops were monitored on a routine basis.

Well, it's hard to convince myself right now that you can actually combine those two into one general coworker model. That's what we're saying. So this section goes into that in some detail about how one needs to look at that.

A good way to, of course, reevaluate if there's a routine program is to go look at the radiological control program documentation. It should spell out who was monitored under what frequency and that sort of

thing.

But then it's -- you have to do a little more than that. You also have to go and make sure that they actually did that. Some sites had very -- well, documents with very good intentions, because of funding or whatever, didn't actually end up following up and collecting the samples that they thought they were. So you need to get some indication that that occurred.

So that's all kind of spelled out here. There's a special category that we put in there, which I think is a special category of routine, although maybe not really, which is people that worked on very short duration campaigns or projects, where it was short enough where they would just take maybe a beginning and an end sample.

And those are probably routine in a way because the project was short enough you could use those data for that specific group of workers to reconstruct their exposures. So that's discussed in here to some extent.

And the last piece of this, applicability of the monitoring data, I think Stu, if you scroll down to the end, maybe I'm wrong. What else is in here? No, that's not -- I think I covered all that.

I meant to say, in the last section under completeness, you also needed to go and look -- we need to look at if you're using data sets provided by the site or epi studies or whatever, you need to kind of make sure that you have -- the data set has all the monitoring data in it.

I mean, if for some reason there are gaps, the computer program didn't collect all the data or it got lost, that needs to be evaluated. And that can be done by going back and looking at some of the original records, if you have them, or some reports that said, in 1955 we took 10,000 samples, or by month, it was this.

You go in your data base and get yourself a good feeling that you have those, that many samples in there, or the fact that

incident samples were always collected separately and aren't in this data base. Well, you need to know that. So that's something that needs to be considered.

Okay, any questions on that? I think we're okay. The third section here is the analysis of the data, and this is basically what we just talked about, how one can generate these distributions using the 50th and 84th percentile to analyze the data. And we would use the 95th percentile of the data.

If you had coworker models, all routine, and you had construction workers in there that were routinely monitored, they would be given the 95th percentile of the exposure because they are presumably a much higher, more highly exposed than say a person who was intermittently present in the plant, during things like walk-arounds, security guards, clerks, that sort of thing. They would get the 50th percentile.

So that's -- this just speaks to that type of a scenario, how you -- what do you do

with the data once it's all good to go, so to 1 speak. 2 It talks about this OPOS scenario, 3 where one person, one statistic for monitoring 4 5 interval, and it refers, actually to Report 53, which is out there, that discusses this OPOS, 6 has been updated to include this one person, one 7 sample, and the fact that it's a time-weighted 8 9 average that goes backwards in time, averaging, rather than forward in time, and a little bit 10 about how to deal with negative values. 11 12 We don't -- we won't use negative 13 values in doing these time-weighted averages. We've come to that conclusion. So that's what 14 15 this section deals with. It's fairly 16 straightforward. Okay, the final section -- oh well, 17 18 just one -- the time interval of the modeled 19 data, we talked about this at some length during 20 the Working Group meetings. 21 Oftentimes we have an annual sample on 22 workers, and we'll be using that. But in some

cases, when the data are sparse, can you lump

some data together? And we feel that you need 1 at least 30 samples in a monitored interval, 2 that one monitored period for good statistical 3 considerations. 4 And if you're going to do that, you may 5 be able to group years together, several years, 6 as long as you can demonstrate that the work 7 practices and processes remain the same over 8 9 that time period. 10 This says here at the end that those intervals should not exceed a three year 11 12 period, unless there's stringent justification 13 for doing so, and that's where it remains. I originally had five years, but I 14 15 went and back and checked our original Report 16 53 and it did say three years. So that's -- at 17 least we're now consistent with our 18 internal documentation. I can't remember all 19 these numbers. 20 So any questions on that, the Okay. 21 analysis section? 22 So now we get to Section 4, which I said is still a little soft. 23 I'm not -- we're

not done with this yet, at least in my opinion. You could tell by what I just described that there's a pretty good process of what needs to be considered on how to stratify the data.

And if any of those conditions are met, I think you just need to stratify. I mean, if you can show that trades workers were incident-driven samples and bioassay, and they were different than the routine process workers, you know, I think one needs to stratify no matter what statistic.

You know, you don't need a statistical analysis to do that, because you've decided, a priori, that these are different populations to begin with. So that leaves the question in my mind open about when one would actually go about doing statistical tests on these data sets.

And we left it at the Working Group discussion level that we would -- we're going to go back and do some example analyses to see. I think it's best accomplished looking at some examples.

I -- right, in my mind right now I'm

1	not really sure how that's going to play out.
2	So right now, this is written very much in line
3	with what Report 53 says, which is this Monte
4	Carlo permutation test or the Peto-Prentice
5	test, it is a statistical test that can be used.
6	I'm 100 percent certain when this
7	would actually, in fact, be appropriate. And
8	so this section is sort of on hold right now
9	until I we get some examples together and can
10	talk a little more concretely about it.
11	I think that concludes my quick
12	perusal through the document.
13	CHAIRMAN MELIUS: Are there comments
14	or questions? Brad, Gen, Loretta?
15	MEMBER CLAWSON: Jim, I understand
16	where you're getting at to this, but to me, this
17	all comes back to the data that we have, and how
18	good it is. This is correct, right?
19	DR. NETON: Yes.
20	MEMBER CLAWSON: I'm looking at that
21	30 percent there, and or 30 samples. And
22	when you're looking at a workforce of maybe a
23	thousand people there, that's I'm not the

1	sharpest tool in the shed here, but it looks
2	like there's some pretty big gaps in there.
3	DR. NETON: Well, and I think there's
4	some caveats in there, that says 30 samples is
5	a minimum but you need to look at the population
6	of monitored
7	MEMBER CLAWSON: Population, okay.
8	DR. NETON: workers and that sort
9	of stuff. Yes, it wasn't the intent that no
10	matter what you could use 30 samples. I agree
11	with you. But again, 30 samples if you had
12	50 people working on a glovebox operation for
13	a year, and you had 30 samples, and they were
14	the highest exposed workers on the glovebox
15	line and you could establish that somehow,
16	maybe that's okay.
17	MEMBER CLAWSON: Okay.
18	DR. NETON: So that's all we're
19	trying to convey there.
20	CHAIRMAN MELIUS: Again, just I
21	think I said this a little bit before, but I
22	think what we've been trying to do is what do
23	we need to look at up front before we get into

stratification? What information do we need to have, and have evaluated, probably more qualitatively than quantitatively?

But -- and with, you know, a fair amount of judgment and a fair amount of information about the site. It's always going to be specific to the site.

But then I think if the -- when I was reviewing one of the earlier back-and-forth reviews from SC&A and, I think, NIOSH's response to it and so forth, I mean, I could come up -- I think they were both right and they were both wrong in the sense that you could come up with scenarios or situations where, you know, whether you could stratify and how you would handle that stratification would be quite different depending on the circumstances at a particular site.

And so I think if we can sort of systematize and get a better set of guidelines on what we need to look at, I think it becomes then easier to decide, is 30 the right number or, you know, and some of these other sort of

more statistical issues. 1 At least we have sort of a --2 3 MEMBER ROESSLER: You need а threshold. 4 5 CHAIRMAN MELIUS: You have а threshold but you also have sort of a factual 6 background that you understand what's going on 7 at that site. So that's what we're trying to 8 9 do, and it's probably why we've more heavily 10 weighted the beginning of this thing, report, and got the implementation guidelines. 11 And I think the other will follow from 12 And I actually think, as the results of 13 our Work Group discussions, some of those 14 15 issues that we were going back and forth on, on OPOS and so forth have become less troublesome, 16 17 so to speak, or less of an issue. 18 So I think it's the, sort of the right 19 way to go. And again, many of these factors 20 probably considered in developing were 21 coworker models, they just weren't 22 explicit in terms of how it was presented to us.

Gen?

1	MEMBER ROESSLER: We have a Board
2	with a wide range of perspectives and
3	background knowledge on this subject, and I
4	think it's really important at this point to
5	help move this forward, that Board Members do
6	submit written comments to Jim in the next
7	couple of weeks. I think that's the most
8	important step in moving this forward.
9	CHAIRMAN MELIUS: Loretta?
10	MEMBER VALERIO: I guess my question
11	is, the coworker models that we're looking at
12	right now are based on chronic exposures,
13	that's correct?
14	DR. NETON: Yes.
15	MEMBER VALERIO: I would assume all
16	of these sites had projects that were short
17	duration, which you did address. At any point
18	do you anticipate that a coworker model for
19	acute exposures would be established?
20	DR. NETON: I didn't cover it in my
21	discussion, but the document does allow for it.
22	Incident-driven coworker models may be
23	appropriate, particularly in the more current

era, when you have very good workplace controls that are -- that can be demonstrated, where there are, you know, continuous air monitors, people frisked in and out of the area, that sort of thing.

And so if you're comfortable that you can believe that there were no upset conditions that occurred that weren't caught somehow, and if that's true, then I think you could use an incident-type model.

But you -- and I think it says in there, you have to be very careful about that. You know, your documentation has to be almost impeccable to be able to do that, but I think it's a -- could be allowed for, particularly post 1990, where, you know, you have, almost everybody is supposed to be on a monitoring program if they had a 100 millirem potential exposure for internal, those sort of things.

And, you know, but you always have to allow for some gaps and technology shortfalls and stuff, so you may be able to do an incident-based model in that scenario.

1	CHAIRMAN MELIUS: Other comments or
2	questions? Any Board Members on the telephone
3	have comments or questions at this point?
4	MEMBER FIELD: This is Bill. I don't
5	have any questions. A lot of this is, has
6	stayed relatively the same over the past year
7	or so, hasn't it?
8	DR. NETON: What was that, Bill? I'm
9	sorry.
10	MEMBER FIELD: I said, most of this is
11	relatively constant. There's not a whole lot
12	new here, is there?
13	DR. NETON: There's a whole lot new in
14	the sense that it's in writing now.
15	MEMBER FIELD: Okay. But this is the
16	same concept that you had discussed previously?
17	DR. NETON: Yes, yes. They've
18	crystallized a little more, and there's some
19	more caveats in there, and maybe some scenarios
20	that I wake up at night and think about
21	MEMBER FIELD: Yes.
22	DR. NETON: and put in there, but
23	in general, it's the same.

But it's nice having 1 MEMBER FIELD: 2 it down on paper that we can provide comments. Thanks. 3 DR. NETON: Yes, okay. 4 CHAIRMAN MELIUS: Yes. I think the 5 -- I mean, the example of, evolution has been 6 on some of the stratification issues between 7 incidence-based and routine monitoring and, 8 9 you know, when is that appropriate, when do they 10 need to be separated and so forth. 11 So, but there's been some changes. 12 But it's, again, getting it down in writing, I 13 think, is what's -- you know, what, as Jim said, is what's most important. 14 15 DR. NETON: I think what's changed, most significantly, is if you recall early on, 16 17 we were trying to come to grips with some 18 practical level of significance and difference 19 between models. 20 We were kind of approaching it from the backwards situation where we were looking 21 at this 100 millirem dose, and then I proposed 22 this model of using the full distribution 23

versus the 95th.

That's all sort of by the wayside right now, because frankly, none of it seemed to work. And I like this approach better where you just identify, do you need to stratify, yay or nay, and then go ahead and do it if the conditions are such that it need be.

Member comments and -- so the, I mean, I think the one thing I'd also mention is that, in terms of the, what do you call it, the rule of 30 or whatever is, it's also, I think we have to remember that when we have a situation where there's very low exposures, residual periods and so forth, I think those you sort of approach differently than you would a situation where you may have very high exposures within a site.

And so that has to be taken into account also. I think what -- as Jim was saying, we were -- we tried -- we started dealing with this on the sort of statistical level, so the problem we got hung up on was just, you know, and some of what we did on sufficient

accuracy, what was helpful for us to understand, but it's just very hard to -- it gets very complicated.

And I'm not sure that the situations

are comparable enough at each site that a very statistical approach is going to be practical.

I think that's sort of what we found. Every site is different enough, has a different set of records.

And I think the other thing we have to remember, take into account, is that even if sort of theoretically we can identify, stratify groups and so forth, we got to be able to place people within those strata. And very often the records just don't exist.

You know, employment records aren't such that you can tell. And, I mean, that's sort of something we need to take into account. But that's often been the problem we've had with a number of the other coworker models in the past.

They're fine, theoretically, and in general the data supports it, but when you go

1	to then apply it to individuals, it just, the
2	information isn't there. And I think we have
3	to think I mean, it doesn't say we can't apply
4	some of these in those situations, but we have
5	to really think carefully how we do that. Yes,
6	Dave?
7	MEMBER RICHARDSON: You know, the
8	rule of 30 is sort of, I think it's the way
9	that the approach is developed, I think it makes
10	sense. And on the other hand you could argue
11	the opposite, that as the data become more
12	finely stratified you may not need as many
13	observations.
14	And it seems like they I guess the
15	extreme would be if you imagined two workers set
16	out to do a task and only one of them was
17	monitored. And that would be sometimes how you
18	would describe coworker settings, that there
19	was
20	DR. NETON: Yes, for example, six
21	workers. Yes.
22	MEMBER RICHARDSON: Right, where
23	it's there was a lot of knowledge that there

was similarity of the task and the work experience and the environment that they were going into, and you would issue one badged. So you wouldn't say you need 30 workers to be badged to, you know, to kind of protect or monitor the 31st.

You might have -- they might work as a team, and then have one observation measured and the other one -- right. But I -- so I mean, it's -- but the problem, as you're saying is, that would be the extreme, where you had a lot of knowledge to place those people into the same time and area and task. You wouldn't need very much information to be sort of confident about understanding their exposure.

And I guess what we're describing is we're using coworker monitoring, not the way that some people do in radiation protection, that would issue a badge to a group of workers who are going to be doing things, but turning the world on its head.

MEMBER ANDERSON: Like we did yesterday, when we had one badge for five slots.

1	But then we loaned our badges
2	CHAIRMAN MELIUS: I okay. I
3	wasn't aware of that.
4	MEMBER ANDERSON: So we have personal
5	experience.
6	DR. NETON: In our part we call that
7	cohort badging.
8	CHAIRMAN MELIUS: Yes. I think the
9	other factor that I have some trouble thinking
10	how I'm not sure how important it always is,
11	but I think we sort of have to take into account
12	is what is the gap we're trying to fill and how
13	much data do we have?
14	If we have really good data for every
15	year but one, and, you know, it's a sort of a
16	production workforce that's, you know, going to
17	be there for a long period of time that we're
18	looking at, I think we're more comfortable with
19	a coworker model and what data that that's based
20	on.
21	If we have, you know, we're missing
22	lots of years on everybody, and a very small
23	percentage of the workforce has been monitored,

and there's a lot of variability, and there's 1 high exposures and so forth, then I think we 2 have to have more stringent criteria in terms 3 of whether we -- how good is the coworker model, 4 because, you know, to do that. 5 And in some ways it's a statistical 6 judgment, but I just don't think we can get 7 there very easily. I think it's more going to 8 9 be judgment, but we need to be able to look at 10 that. Andy? Yes, I was just 11 MEMBER ANDERSON: 12 looking at this and thinking in terms of, do you 13 see us being able to, in a general sense, convert this into a bit of a checklist? 14 15 I mean, when we get your SEC and you 16 say, well, you know, here's how much data there is there, and we believe we can only do this, 17 18 you know, we're denying an SEC because we can use a coworker model, then we have to start to 19 20 try to qualify -- is that useful? 21 Can you convert this, in a general sense, into, you know, a checklist like we've 22

done with some of the -- I mean, you can't, I

1	think
2	DR. NETON: Yes. I would certainly
3	entertain any suggestions to do so.
4	MEMBER ANDERSON: Well, that's I
5	don't know. I was but that's what I was
6	talking to the
7	DR. NETON: But yes, we talked about
8	that earlier, you know, a table or something
9	like that to and that's possible. I mean,
10	it would be a general checklist because again,
11	we're not trying to cover this
12	MEMBER ANDERSON: A descriptive
13	checklist, yes
14	DR. NETON: We're not trying to cover
15	this
16	MEMBER ANDERSON: like the
17	30-number or what are the characteristics that
18	you're hoping to fill here, rather than just
19	saying we're going to and here's the model
20	we're going to use, and we I mean, that's
21	DR. NETON: Yes. It could be fairly
22	easily converted to some sort of but it
23	wouldn't be perfect, because again, it's a

1	MEMBER ANDERSON: No, no. It
2	wouldn't be
3	DR. NETON: qualitative judgment.
4	MEMBER ANDERSON: you have to meet
5	these, but it would be helpful as a descriptive
6	thing when we're looking at these, to say you're
7	going to I was looking at and I'm trying to
8	I don't know if I could do that, but I'm
9	asking you to do it.
10	DR. NETON: I think I it could be
11	done.
12	MEMBER ANDERSON: Yes, okay.
13	MEMBER ZIEMER: Yes, I don't think
14	it's quite a checklist but we these are
15	criteria, and I think we would expect NIOSH and
16	SC&A both to look at data sets, and examine how
17	each of these issues was addressed for a given
18	situation or a given site. So if that's a
19	checklist, it
20	MEMBER ANDERSON: Well, that's what I
21	meant.
22	DR. NETON: Follow yes.
23	MEMBER ZIEMER: We have other

1	criteria that we use, like the surrogate data
2	issue. And it's not quite a checklist, but you
3	have to say, how did you evaluate against these
4	criteria?
5	DR. NETON: That's true.
6	MEMBER ZIEMER: And then we have to
7	examine whether or not we feel that that's met
8	some sort of bar or a test level, you know.
9	DR. NETON: Not unlike what we do for
10	surrogate data. I mean, there are four or five
11	criteria, and we drill down through them and
12	say, okay, this is this met, is this met, is
13	that met, is
14	MEMBER ANDERSON: I think your
15	categories here fit that
16	DR. NETON: They do.
17	MEMBER ANDERSON: process. So I
18	don't think you're missing anything.
19	CHAIRMAN MELIUS: Okay. The
20	heartburn question, or the one that'll keep Jim
21	up at night, and Stu, is, well, we've come up
22	with these criteria, guidelines, whatever we
23	want to call them, what does this say about past

coworker data sets?

DR. NETON: I've already thought about that.

CHAIRMAN MELIUS: Yes. And I think
-- and again, it's -- you know, is it worth -you know, when should we go back? Do we need
to go back and, you know -- can we then -because I suspect, as I'm pretty sure that a lot
of these, you know, criteria or guidelines,
whatever you want to call them, have been taken
into account.

We may not have all -- may have been not documented to us, or the documentation may not be as explicit as we thought, but -- or might want now, but it's, it may still be there, and they may not. But we've changed, you know, the criteria on stratification, so to speak.

That may be more of an issue, but it may not. I don't know. I don't know what situations -- again, because often some of the practical issues about a site, particularly would the employment records support a differentiation between different types of

workers, you know, may, you know, sort of have 1 obviated stratification. 2 So, and they may already be SECs or 3 But I think we can cross that 4 whatever. 5 bridge, and Jim will have a few sleepless 6 nights. 7 DR. NETON: That's one thing that I've been thinking about. You just mentioned 8 9 it, Dr. Melius, is that many of the coworker 10 models that we developed early on, the sites have become SECs for very large portions of 11 12 where the models apply. 13 CHAIRMAN MELIUS: Yes. DR. NETON: And then what does that 14 15 mean in terms of sufficient accuracy, you know, 16 that kind of thing for the non-presumptive 17 cancers that we're reconstructing. I'm 18 wrestling with that right now. 19 CHAIRMAN MELIUS: Yes, yes. 20 MEMBER MUNN: But it's still unlikely 21 that there are any salient criteria that 22 haven't been a part of the conversation. 23 fact that they aren't a part of our written documentation doesn't mean that they haven't been considered and addressed in some way. But -- yes, we still don't know what they are.

CHAIRMAN MELIUS: Yes. Yes, that's what I was saying. I think the, maybe the more relevant question is have they been consistently applied.

MEMBER MUNN: Yes.

CHAIRMAN MELIUS: And I think that's -- I think that's what we found, for example, I think, with surrogate data, was that we -- and even with the SEC evaluations, we -- there was nothing new in either of those documents, in terms of what had been done, but there were, you know, a few sites that, where those that there had been, you know, some pretty significant inconsistencies in our approach.

And some of that's just time. Some of it's, you know, information, what was available at one time and not at another and, you know, over the 102 meetings, we -- the Advisory Board's decision-making has certainly changed or evolved. Evolved, that's a better word for

1	it.
2	MEMBER MUNN: Absolutely.
3	CHAIRMAN MELIUS: Okay. Any other
4	questions, comments? Okay. I saw you sitting
5	on the edge of your seat there. We're in
6	we've been so much trouble recently or
7	something that we have two lawyers here today
8	to keep an eye on us.
9	MS. LIN: So if any Board Member felt
10	compelled to submit a written comment
11	addressing this document or this, the
12	application of it, please coordinate with Ted
13	Katz so we can preserve the Board's
14	deliberative process.
15	CHAIRMAN MELIUS: Well, that was
16	easy. Yes. We've got a few minutes. Any
17	items we can do, or work items we can do quickly?
18	Meeting times?
19	MR. KATZ: Meeting times? We can do
20	that. Yes, sir. So, let me just remind you
21	all of what we have already scheduled. Yes,
22	I'm sorry. Thank you.

So what we already have scheduled is,

1	looking forward, January 6th, we have a
2	teleconference. Then March 25th and 26th we
3	have a meeting, place to be determined today.
4	And the next day, 27th I mean, NIOSH
5	has said there may be a lot on the plate for that
6	meeting, so that actually is that's a
7	Wednesday and Thursday, 25th and 26th of March.
8	I don't know whether, if we need to eat into,
9	halfway into Friday, that might be possible for
10	Board Members. We didn't really check about
11	that at the time, but we'll see.
12	Then the next teleconference is June
13	9th, and then the next Board meeting July 23rd
14	to 24th. So that's what we have
15	MEMBER ANDERSON: What was your first
16	meeting?
17	MR. KATZ: I'm sorry. January 6th is
18	a teleconference, 11 a.m.
19	CHAIRMAN MELIUS: And the second
20	teleconference?
21	MR. KATZ: The second is June 9th.
22	MEMBER ANDERSON: Okay.
23	MEMBER VALERIO: And July is the 23rd

1	and 24th?
2	MR. KATZ: July 24th right, for
3	now, yes, 23rd through 24th, that's a Thursday,
4	Friday, I believe, so that's as far as it would
5	go. So
6	CHAIRMAN MELIUS: And both of those
7	the next two in-person Board meetings we have
8	to decide on a location that
9	MEMBER ANDERSON: Exactly, yes.
10	MR. KATZ: Right. Well, for the one in
11	March, we should decide today.
12	CHAIRMAN MELIUS: Yes, definitely.
13	MR. KATZ: Right. So we talked about
14	Richland is a possibility, for Hanford. I
15	mean, the other two sites that come to mind,
16	well at least one that may be right, I'm not
17	sure. I'm always a little unsure, but I
18	thought last I heard, Rocky Flats might be
19	ready by March. Has that changed?
20	MR. RUTHERFORD: I don't know that
21	it's changed. We have four or five issues that
22	are open, and we have papers out on three of
23	those, with another paper coming out shortly,

probably within a week or two weeks. 1 And the only issue that's outstanding 2 is this data falsification issue, which we're 3 waiting for formal release of documents by the 4 5 FBI. And that's kind of up in the air, that one, you know, because I just don't know, 6 know, that's another agency, and how quickly 7 they're going to get those released, so. 8 9 MR. KATZ: Okay. But then it sounds like there's plenty of -- be plenty of material 10 for a Work Group meeting and --11 12 MR. RUTHERFORD: Oh yes. 13 MR. KATZ: without that, depending on what comes of that, there may be 14 15 good fodder for the Board meeting. 16 And so anyway, we have those two. 17 Kansas City, my sense is that that's -- there's 18 still a lot of work ongoing and that's not going 19 to be ready by then. 20 CHAIRMAN Hanford, MELIUS: my 21 understanding is there's active data that has 22 been gathered, and is in process, and there's 23 potential for Board action.

1	MR. RUTHERFORD: Yes. Yes, in fact
2	there's been interviews, data captured, and
3	there's another data capture actually
4	scheduled right now for early December. And
5	there is an open issue that could drive some
6	action for the March meeting.
7	MR. KATZ: So we have one bid from
8	Brad for Hanford. Any reason anyone, any
9	reason not to go to Richland?
10	We'll need to settle it today, because
11	the process of making arrangements, that's got
12	to get going.
13	CHAIRMAN MELIUS: Well, let's talk,
14	throw Rocky into that. What I'm not sure about
15	is how connected these issues are, in terms of
16	decision-making. The Hanford one, the one I
17	know about, is should be relatively
18	straightforward.
19	MEMBER FIELD: Can your hear all
20	that?
21	CHAIRMAN MELIUS: And it's a single
22	issue, but the Rocky ones, I'm not sure where
23	the data falls, but how that ties in to some of

1	the other issues and what will be
2	MR. RUTHERFORD: Well if you yes.
3	If you look at the years associated with the
4	potential falsification, I mean, it lines up
5	with the years that we're looking at with other
6	issues.
7	CHAIRMAN MELIUS: Oh, okay.
8	MR. RUTHERFORD: So, I mean, they're
9	all roughly in the same time period that we're
10	looking at, you know.
11	CHAIRMAN MELIUS: And I think the
12	other, maybe another consideration is that
13	where we hold the meeting doesn't preclude the
14	Board taking action on the site. I think it's
15	as much, do we need where have we been
16	recently and where do we where would we,
17	might benefit from further input.
18	We haven't been I think we've been
19	to Rocky a lot more recently than Hanford.
20	MR. KATZ: Yes. It's been a few
21	years for Hanford.
22	CHAIRMAN MELIUS: Yes. Since we've
23	visited Wanda there.

1	MR. KATZ: Yes. And Josie, yes.
2	CHAIRMAN MELIUS: And I'm not sure
3	which one is easier to get to at the end of
4	March, or get out of, I guess is the
5	MR. KATZ: Well, Wanda made
6	assurances, previously, that Hanford would be
7	fine, Richland would be fine in March.
8	CHAIRMAN MELIUS: Yes, well Brad
9	claims it never snows in Idaho Falls either.
10	Hanford? Yes, okay. Sounds good.
11	MR. KATZ: Okay. Let's do that,
12	then. Very good. Thanks. That's helpful.
13	So we've settled that. Now, just for
14	scheduling further out, so I have, for the next
15	teleconference, again, the last meeting now
16	scheduled, the latest meeting is July 23rd
17	through 24th.
18	We need a teleconference, and
19	ballpark, you know, timing for that would be
20	September 21st, that week. But it can fall on
21	either side of that week, too. September 21st
22	is about the right timing. So look at that week
23	first. If that week's not good, then we can go

1	before or after, too.
2	MEMBER MUNN: I suggest the previous
3	week, the week of the 14th.
4	MR. KATZ: You're not available the
5	week of the 21st, is that what you're saying,
6	Wanda?
7	MEMBER MUNN: No, I could do it.
8	Just September, but it seems the preceding week
9	might be a little easier.
10	CHAIRMAN MELIUS: Okay. Anybody
11	have problems with either week, I guess is
12	David, do you have issues with
13	MEMBER KOTELCHUCK: No, I was looking
14	
15	CHAIRMAN MELIUS: The other David,
16	but you should
17	MEMBER KOTELCHUCK: Oh, excuse me.
18	CHAIRMAN MELIUS: Yes, but
19	MEMBER KOTELCHUCK: I was looking at
20	my the I was looking at Rosh Hashanah and
21	Yom Kippur for some of us.
22	CHAIRMAN MELIUS: Yes, that's the
23	week of the 14th. I've got it on my calendar.

1	MEMBER KOTELCHUCK: Thirteenth, 14th
2	is Rosh Hashanah, so that Yom Kippur would
3	occur, if somebody will help me
4	MEMBER MUNN: On the 23rd.
5	MEMBER KOTELCHUCK: Pardon?
6	MEMBER MUNN: On the 22nd.
7	CHAIRMAN MELIUS: 22nd, 23rd.
8	MEMBER KOTELCHUCK: Oh good. Okay.
9	MEMBER MUNN: One of the reasons I was
10	suggesting the preceding week.
11	MEMBER KOTELCHUCK: 22nd, 23rd, yes
12	there it is.
13	MR. KATZ: Okay. So recall, this
14	just a teleconference. It's just that 11 a.m.
15	call.
16	MEMBER KOTELCHUCK: Right. So
17	Monday would certainly not Monday the 21st
18	is not
19	CHAIRMAN MELIUS: Yes.
20	MR. KATZ: Yes.
21	MEMBER MUNN: Yes, it is.
22	MR. KATZ: Yes. It's just a call,
23	and it's we can either do it Wednesday,

1	Monday, whatever.
2	MEMBER KOTELCHUCK: That's good.
3	MEMBER MUNN: Yes. Wednesday would
4	be September 16th, the preceding day.
5	MR. KATZ: Well, the week of okay.
6	MEMBER KOTELCHUCK: That would work,
7	the 16th.
8	MR. KATZ: So is that good, the 16th?
9	MEMBER RICHARDSON: The 16th is not
10	good for me.
11	MR. KATZ: No, no, not good. But
12	what about the
13	MEMBER RICHARDSON: I can do anything
14	else, but
15	MR. KATZ: following week, the
16	23rd?
17	MEMBER MUNN: 23rd?
18	MR. KATZ: Is that a
19	CHAIRMAN MELIUS: Of September?
20	MR. KATZ: Yes. Teleconference.
21	Is that good with everybody? Dave?
22	MEMBER KOTELCHUCK: One second.
23	Looks good.

1	MR. KATZ: Looks good, he says.
2	CHAIRMAN MELIUS: Okay.
3	MR. KATZ: Okay. So let's do that.
4	Bill Field, is that okay with you, too?
5	MEMBER FIELD: Yes, that works fine.
6	Thanks.
7	MEMBER KOTELCHUCK: The usual time?
8	MR. KATZ: Yes, 11 a.m. Eastern time.
9	MEMBER KOTELCHUCK: Okay, so that's
10	September 23rd, 11 a.m.
11	MR. KATZ: Okay. And then for the
12	next in-person meeting, the right ballpark is
13	a year from now, November 2nd, the week of the
14	2nd, the 9th, the 16th, that's the ballpark.
15	Get it in before Thanksgiving for sure.
16	MEMBER ANDERSON: Not the first week.
17	MR. KATZ: Not the first week. So
18	the week of the 9th, maybe?
19	CHAIRMAN MELIUS: We've got
20	Veteran's day in the middle of that week,
21	Wednesday.
22	MR. KATZ: Of the 9th?
23	CHAIRMAN MELIUS: Yes.

1	MR. KATZ: It's on a Wednesday?
2	CHAIRMAN MELIUS: It's the 11th, yes.
3	It's always on the 11th.
4	MR. KATZ: Always on okay.
5	CHAIRMAN MELIUS: It's one of the
6	holidays we actually support on the actual day.
7	MR. KATZ: The actual day, right,
8	regardless of what day of the week.
9	CHAIRMAN MELIUS: There's the 4th,
10	and Christmas and a few others, but
11	MR. KATZ: What about the week of the
12	16th?
13	MEMBER ANDERSON: Of November?
14	MR. KATZ: Yes.
15	MEMBER ANDERSON: That would work
16	better for me. Towards the end of that week.
17	CHAIRMAN MELIUS: So 18th, 19th or, I
18	mean
19	MEMBER MUNN: Wednesday and
20	Thursday.
21	CHAIRMAN MELIUS: Wednesday,
22	Thursday or
23	MEMBER MUNN: Eighteenth and 19th,

1	yes, that would be
2	MR. KATZ: Eighteen, 19 are we
3	saying? Okay. And Bill, on the phone?
4	November 18th
5	MEMBER FIELD: That sounds good.
6	MR. KATZ: Okay, super.
7	MEMBER KOTELCHUCK: Now this is the
8	are we talking about
9	MR. KATZ: This is an in-person
10	meeting, November 18th and 19th of next year.
11	Okay. Okay, that takes care of scheduling.
12	CHAIRMAN MELIUS: It's also the
13	September, you know, may be, I mean not to
14	predict anything politically, or not to let
15	politics intrude on our efforts. Okay.
16	MEMBER KOTELCHUCK: This meeting's
17	going to be done for
18	CHAIRMAN MELIUS: Absolutely.
19	Absolutely. It's that's I mean, will
20	last for a whole fiscal year. I mean, I'll be
21	meeting through the following September before
22	we get out of there.
23	MR. KATZ: Okay. That's we've run

1	out of time.
2	CHAIRMAN MELIUS: When I was in
3	NIOSH, I had somebody who was trying to get out
4	to the crab processing places out in Kodiak, and
5	I think spent about a month in Alaska trying to
6	do the make the trip.
7	Okay, why don't we take a break? A
8	reminder, we do have, if you have nothing more
9	to do after you eat your lunch, you can look at
10	the public comments from the last meeting,
11	because we'll be going over those.
12	And then also prepare your Work Group
13	report, and any or reports, and Subcommittee
14	reports, and also you might want to look at both
15	the NIOSH schedule for reports that they that
16	went around from what, with what Ted sent out.
17	And then SC&A sent out their updated
18	scheduling and so forth as a separate email,
19	that I believe everybody's
20	MR. KATZ: Right.
21	CHAIRMAN MELIUS: gotten. So we
22	can try to be prepared. And but the
23	highlight of the afternoon, get all prepared,

1	the highlight will be the first at 1:30, so
2	be on time, LaVon Rutherford will give us his
3	SEC update.
4	MEMBER MUNN: Cliffhanger.
5	CHAIRMAN MELIUS: Cliffhanger, lots
6	of questions, should be a very volatile session
7	so be prepared. You don't want to miss it.
8	MR. KATZ: Sharpen your knives.
9	
10	(Whereupon, the above-entitled
11	matter went off the record at 11:56 a.m. and
12	resumed at 1:36 p.m.)
13	CHAIRMAN MELIUS: Welcome back and
14	we'll now move on with our agenda. Let me check
15	on people on the line. Okay.
16	MR. KATZ: I'm getting to you.
17	CHAIRMAN MELIUS: I'm only doing it,
18	he said like it was good advice. And I'll now
19	let the Designated Federal Official do his
20	designated duties.
21	MR. KATZ: Thank you. Thank you very
22	much. Just checking, first, roll call. I
23	know who's in the room. Everyone's in the room

1	who was here before. But on the line, Bill, are
2	you with us again?
3	DR. FIELD: I'm present
4	MR. KATZ: Dr. Bill?
5	DR. FIELD: and attending.
6	MR. KATZ: Super. And we were
7	missing before Mr. Griffin, is he still
8	missing? Is he on the line? Okay. And how
9	about Dr. Poston? Okay. So that takes care
10	for roll call then.
11	CHAIRMAN MELIUS: Is Dr. Lockey here?
12	MR. KATZ: Lockey, we knew.
13	CHAIRMAN MELIUS: Yes, but if you do
14	the roll call
15	MR. KATZ: Okay. And Dr. Lockey, are
16	you on the line?
17	No Dr. Lockey either.
18	CHAIRMAN MELIUS: Okay.
19	MR. KATZ: And let me just a couple
20	other things. Remind folks who've joined us on
21	the phone to mute your phones. Press *6 if you
22	don't have a mute button, that'll mute your
23	phone. And press *6 again to take it off of

1	mute. But please keep it on mute except when
2	you're addressing the group.
3	CHAIRMAN MELIUS: Public comment.
4	MR. KATZ: And one other thing.
5	Exactly, public comment. Thank you, Dr.
6	Melius. We have a public comment session that
7	begins at 4:30 and goes at least until 5:30 or,
8	well, until 5:30 or until we run out of public
9	comments. That comes before.
10	So for people who've joined us in the
11	room, there's a sign-up book outside. If you
12	want to make public comment during the public
13	comment session, please sign the book.
14	For people on the line, you don't need
15	to sign in. We will get to folks on the line
16	after we've gotten through everyone who's in
17	the room here during public comment session.
18	And that's it. Thank you. Dr. Melius.
19	CHAIRMAN MELIUS: Yes. And just one
20	more thing on public comment. I'm not sure
21	we'll do it today, but there will be a
22	presentation on Santa Susana at 4:00 and
23	immediately following that presentation we

will start the public comment period. So if it starts a little early, that would be fine.

MR. KATZ: Yes.

CHAIRMAN MELIUS: Yes. Good. And

now I'd like to introduce the highlight of the meeting, the SEC petitions status update. LaVon Rutherford, I believe it is.

MR. KATZ: Yes.

MR. RUTHERFORD: Thank you. It's nice to be the highlight, that's for sure. I'm going to give you a Special Exposure Cohort update and then I'm going to take all the drillings and the questions that you guys are going to have afterwards.

All right. The purpose, obviously, as we do the -- I'm usually loud enough anyway, but that's okay. We do this at every Advisory Board meeting. We give the update of upcoming SEC petitions and existing petitions, petitions that are in different phases. This gives the Board updates and allows them to prepare Work Group meetings and other Advisory Board meetings.

Okay. As of October 28th, we had a little bit of an increase here in the number of petitions we received. We're up to 222. We have four petitions in qualification process.

If you recognize that we went quite some time without receiving a petition and here recently we've gotten, I think, about six in the last four or five months. And you can see the status on the existing petitions. We have two that are in the evaluation process right now.

petitions The that in are qualification: Westinghouse Electric Corporation, Bloomfield -- this is for the residual period at Westinghouse. It is going to qualify. I will let you know that. there is some, you know, we have found some issues with, not the petition, but in our early reviews of documentation, we actually found indication there may have been work involved at Westinghouse Electric Corporation. During this residual period they did some work for Fernald.

And so we have actually provided that

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documentation to the Department of Labor so they can evaluate if those actually should be covered operational years instead of residual years.

SEC 220 is for Y-12. This is 1944 to '45. You guys, I think everybody knows that we already have an SEC for 1944 and '45 period. highly Therefore, it's unlikely this petition's going to qualify. The only way this petition would qualify is if they presented evidence that incidents such that, you know, exposure -- I mean, so -- I can't think of the What's the right word I'm looking for? It's such a heavy word, Presence. There. presence. If we had a incident, such as criticality or something like that, that ultimately we could move from the 250 days to presence. They haven't provided that on this one, so it's unlikely that it would qualify.

Lawrence Livermore National Lab, this is the post-SEC years at Lawrence Livermore. This is in the early stages of qualification, but it does look like it will qualify.

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And then we just recently got a Grand Junction Operations Office. We're just now going through that. Most of you will remember we're already evaluating those years at this time anyway.

Two petitions that have qualified and we're moving forward with the evaluation: Dow Chemical Corporation. This is actually here in California, 1947 to '57. We are almost finished with this evaluation at this time. We did get slowed up a little bit with some funding issues at OSTI, you know, to look at some of the documents that Dow had there.

But this Evaluation Report should be out within the next month or two. Anyway, it says January 2015, but that's -- you know, hopefully we'll get those documents sooner than that.

Idaho National Laboratory, this evaluation's been ongoing. There's been a lot of work going on both with the Site Profile work and concurrently the SEC evaluation that's going on at the same time. We expect to complete that evaluation by February 2015.

There's still, obviously, going to be a lot of work going on with that one.

And then Kansas City Plant, we presented that some time ago. The Board has been reviewing that, and the Board Work Group and SC&A. There's been a lot of activity with that one was well: interviews, on-sites and data captures and such.

We have a number of sites that have portions of their petition evaluation that are open. They still need to be resolved. Fernald, we have 1984 to 1989. I think they're getting very close to closing things out at Fernald.

Grand Junction's Operations Office, the '75 to 2006. This one will be presented. We are going to present an addendum or revision depending on how that lays out at the March Board meeting.

Hanford, as mentioned earlier in the meeting, there's been a lot of work going on with the '84 to '90 period at Hanford. And we hope to be able to take some action on that one

as well at the March meeting.

Los Alamos National Lab, this one's been a struggle. We really tried to -- the post-1994 period, '95 period, we've taken an approach of this is a 10 CFR 835 era, and we've taken the approach to see how the site is implementing 10 CFR 835. We're struggling a little bit getting the documentation from them on that. We went back and forth and we decided to take a project or something that was going on during that period, maybe an exotic, and look how they were complying with 835 just to see that, you know, that they were following 835 and that dose reconstruction would be feasible.

We did just recently get some information from them and hopefully we'll be able to move forward with that.

Rocky Flats Plant, as I mentioned earlier today, we have roughly five items, open issues. And of those, we've completed papers on three. Another paper will be out very shortly and we should be able to have a Work Group meeting. After SC&A's had a period of

time to review that neptunium report, we ought to be able to have a Work Group meeting.

Sandia National Lab-Albuquerque, this, again, we're looking at the 10 CFR 835 implementation at the site. It has slipped somewhat, mainly because of the priorities that we have with other activities. Obviously, new SEC petition evaluations, because of the 180 days, are going to take precedence. And so certain things, we adjust priorities based on that, based on what the Board is currently really wanting or looking at. And so the post-'94 period at Sandia has slipped a little bit.

Santa Susana Field Laboratory, I think I will let Dr. Hughes handle this one later on.

And Savannah River Site, continues to be a lot of activity at Savannah River. We were slowed down considerably because of classification reviews on documents. I think we've got that issue -- or at least we've got a path forward worked out. And we should get

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some of the documents and things released here soon so we can start making some progress there.

St. Louis Airport Storage Site, this one's kind of in limbo, mainly because we presented our evaluation. And maybe what we ought to do is, you know, we indicated the '72-'73 period, 1984-'98 period, we felt dose reconstruction was feasible.

It might be appropriate at the next Board meeting or, you know, if there's too much on the Board meeting, maybe during the Board conference call, that I provide a status update, you know, and basically get things moving on that period again.

And potential 83.14s, again, these have been on the plate for a while, mainly waiting on a litmus claim that we could move them forward. There's really no claims that are being negatively affected by this, us waiting, because there are no claims, you know, in that period. 1945 to 1948, that was the old Division Z at LANL. It's now Sandia-Albuquerque. And as soon as we get a

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1	litmus claim, we'll move that forward.
2	The Dayton Project was a facility
3	designation change so we had a limbo period of
4	nine months. However, currently all the
5	claims are covered by another, existing SEC, so
6	we have no litmus claims there as well. And
7	that's it. Questions?
8	CHAIRMAN MELIUS: Yeah, Paul.
9	MEMBER ZIEMER: LaVon, on St. Louis,
10	can you remind us when the original petition
11	came to us and what action did we take?
12	MR. RUTHERFORD: Yeah, I can do a
13	brief reminder on that because I don't
14	MEMBER ZIEMER: I don't need all the
15	details but
16	MR. RUTHERFORD: I was the one who
17	presented it, so I know a lot of it. That
18	actually it's probably been, I don't know,
19	four or five years ago that it was presented
20	originally. We recommended adding a Class
21	during the operational period. There was a
22	period of time, basically, the site was a

They had some of the K-65

storage site.

materials, the African ores that were stored out there for a very brief time. Some worker was doing some raffinates. We ultimately added a Class for that operational period.

Then there went to a stagnant period where it was basically closed down, and then they went through a clean-up period in the 1972-'73, I think, timeframe. And then there is additional clean-up work that went on later on, if I remember correctly. That may not be totally, you know, accurate, but it's generally in that manner.

And what we found was, during the '72-'73 time period, or during, you know, those later years, we had additional information. We had monitoring data that allowed us to dose reconstruction. But there was never a formal recommendation by the Board on that period.

MEMBER ZIEMER: Okay. That was really what my question was. Maybe Dr. Melius remembers. So it was that latter period, and did we send it back for some review or did we do anything?

1	MR. RUTHERFORD: No.
2	MEMBER ZIEMER: We didn't take action?
3	MR. RUTHERFORD: Yeah, there was a
4	period of time, you know, that we didn't take
5	action on some of the residual periods and
6	things like that. And this was kind of a
7	residual period. And there was never really a
8	follow-on on that one.
9	CHAIRMAN MELIUS: What's a A kind of
10	a residual period?
11	MR. RUTHERFORD: Well, when I say a
12	kind of a residual period, I mean, because there
13	was some remediation work that went on in that
14	period as well, in addition to that residual
15	period.
16	CHAIRMAN MELIUS: Okay. And have we
17	had SC&A review that?
18	MR. RUTHERFORD: I do not believe so.
19	CHAIRMAN MELIUS: Okay. So, maybe
20	it's not appropriate to do it today, but if you
21	think you have enough information, maybe we can
22	get this moving ahead.
23	MR. RUTHERFORD: Okay.

1	CHAIRMAN MELIUS: Is there enough
2	information in the Evaluation Report to form
3	the basis for evaluating the residual period,
4	or is there additional information that's not
5	in the report?
6	MR. RUTHERFORD: No, I think it's all
7	there. I think. And all the supporting
8	documents are referenced and such.
9	CHAIRMAN MELIUS: Okay.
10	MR. RUTHERFORD: If you would like, I
11	could put
12	MEMBER ZIEMER: Well, I was wondering
13	if there was an actual recommendation from
14	NIOSH and if we just chose not to act. And did
15	we specifically choose not to act?
16	MR. KATZ: I think we just covered
17	MEMBER ZIEMER: Or did we just forget
18	to act or what happened? I just don't
19	remember, but I can attribute that to my age.
20	But somebody needs to know what happened.
21	CHAIRMAN MELIUS: Well, LaVon is
22	fairly young.
23	MEMBER ZIEMER: I don't need to know

1	today, but I'd like a follow up on
2	CHAIRMAN MELIUS: When he was a high
3	school senior, he went to the same football
4	games I was going to in Cincinnati, in '83.
5	(Laughter.)
6	MR. RUTHERFORD: What I can do is
7	provide the Board and SC&A, basically, a
8	summary of, you know, where we were when and,
9	you know, the dates and also point to
10	MR. KATZ: The transcript.
11	MR. RUTHERFORD: the documents and
12	such.
13	MR. KATZ: Well, and the transcript
14	from the Board meeting.
15	MR. RUTHERFORD: Oh yes, exactly.
16	MR. KATZ: But basically the
17	transcript so it wasn't set aside to be
18	addressed later, but the Board only
19	specifically took the action that was being
20	recommended, which was to add the Class for the
21	operational period. And it just wasn't spoken
22	to.
23	MR. RUTHERFORD: Right. Okay.

1	MEMBER ZIEMER: But there was another
2	recommendation. That's
3	MR. KATZ: Well, there was a
4	recommendation, but it wasn't spoken to, is
5	what I'm saying. The Board didn't speak to it.
6	CHAIRMAN MELIUS: So, what I would
7	suggest is that for the Board call, the next
8	call, can you put together a short presentation
9	on the residual period?
10	MR. RUTHERFORD: Yes, I can.
11	CHAIRMAN MELIUS: And then we'll do
12	that and we can either take action at the Board
13	call or we can refer it on for additional
14	MR. RUTHERFORD: Yes.
15	CHAIRMAN MELIUS: We can either
16	accept the recommendation or we can refer it on
17	for further review. And I think that would
18	probably be a better way of doing it. Does
19	everybody agree?
20	MR. RUTHERFORD: Yes.
21	MEMBER ZIEMER: Thank you.
22	CHAIRMAN MELIUS: Okay. Yes, thank
23	you on that.

1	Other questions for LaVon? So LaVon,
2	could you just sort of go over what is going to
3	be available for the March meetings?
4	MR. RUTHERFORD: Yes.
5	CHAIRMAN MELIUS: And I hope the
6	transcriber
7	MR. RUTHERFORD: Can you pull that
8	back up, please?
9	CHAIRMAN MELIUS: listens very
10	carefully here and keeps track of this.
11	MR. RUTHERFORD: I'm waiting for my
12	presentation to come back up again so I can go
13	back. No, actually, we will be presenting Dow
14	Chemical Walnut Creek Petition Evaluation
15	Report. We will plan to present the Idaho
16	National Lab Evaluation Report. The Grand
17	Junction Operations Office, we plan to present
18	that as well.
19	Then I expect action to be taken on
20	Hanford. Some kind of action at least. I believe
21	so, anyway.
22	There potentially could be action on
23	Rocky Flats, just depends on the release of

1	documents and where the Work Group goes on that
2	one. So there's four or five, anyway.
3	CHAIRMAN MELIUS: Okay. What is the
4	timing on Grand Junction?
5	MR. RUTHERFORD: When will the
6	addendum or the evaluation revision, whichever
7	way we end up going, when will that be out?
8	CHAIRMAN MELIUS: Yeah.
9	MR. RUTHERFORD: January/February
10	timeframe.
11	CHAIRMAN MELIUS: Okay.
12	MR. RUTHERFORD: That
13	CHAIRMAN MELIUS: Go ahead.
14	MR. RUTHERFORD: I was going to say,
15	that report actually would've been out earlier,
16	but in our additional reviews we found some
17	thorium sources that were not previously
18	recognized that we need to look at that.
19	CHAIRMAN MELIUS: I'm just trying to
20	determine are there any of these that are
21	straightforward enough that would be done in
22	time for our January conference call?
23	MR. RUTHERFORD: Well, I'll see if

1	Jim agrees with me. I think Dow Walnut Creek
2	would be pretty yeah, I think Dow Walnut
3	Creek would be pretty straightforward enough.
4	I think it'll be ready.
5	CHAIRMAN MELIUS: Well, when?
6	Because our call is early in January.
7	That's
8	MR. KATZ: Very early in January.
9	CHAIRMAN MELIUS: Yes. So it would
10	have to be before the holidays.
11	MR. RUTHERFORD: No, that's not going
12	to work.
13	MR. KATZ: Yeah, I don't think so.
14	CHAIRMAN MELIUS: Okay. That's
15	fine. Just checking. So, Dow, Idaho, Grand
16	Junction and Hanford, okay.
17	Any other questions for LaVon, here,
18	now that he's got us confused? Okay. You're
19	off the hook for now.
20	Okay. We now have our Board work
21	session. And we've completed part of it. So
22	we will start with the public comment session
23	from our last Board meeting.

1	And you all should have gotten two
2	documents, one is the summary spreadsheet and
3	the other is the transcripts that sort of back
4	that up and provide a little bit more detail on
5	that.
6	And I will go through this relatively
7	briefly, but feel free to interrupt me or if you
8	have questions and so forth. There were a
9	number of first comments had to do with Simonds
10	Saw and Steel. And there was some questions
11	about the basis for the dose reconstruction and
12	the follow-up clean-up there. Those have been
13	addressed and responded to.
14	MR. KATZ: Which document?
15	CHAIRMAN MELIUS: It's a
16	spreadsheet.
17	MR. KATZ: A spreadsheet, Excel.
18	CHAIRMAN MELIUS: And it's got a
19	funny name to it. Yes, what is BPCP? Board
20	MR. KATZ: Board public comment
21	session, or whatever the comment
22	presentation.
23	MEMBER ZIEMER: Oh, it's a

1 spreadsheet. I've got it. CHAIRMAN MELIUS: Okay. Then we 2 have a set of three comments regarding the Santa 3 And those have all 4 Susana site. 5 addressed, including some follow-up I think we=ll probably hear about a little bit later 6 7 today. There's some questions on the INL site 8 9 and comments from one of the people at that meeting. And, again, I think these are all 10 straightforward in terms of being addressed. 11 12 There's actually a whole series of 13 comments on INL that we heard, which were, as I recall, very helpful in terms of further 14 15 background on that site. And we'll probably be 16 talking more about it. It was guite a long 17 comment, as you may remember. 18 Okay. Then we have some general 19 comments, some comments on Hooker, General Steel Industries and on Dow Madison. And these 20 21 are being followed up on or in the process of

Some comments on the Blockson site,

being followed up on.

22

1	which was more of an issue having to do with DOL
2	and sort of a facility designation. There was
3	another comment on the Blockson site and,
4	again, sort of a what-if, procedural issue,
5	which, again, I think was addressed actually
6	right at the meeting by LaVon.
7	And there was some additional
8	follow-up reported from the April Board
9	meeting. It was something new. You're adding
10	Boulder, Ted?
11	MR. KATZ: It was probably commented
12	on that we would follow-up on that.
13	CHAIRMAN MELIUS: Oh, okay. Okay.
14	That's the first I've seen. Longer term
15	follow-up. But, again, there was a follow-up
16	to some comments made at the April Board meeting
17	and a conference call and NIOSH and ORAU had
18	followed up and addressed that.
19	So any Board Members questions or
20	comments on that? I think the level of
21	documentation's good and okay. I don=t
22	think we need any further action on that.

Now we'll move onto Subcommittee and

1	Work Group reports. Josie's not here. The
2	first one I have on my list, which was off the
3	website, is on Brookhaven. I don't think
4	there's much going on there. Henry, do you
5	MEMBER ANDERSON: Nothing.
6	CHAIRMAN MELIUS: Nothing. I think
7	it's
8	MEMBER MUNN: there's no meeting.
9	MEMBER ANDERSON: No.
10	CHAIRMAN MELIUS: Fernald, Brad?
11	MEMBER CLAWSON: On Fernald, we've
12	really got one outstanding issue that we're
13	still working on. That's the thorium. That's
14	in SC&A=s hands. They're supposed to have a
15	paper for us in about a month or so.
16	CHAIRMAN MELIUS: Okay. And that's
17	on schedule, John?
18	MR. STIVER: Yes, it is.
19	CHAIRMAN MELIUS: Okay. Well,
20	Hanford, we've talked about there's some active
21	evaluation going on and data gathering
22	regarding some issues out at Hanford. And we
23	will be doing a follow-up there.

1	So we can expect a Work Group meeting
2	before the March meeting. And I expect we'll
3	be in a place to take action at the March meeting
4	on that. I think that I understand from both
5	Sam and from talking to Arjun about that.
6	Thank Sam for his communication on that one.
7	Idaho, Phil?
8	MEMBER SCHOFIELD: There are
9	MR. KATZ: Use the mic, please.
10	MEMBER SCHOFIELD: Oh, okay. There
11	are more worker interviews scheduled in about
12	a week and a half that will hopefully shed a lot
13	more light on some of the areas that we're kind
14	of weak on. And that's really where we stand
15	at this point.
16	CHAIRMAN MELIUS: Okay. And then we
17	have the SEC Evaluation Report for March, so
18	that will pull that together. Okay. And
19	we've been assured that that's on schedule?
20	MR. RUTHERFORD: Yes.
21	CHAIRMAN MELIUS: Okay.
22	MR. RUTHERFORD: But it's on schedule
23	now. There's always things we can come up

I would like to point out this is a very 1 unique situation in that we have a Site Profile 2 review going on at the same time that an SEC 3 evaluation is going on. So it has been a 4 coordination trick, I can tell you. 5 CHAIRMAN MELIUS: Well, and it's also 6 combining two sites and --7 MR. RUTHERFORD: 8 Yes. 9 CHAIRMAN MELIUS: Yeah, which were 10 initially evaluated as separate sites and now have been combined. 11 12 MR. RUTHERFORD: Actually, we have to 13 separate them from -- initially, the idea was that they thought they were going to be able to 14 It's going to be two 15 do it as one petition. 16 petition evaluations. The second petition 17 evaluation will be for the Argonne Lab itself 18 because, regulation-wise, we can only do it by 19 site, you know, for a single site. 20 So we've got a petitioner providing a 21 second petition for that separate site, which will qualify and we'll move it forward. 22

quess I should have mentioned that earlier.

1	CHAIRMAN MELIUS: Yeah. Yeah.
2	(Laughter.)
3	MR. RUTHERFORD: I apologize for
4	that. I forgot about that, I guess.
5	CHAIRMAN MELIUS: Okay. Got me
6	confused. So what's the timeframe on the
7	second petition?
8	MR. RUTHERFORD: It will not be ready
9	for the March meeting, but it should be shortly
10	thereafter just because we're doing data
11	gathering for both sites.
12	So, the interviews, the data captures
13	and all are going on concurrently. I just
14	don't think that, from a schedule standpoint,
15	we'll be able to produce both of them at the same
16	time in order for the March meeting.
17	CHAIRMAN MELIUS: Okay. So, don't
18	go away.
19	MR. RUTHERFORD: I'm not.
20	CHAIRMAN MELIUS: Do they overlap?
21	What extent do the petition evaluations
22	overlap, in terms of
23	MR. RUTHERFORD: Years, are you

1	talking about or
2	CHAIRMAN MELIUS: Years, operations?
3	MR. RUTHERFORD: Well, it's not clear
4	yet. And this is, again, this is really
5	difficult because this facility sits inside of
6	the main facility
7	CHAIRMAN MELIUS: Yeah, right.
8	MR. RUTHERFORD: And also this is very
9	similar to Y-12. If you remember back when we
10	did the early years at Y-12, we had facilities
11	at Y-12 that were turned over to Oak Ridge
12	National Lab and they were doing work with
13	calutrons and cyclotrons.
14	And, so, in this situation, you've got
15	working going on at Idaho that it's actually
16	being done by the Argonne crew, but at the Idaho
17	facility. So there's a lot of little nuances
18	that are going to make it difficult.
19	CHAIRMAN MELIUS: Okay. Yeah. I'm
20	also thinking, in terms of the review, where
21	we've essentially combined the Site Profile
22	reviews, and now we're separating them out

again.

1	MR. RUTHERFORD: Yeah, right.
2	CHAIRMAN MELIUS: Yes. Yes.
3	Thank you, LaVon. K-20, gaseous
4	diffusion plants. Phil, anything to report on
5	them?
6	MEMBER SCHOFIELD: Nothing to report
7	this time. We're kind of a little in the dark
8	here for recently, but we need to get a meeting
9	put together and try and get those closed out
10	if at all possible.
11	CHAIRMAN MELIUS: Yeah, it's the Site
12	Profile.
13	MEMBER SCHOFIELD: This is the Site
14	Profile issues.
15	CHAIRMAN MELIUS: Kansas City,
16	Josie's not here. Brad, do you want to?
17	MEMBER CLAWSON: Yeah, we've had
18	quite a bit of data capture up through there.
19	Matter of fact, it was just a little while ago.
20	We're proceeding on with the work on it. We're
21	waiting kind of, and it's in NIOSH's court to
22	respond that they have to put their mark in the
23	sand, but we've had some good data capture up

1	there and we'll go on from there.
2	CHAIRMAN MELIUS: Good. Timeframe,
3	do we have one?
4	MR. RUTHERFORD: Well, unless I'm
5	wrong here, I mean, we provided our evaluation.
6	It's now in the Board and Work Group=s hands to
7	provide a response to the evaluation.
8	Now, there is a lot of work that's
9	going on to gather interviews and such, but we
10	have committed to now products beyond
11	because we haven't gotten anything from SC&A or
12	anything to review at this point.
13	MEMBER CLAWSON: Okay. Well, I was
14	understood from Pete that since we've dove into
15	this a little bit that there's the ER is being
16	revised.
17	MR. RUTHERFORD: Okay. Pete has not
18	said anything to me about that.
19	MEMBER CLAWSON: Okay. Well
20	MR. RUTHERFORD: And this is the very
21	first time I've ever heard. I'll talk to Pete.
22	MEMBER CLAWSON: No, I could be wrong
23	on that but, you know, we're proceeding on, so

1	we're trying to get into where we can get into
2	a Work Group and then move on from there.
3	CHAIRMAN MELIUS: And does SC&A have
4	anything to add or Joe?
5	MR. FITZGERALD: Yeah, we're still in
6	the issue resolution. I mean, we're
7	identifying issues as we go. There's new
8	issues that weren't covered in the ER that we
9	have now identified.
10	So, this is a transition period where
11	we've gone from the ER to one of actually the
12	Work Group and SC&A identifying issues,
13	bringing them to the floor. It's being done in
14	conjunction with NIOSH, so there's a lot of
15	interchange. But, how should I put it, the
16	dust hasn't really settled on what the issue
17	slate is, but we're getting close to having
18	that. So we should be able to
19	CHAIRMAN MELIUS: And as you're
20	identifying issues, are you sort of separating
21	out SEC issues from Site Profile issues?
22	MR. FITZGERALD: That's, I mean,
23	that's the process. I think, really, in a lot

1	of the research and interviews and everything,
2	we've done a lot. It's directed to sifting out
3	what may have been Site Profile issues earlier
4	in the year to ones that are standing as
5	potential SEC issues.
6	CHAIRMAN MELIUS: Okay.
7	MR. FITZGERALD: And that potential
8	part is taking some time to really get a feel
9	for it.
10	CHAIRMAN MELIUS: And some of the
11	potential ones may not be fully addressed in the
12	SEC Evaluation Reports?
13	MR. FITZGERALD: That's right. And
14	that means you're doing fundamental research
15	onsite. So this is new stuff that's really
16	being looked at.
17	CHAIRMAN MELIUS: Okay.
18	MR. KATZ: So the next product will be
19	an SC&A evaluation review.
20	CHAIRMAN MELIUS: Right. And, yeah,
21	so there'll be an SEC evaluation review and
22	that'll be the basis for a Work Group meeting.
23	Okay. Good. Lawrence Berkeley?

1	MEMBER ZIEMER: NIOSH is still
2	reviewing information from the most recent data
3	captures there. And I was just checking my
4	emails, and I didn't get to the right one, but
5	Dr. Hughes is here. But I believe she
6	indicated to me that it will probably be early
7	next year, maybe January, before NIOSH finishes
8	the last White Paper. And she's nodding yes,
9	that that's correct.
10	Now, we have several White Papers
11	already that have been prepared earlier and
12	those have actually been also reviewed by SC&A,
13	but we're waiting for this final group of White
14	Papers so we have all the issues from Lawrence
15	Berkeley. And then we'll have an opportunity
16	for SC&A to review those and then we will meet.
17	CHAIRMAN MELIUS: Okay. Thank you,
18	Paul.
19	Linde. Gen, is there anything?
20	We're done, right? So that should be inactive?
21	MR. KATZ: Yes.
22	CHAIRMAN MELIUS: Okay.
23	MR. KATZ: It's inactive.

1	CHAIRMAN MELIUS: It doesn't say so
2	on the website.
3	Okay. LANL? Mark's not here. I
4	don't know if there's any yeah. You had it
5	on your
6	MR. RUTHERFORD: Yeah, I think I
7	provided the update. We're working that 10
8	CFR. 835 implementation with this site and we
9	just got all the information from them back in,
10	I think, October.
11	And so we should be able to move
12	forward here and provide something to the Work
13	Group. You know, I don't know, I don't want to
14	commit to a date, but it'll be soon because
15	they're still reviewing how much information we
16	got. Okay.
17	CHAIRMAN MELIUS: How about an, you
18	know, an estimate?
19	MR. RUTHERFORD: A couple months.
20	CHAIRMAN MELIUS: Okay. It's in the
21	transcript.
22	MR. KATZ: I think we got him.
23	CHAIRMAN MELIUS: We'll remember.

1	Okay. Mound. Josie's not here
2	also. Brad, do you have any update? I'm not
3	sure there's much activity there.
4	MEMBER CLAWSON: There hasn't been
5	much activity at all on that.
6	CHAIRMAN MELIUS: Have we completed
7	the Site Profile? I don't recall.
8	MEMBER CLAWSON: No, we
9	CHAIRMAN MELIUS: Oh, here, Jim
10	has
11	DR. NETON: There's still a hold up on
12	our issuance of the review of the external dose
13	section of the Site Profile due to
14	neutron-photon ratio issues.
15	CHAIRMAN MELIUS: Oh, right.
16	DR. NETON: Dr. Taulbee's working on
17	that and his schedule's been pretty full, but
18	I think it's high on his priority list right
19	now.
20	CHAIRMAN MELIUS: Okay. Nevada Test
21	Site. Brad?
22	MEMBER CLAWSON: Yeah, we've got a
23	Work Group meeting coming up on that, I believe,

December 3rd. And all of these are Site 1 Profile issues. 2 SC&A has reviewed the matrix. We've 3 4 just got to sit down with NIOSH and basically 5 close out the Site Profile issues. That should be it for Nevada Test Site. 6 7 CHAIRMAN MELIUS: Good. Thank you. When I saw the announcement on the Work Group 8 9 meeting, I expected it to get recalled, that it 10 had the wrong name on it. We hadn't seen Nevada Test Site for a while, so thanks for keeping 11 12 that moving, Brad, and everybody involved in 13 that. X-10, Ridge National 14 0ak 15 Laboratories. Gen? MEMBER ROESSLER: Dr. Taulbee's not 16 17 here and I've been waiting for word from NIOSH. 18 Does anybody else have any update on that? 19 MR. RUTHERFORD: Yeah, we completed 20 everything with the petition evaluation 21 before. This is now, there was a post period 22 after the SEC period that we were looking at additional works and exotics and things. 23

1	And we went and retrieved a number of
2	log books that identified air sampling and
3	such. We've been working through those. We
4	also got into the difficulty of getting the
5	documents released from a classification
6	standpoint, so that slowed us down. But we
7	have all the documents now and we can move
8	forward with that.
9	The problem we got is, to be honest,
10	is resources. We're balancing priorities
11	right now. And, you know, if we see something
12	that's clearly looks like it's going to be an
13	SEC issue, we'll move that to the forefront.
14	But right now, we're working through
15	all those documents and, hopefully, we'll have
16	something to the Work Group. I'm not going to
17	commit more than in the next six months.
18	CHAIRMAN MELIUS: Okay.
19	MR. RUTHERFORD: I just looked and
20	November, December, January are rough right
21	now.
22	CHAIRMAN MELIUS: Okay. No, I
23	think, as I recall, when we did the original SEC

1	evaluation there was uncertainty as to whether
2	there was uncertainty as to whether there would
3	be other ones sort of going forward there. I'm
4	not sure I'm remembering the competing
5	priorities discussion, but at least the general
6	basis for it, you said. Gen, do you have
7	anything to add or is that
8	MEMBER ROESSLER: Thank you. No,
9	we're just waiting.
10	CHAIRMAN MELIUS: Okay. Pacific
11	Proving Ground, Dr. Lockey isn't here. I don't
12	believe there's been a meeting or
13	MR. KATZ: No.
14	CHAIRMAN MELIUS: Okay. All I do is
15	keep getting emails about when is the site
16	visit. I'll probably get more of those.
17	MEMBER MUNN: You know, we all want to
18	go.
19	CHAIRMAN MELIUS: Dr. Lemen's going
20	to be out in that general area later this month,
21	Australia and Indonesia. So maybe you can take
22	a sail over them.
23	MEMBER LEMEN: I may just drop by.

1	CHAIRMAN MELIUS: Yes.
2	MEMBER LEMEN: Then they won't have
3	to have a site visit. I'll just go over.
4	CHAIRMAN MELIUS: No, well
5	(Simultaneous speaking.)
6	CHAIRMAN MELIUS: Okay. I skipped
7	over Pantex.
8	MEMBER CLAWSON: Yeah, we had a Work
9	Group meeting, along with Fernald here, about
10	a month, month-and-a-half ago. Everything on
11	Pantex is pretty well taken care of, the Site
12	Profile issues.
13	But we still have the neutron-photon
14	ratio issue that, I believe, has been our
15	overarching issue several times. And that's
16	the only thing that we have left on that.
17	DR. NETON: Actually, we decided not
18	to use the neutron-photon ratio approach at
19	Pantex because of some issues that we had with
20	that. And we're now redeveloping a coworker
21	model just based on the actual neutron dose
22	distributions. And that's in the works. It's

a couple months out. Yeah, and these are for

1	non-SEC cancers because Pantex is an SEC
2	after a certain period of duration of the
3	covered period.
4	
5	CHAIRMAN MELIUS: Okay.
6	MEMBER CLAWSON: So once we have that
7	paper, we'll finish up Pantex.
8	CHAIRMAN MELIUS: Okay. Good.
9	Thank you both. And, Jim, we'll also remember
10	a couple months out.
11	Pinellas?
12	MEMBER SCHOFIELD: We haven't done
13	much on Pinellas right now. It's kind of like,
14	just like the gaseous diffusion plants, and
15	we'll hopefully get together and close that one
16	out. I don't think there's a whole lot left on
17	that that we have.
18	CHAIRMAN MELIUS: Jim's going to
19	complicate things.
20	DR. NETON: Well, this is going to
21	sound like a broken record about Pinellas, but
22	there's only issue remaining at Pinellas,
23	outstanding, and that's the reconstruction of

tritide doses. And we're still trying to figure out whether they filtered the smears before they measured them or not, which, if they did, causes some issues with trying to reconstruct the tritide exposures.

That's a little bit out, though, on the schedule and that's one of those prioritization-type issues. I think it's out into January next year. So early next year, it's on schedule, is my recollection.

CHAIRMAN MELIUS: What's involved in finding that out?

DR. NETON: I think there's some more interviews that have to be done. They're searching through the health physics documentation, the records. Because we just have one indication that they did filter these samples. But there's got to be some other health physics documentation to substantiate that somewhere, why they did that in the first place, you know, or maybe they didn't and there's other documentation to address that.

CHAIRMAN MELIUS: Okay. Thanks for

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1	the explanation.
2	Rocky Flats I think we've already
3	pretty much addressed. Probably should have a
4	Work Group meeting between now and the
5	MEMBER MUNN: Meeting in December.
6	CHAIRMAN MELIUS: Oh, it's already
7	scheduled. Okay. Good.
8	Sandia, Dr. Lemen?
9	MEMBER LEMEN: I don't have anything
10	new to report. I'm waiting for Sam. Does Sam
11	got anything new?
12	MR. RUTHERFORD: Yeah, I think I
13	actually talked about that a little earlier,
14	that we did get a number of documents from
15	Sandia, actually, back in September.
16	We are actually supposed to get more
17	documents later on this month. But I think our
18	schedule right now doesn't have us really
19	completing things until sometime in April of
20	next year.
21	MEMBER LEMEN: So we haven't planned
22	any Board Working Group meetings until after we
23	get data back from them.

1	CHAIRMAN MELIUS: Right.
2	MR. KATZ: Can we go back to Rocky
3	Flats? Is that Rocky Flats you said we have a
4	meeting in December? No, we have a Dose
5	Reconstruction meeting in December.
6	MR. RUTHERFORD: Nevada Test Site.
7	MR. KATZ: Nevada Test Site and
8	Fernald. Those are in December. No Rocky
9	Flats meeting in December. We have not
10	schedule a Rocky Flats, because I've not
11	contacted Mark about this.
12	MEMBER MUNN: I thought he had
13	scheduled.
14	MR. KATZ: No. We have NTS on the
15	3rd, Fernald on the 4th and Dose Reconstruction
16	on the 8th. That's it.
17	CHAIRMAN MELIUS: Okay.
18	MR. KATZ: Anyway. I just wanted to
19	get that straight.
20	CHAIRMAN MELIUS: Savannah River,
21	Mark isn't here. And I think we're waiting for
22	some NIOSH reports?
23	MR. KATZ: I think so. Yes, we're

1	waiting on NIOSH. Well, SC&A's been working on
2	matters too. I don't know whether they have a
3	report coming out too.
4	CHAIRMAN MELIUS: Yes, we have an
5	SC&A report that we're waiting for for the SEC
6	evaluation, which is a coworker.
7	MR. KATZ: Exactly.
8	MR. RUTHERFORD: And we were waiting
9	for a number of documents to be released from
10	the site from classification review. Now that
11	those documents are slowly coming out, we'll be
12	able to finish up some other papers that we're
13	working on.
14	CHAIRMAN MELIUS: Timeframe?
15	MR. RUTHERFORD: Let's see what Dr.
16	Taulbee has in here. I don't see an update on
17	any. I'll have to get back with the Board on
18	that one.
19	CHAIRMAN MELIUS: Do you have one?
20	MR. RUTHERFORD: I don't see it.
21	MEMBER ZIEMER: There's a date of
22	March 2015 up there.
23	MR. RUTHERFORD: Oh yeah, March of

2015. 1 CHAIRMAN MELIUS: And I also think 2 our sort of coworker guideline issue may impact 3 on this also. 4 5 Scientific Work Issues Group. David. 6 MEMBER RICHARDSON: Since the last 7 meeting I had a chance to talk with NCRP about 8 9 -- we had a topic that we were interested in 10 which was biological effectiveness 11 low-energy photons and tritium. And the NCRP 12 was working on a report on that. I had some back and forth with them 13 about whether we could get a draft of that 14 15 report, or at least the first chapters of it, to review. And, unfortunately, they have a new 16 17 executive director who said that they're not 18 going to release any of the material until 19 publication. They're behind on publication and it 20 21 had been intended to be out by now. 22 hopefully by early 2015 there will be a report

that's available for us to review on that.

1	And the suggestion, again, had been to
2	get maybe David Kocher to come and at least
3	introduce the report and present sections of it
4	for us.
5	CHAIRMAN MELIUS: Okay. Do we want
6	to do that for a Board meeting or for a Work
7	Group meeting?
8	MEMBER RICHARDSON: I think for a
9	Work Group meeting.
10	CHAIRMAN MELIUS: Okay. Jim, you're
11	shaking your head. Is it we=re doing it for a
12	Board meeting or
13	DR. NETON: Well, I think it could
14	start as a Board meeting and then eventually
15	escalate it through a Work Group.
16	CHAIRMAN MELIUS: Okay.
17	MEMBER RICHARDSON: Yes, I was hoping
18	we could digest it some and then maybe
19	CHAIRMAN MELIUS: Okay. SEC Work
20	Group. I think we've got three things
21	underway, if I remember correctly. One is the
22	coworker sufficient accuracy issue which we're
23	already talking about today. And I think you

have a pretty good idea where that is.

Secondly, we have a SC&A review of a Savannah River coworker issue that I think is -- I can't remember if it's just out from DOE review or where. It's very close. Just got it, okay. I'm still on the bad list for my CDC email. So I will get it when I get back to the office next week.

And then the third issue we have.

Once upon a time a long time ago the SEC evaluation group looked at the Dow Madison SEC and so forth. And we now have a follow-up. We have a Site Profile and we have also a PER.

So if this is okay with my fellow Subcommittee chair, Work Groups, I think we would like to task SC&A to really review both, the PER and the Site Profile.

And my understanding, one, that this is sort of a priority, available resources issue also. And my understanding is that NIOSH now has the timing appropriate, is that true, Stu? At least that's what Ted tells me after talking to you.

1	MR. HINNEFELD: Right. The Site
2	Profile has been revised. So there's a revised
3	Site Profile published, so that's certainly
4	available to review. PER is underway now.
5	MR. KATZ: It's about the come out.
6	MR. HINNEFELD: I mean, we haven't
7	actually considered all of those. So
8	certainly the Site Profile revision is
9	available for review.
10	CHAIRMAN MELIUS: It seems sensible
11	to me to combine the two. Is that I'm not
12	familiar with the PER.
13	MR. HINNEFELD: Well, the PER review
14	typically kind of does the revised Site Profile
15	review anyway. You know, they kind of look at
16	the changes that were done and were the changes
17	to the Site Profile appropriate? I think
18	that's one of the tasks, isn't it?
19	And so to our way of thinking, whether
20	they're combined or not, you know, is kind of
21	irrelevant to us. So I suppose it could be
22	combined as one assignment and then if the PER
23	cases aren't completely worked they would, you

1	know, SC&A would just have to wait until they
2	get that sub-task until the cases are worked.
3	CHAIRMAN MELIUS: Yeah, why don't we
4	get it assigned? Dr. McKeel will rest easier
5	and we can get this going. And then face this
6	when the review is done, at the appropriate
7	timing we can then decide if all this Site
8	Profile issues, sort of new issues, revolve
9	around the PER. Then, you know, the chair of
10	the Work Group would be glad to send this all
11	over to the Procedures Subcommittee for action.
12	But if there are others, we can sort
13	of work that out when we get to that point.
14	MEMBER MUNN: We're looking forward
15	to your action.
16	CHAIRMAN MELIUS: Our actions. It=s
17	a group decision. Okay. I hope we haven't
18	confused everybody by that.
19	Paul, maybe our hardest working Work
20	Group, TBD-6000.
21	MEMBER ZIEMER: Well, TBD-6000, I
22	want to report on two different facilities.
23	First of all, General Steel Industries. You

recall that Appendix BB, which 1 basically, what you might call the Site Profile 2 for General Steel Industries, Revision 1 of 3 that was issued this past summer. 4 SC&A was tasked to review the revision 5 and they have just completed that review. 6 We just got the review within the last few days, 7 actually. And I believe once NIOSH has a 8 9 chance to review that and respond, the Work 10 Group will meet. Now, this does also raise the issue of 11 12 the process of a PER, as well, because there's 13 not a PER for this one yet, either. It was my understanding that NIOSH may be wanting to 14 await the review of this one before they 15 16 actually issue a PER in case there is an

I'm not sure which is the best way to do this because in the past some of these PER reviews were actually reviews of the revisions themselves.

additional revision or two.

But we have the revision. I think we'll be prepared fairly soon. Although, I

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didn't see it on the NIOSH worksheet yet when they would have a chance to respond.

My early review of the SC&A report, and I haven't had a chance to review it in complete detail, a lot of the comments were wording things in there, but there is one thing, at least, that's a little more substantial. So the Work Group will have to take a look at that.

But at the moment, we're proceeding just with what we have before us. And it will be up to NIOSH how they want to proceed with the PER in the process.

But that's where we are in GSI. And, personally, I'd like to get the comments closed as quickly as we can because this has been a long process and we want to make sure that there's a sooner rather than later opportunity for any revised dose reconstructions to be handled. Most of them have already been done and many of them, I'm sure, will have to be redone. So we'll need to come to closure on that.

The other thing I'll report on is Simonds Saw, which is also a TBD-6000. Just

within the last few days, I think late last
week, we received Rev 2 of what constitutes
their TBD. This is not an SEC issued Site
Profile and we just received that from NIOSH a
few days ago. SC&A will have to review that
yet. But just to let you know that that is in
the works and at some point we'll have to see
if there's any issues yet on Simonds Saw on the
Site Profile.
CHAIRMAN MELIUS: Okay. Good.
TBD-6001, otherwise known as the Uranium
Refining Atomic Weapons Employers Work Group,
nothing to report?
MEMBER ANDERSON: Nothing to report.
CHAIRMAN MELIUS: Okay.
MEMBER ANDERSON: We've got
additional assignment sites, but they're not,
I don't think, ready for us to look at yet.
CHAIRMAN MELIUS: Okay. Surrogate
Data is in the same position. We have nothing
thing active for that. We're not expecting
anything to be active about, but you never know.
Weldon Spring?

1	MEMBER LEMEN: I have nothing new to
2	report on that.
3	CHAIRMAN MELIUS: What do we have
4	that's old? I don't
5	MEMBER LEMEN: I don't have anything
6	that's old either, unless NIOSH does.
7	CHAIRMAN MELIUS: We haven't done a
8	Site Profile review there, have we?
9	MEMBER LEMEN: I think everything
10	just kind of stopped after the SEC.
11	MR. HINNEFELD: No, I don't recall
12	any Site Profile issues being open from the SEC
13	process.
14	MEMBER LEMEN: That's correct.
15	MR. HINNEFELD: When we finished the
16	SEC process, I didn't think there were any Site
17	Profile issues left. That's my recollection
18	anyway.
19	CHAIRMAN MELIUS: Is there a Site
20	Profile review or was it just an SEC review?
21	MR. HINNEFELD: Well, there was
22	MR. KATZ: Combined maybe.
23	MR. HINNEFELD: Yeah, when an SEC

1	review is done, you know, and the collection of
2	issues are made, sometimes those are parceled
3	into Site Profile issues and SEC issues.
4	And sometimes we'll resolve the SEC or
5	add an SEC Class and get rid of the SEC issues
6	and still have Site Profile issues remaining.
7	But I don't recall that there were any Site
8	Profile issues remaining from the Weldon
9	Springs work.
10	CHAIRMAN MELIUS: But if SC&A, and
11	this is a hypothetical, if SC&A has not done a
12	Site Profile review, then there might be SEC
13	issues. They may have focused just on SEC
14	issues and not focused at all on Site Profile
15	issues, which is why we do separate Site Profile
16	reviews. And I just don't recall on this one
17	what was done.
18	MR. HINNEFELD: I don't recall
19	either. I don't know that
20	CHAIRMAN MELIUS: Well, maybe John
21	Stiver has some.
22	MR. STIVER: After the SEC was closed
23	out, I believe we had a meeting in September of

1	2012. And then all the TBDs were revised after
2	that, in 2013, and we have not looked at those
3	yet.
4	I think it's probably, mainly, some of
5	the superficial changes to incorporate the SEC,
6	but I don't know because we haven't really
7	checked on that yet.
8	CHAIRMAN MELIUS: Okay. Should we
9	task or
LO	MR. KATZ: Why not? Why not just to
L1	confirm that. If it's superficial, it'll be
L2	easy.
L3	MR. STIVER: Yes, I mean, it's a
L4	matter of an afternoon. One afternoon looking
L5	at it and see if there's anything on it.
L6	MEMBER LEMEN: So if you task the
L7	SC&A, does that mean should we have another
L8	Working Group meeting after that?
L9	MEMBER MUNN: It depends.
20	MR. STIVER: In any case there are TBD
21	revisions out there that we have not seen yet.
22	CHAIRMAN MELIUS: Okay. So we'll
23	task SC&A to review those. Okay.

1	MEMBER LEMEN: And then should the
2	Working Group wait until SC&A reviews it and
3	then take action after that?
4	CHAIRMAN MELIUS: Yeah, yeah, NIOSH
5	is going to have to respond to their reviews.
6	And this is down the road a bit, but that
7	MEMBER LEMEN: I just want to make
8	sure there's nothing from the Working Group
9	you're expecting.
10	MR. KATZ: Nothing yet.
11	CHAIRMAN MELIUS: Nothing yet.
12	And Worker Outreach, Josie's not
13	here, so I think we'll put that off until the
14	next meeting. And there is some follow-up that
15	we need to do in that, but Josie needs to should
16	be present for that.
17	And then we have our two
18	Subcommittees. Dose Reconstruction Reviews.
19	David?
20	MEMBER KOTELCHUCK: Okay. Well, our
21	last
22	CHAIRMAN MELIUS: Can you speak into
23	the microphone

1 MEMBER KOTELCHUCK: Our last teleconference meeting was October 29th, which 2 Wanda kindly chaired in my absence, and much was 3 We almost finished 10 through 13. 4 Actually, have 14 findings 5 we remaining in the so-called DCAS sites file. 6 We have a next meeting, as Ted noted, on December 7 And we will finish up the findings at that 8th. 8 9 time and then begin, finally, the 14th set. And that will allow us also to begin working on 10 the audit report, finally, for 10 through 13. 11 12 And work is going on on the blind cases 13 and work is continuing on NIOSH work on the 9th up through the 19th set. So that's the report 14 15 and we're moving along. 16 CHAIRMAN MELIUS: Paul. MEMBER ZIEMER: I'd like to ask my 17 18 periodic question. And that is, what are the 19 plans to report to the Secretary on the 20 scientific validity? Is that what you were 21 talking about? MEMBER KOTELCHUCK: 22 Yes, that's what 23 I was talking about, the audit.

1	MEMBER ZIEMER: Okay.
2	MEMBER KOTELCHUCK: Oh yes,
3	absolutely. We're behind.
4	MEMBER ZIEMER: Yes. That's okay.
5	MEMBER KOTELCHUCK: We had hoped to
6	begin earlier, but we will do it now and try to
7	expedite it.
8	MEMBER ZIEMER: I just wanted to make
9	sure I understood that.
10	CHAIRMAN MELIUS: And I will ask my
11	periodic question, which is what about the
12	blind reviews?
13	MEMBER KOTELCHUCK: The blind reviews,
14	we've gone over, I believe, four out of the six.
15	It's been put back on our schedule because we
16	want to get 10 through 13 completed so we can
17	do the report to the Secretary.
18	The blind reviews, as you remember
19	from earlier meetings, for the four cases that
20	we reviewed, the blind reviews were identical
21	or compatible. We haven't moved ahead on the
22	others. We will now be able to, however, when
23	we finish 13.

1 CHAIRMAN MELIUS: Okay. Any other questions, comments, anything from NIOSH or 2 3 SC&A on that? Okay. MR. KATZ: Well, just to update you, 4 5 SC&A has been assigned. They're doing six plan reviews now, additional ones. 6 And they've also been assigned their 21st set of dose 7 reconstructions, which will take them through 8 9 March. So that's also happened in this 10 interim. CHAIRMAN MELIUS: Procedures Review. 11 MEMBER MUNN: Procedures has not met 12 13 since I gave you a fairly concise report during our September Board teleconference. 14 We are scheduled for Tuesday, November 25. 15 So we will be meeting later this month. 16 17 At our last meeting, we had a number 18 of PERs that we looked at and we have a gaggle 19 more that's coming up for us. We also took a 20 look at several OTIBS that we were attempting to close out last time, some of which we did. 21 22 They included ingestion, inhalation 23 of plutonium and internal dosimetry issues,

1	including some internal doses related to gross
2	alpha and gross beta. But most of our focus,
3	I think, will probably be on PERs during this
4	upcoming Procedures meeting later this month.
5	CHAIRMAN MELIUS: Any questions or
6	comments? Yes, Paul.
7	MEMBER ZIEMER: Actually, if I might
8	be permitted to ask a question of David on the
9	previous report. We have 21 total reviews we
10	will have finished with this last group, 21
11	groups of dose reconstruction reviews. Is it
12	21?
13	MEMBER KOTELCHUCK: No, we've been
14	working on 10 through 13 sets
15	MEMBER ZIEMER: Yes, but which ones
16	which group is ready for SC&A's completing
17	or
18	MR. STIVER: Well, we have delivered
19	up through set 19 now. Set 17 were the six
20	additional blinds that Dr. Kotelchuck was
21	talking about that. And since then we've been
22	tasked to do Set 20, which are another
23	additional set of blinds.

1	MEMBER ZIEMER: Okay. But
2	MR. STIVER: And then 21 through 30
3	are standard dose reconstructions.
4	MEMBER ZIEMER: Okay. So with
5	those, how many total cases have we reviewed?
6	Is it somewhere around four to 500?
7	MEMBER KOTELCHUCK: Let's see.
8	Total? You mean from the beginning?
9	MEMBER ZIEMER: Yeah.
10	MR. STIVER: The table I put in
11	there
12	MEMBER KOTELCHUCK: Over 200. I'm
13	just looking at the
14	MR. KATZ: No, no, no. David, John
15	has the numbers. We talked about this
16	recently.
17	MR. STIVER: Yeah, in the review that
18	I sent out to you guys, there's a table on Page
19	15. And the total number of cases up through
20	Set 19 is 468. Findings are nearing completion
21	through Set 13. That still leaves 14 through
22	19 including
23	MR. KATZ: So though Set 21 it'll be

about 500. That's correct.

MEMBER ZIEMER: Okay. So about 500, what I'm trying to get at is to get an update on what percent of the total cases that have been reconstructed have we been able to review and whether our original goal of two-and-a-half percent was even realistic. I have a feeling it wasn't and I don't know if we'll be able to achieve, ever, what we thought we could ten years ago, or more than ten years ago, twelve years ago, and whether or not we should.

I don't think we need to do it today, but I'm wondering if the Dose Reconstruction Subcommittee might advise us as to what is realistic, so that we have on record --

MEMBER KOTELCHUCK: Sure.

MEMBER ZIEMER: Maybe we need a new goal or we have to do more or change the goal.

MEMBER KOTELCHUCK: Yeah. I mean, to respond just to what you're saying, I mean, I've looked at the numbers and we're really running around one percent. Maybe one-point-something, at least half of

two-and-a-half-percent. And that's 1 we've been running and it's been very slow. 2 But, I mean, well, 3 CHAIRMAN MELIUS: we've all talked about this, but it's not a 4 5 static methodology and it's very complicated. So the methods that were used for the first 6 hundred were different than the last hundred. 7 MR. KATZ: But are we doing it any 8 9 faster? I don't --10 CHAIRMAN MELIUS: No. Well, and so I quess the follow-up, and reason for my question 11 12 was about the blind reviews is, is there a 13 better method that we could be using or a different methodology we should be using that 14 more efficient? 15 miqht be Or a mix 16 methodologies and approaches that might be more 17 efficient in terms of assuring the quality of 18 that, but also identifying any remaining 19 problems. 20 I think most of our problems have been 21 -- at least my estimate is just sort of this fact 22 that you've separated the dose reconstruction

reviews from the Site Profile, SEC issues and

so forth. We go at them sort of differently. 1 And Procedures and so forth. But is there some 2 way of taking that into account? 3 Is there something else we should be able to do in 4 methodology. 5 But I think to get to that, I think, 6 when we last talked about this, which is 7 probably a couple years ago, was the issue of, 8 9 one, we needed to have a report to the Secretary 10 or something like that would summarize this, at least for the more recent reviews to be able to 11 12 evaluate it. 13 secondly, the blind reviews would also be helpful in terms of helping to 14 evaluate what other methodologies might be 15 16 Not that we could ever do, you know, two-and-a-half percent blind reviews, 17 18 might point to issues that would come up. 19 So I think we could try to aim for, you 20 know, pulling our -- so really looking at our 21 methodology again as we're doing this audit. 22 MEMBER KOTELCHUCK: Well, as both the

chair and also a new person on the Committee,

we have not had, with the blind reviews that 1 have been completed and that we've looked at, 2 there seems to be no problem, that we're doing 3 4 a good job, there is agreement. 5 And the Sets 10 through 13 have been very, very slow. And at least from my point of 6 7 view, that's most of the time I've been on the and the Subcommittee, 8 group we've 9 discussing those. 10 And so the push has really been to get those out so that we can begin to work on the 11 12 I believe we can come back onto the 13 blind review cases fairly quickly. small in number and there hasn't been a problem. 14 15 So with respect to what the Chairman is saying, we will go back to blind reviews, but 16 17 we just had to get 10 through 13 off our plate. 18 And that has been, I felt, an imperative, 19 because we could not even begin to talk about 20 the report to the Secretary, which has a high 21 priority. 22 CHAIRMAN MELIUS: I'm not being

critical of both the priorities.

23

I was just

trying to address how do we address the bigger
issue, which Paul raised also. Is there some
change? And not to reflect on the work that
you've done or the Subcommittee or the people
involved in the reviews, but it's been a long
time and we need to look at that. And we
recognize that we need to at least get that 10
to 13 audit, whatever we're calling it,
completed as a priority and then be able to move
on.
MEMBER KOTELCHUCK: I'm not feeling
that you're attacking the Committee or
anything
CHAIRMAN MELIUS: Yeah, yeah. Okay.
MEMBER KOTELCHUCK: but just
trying to say how we sort of established
priorities ongoing. And getting something to
the Secretary, as I've said, is a high priority.
CHAIRMAN MELIUS: Yeah.
MR. KATZ: And if I could just add
something related to Paul's point, to put a very
fine point on it. We're falling, despite the
fact that we've really thrown ourselves at this

harder since -- although we've had some quorum issues at times that have gotten in the way, we're actually losing ground in terms of the difference between where the Subcommittee is in reviewing cases and the SC&A's production reviews.

CHAIRMAN MELIUS: Right.

So we definitely have to MR. KATZ: have this sort of better thinking about how we go about this in a big sense, so I totally concur.

> CHAIRMAN MELIUS: Stu.

MR. HINNEFELD: I was going to offer one thing to keep in mind when you talk about the original objective of two-and-a-half percent, or whatever, is that for many years now the Subcommittee has selected cases with PoCs over 40 percent or over 45 percent, which is a very tiny minority of the total cases.

Ι bet that=s far less than two-and-a-half percent of the dose reconstructions we do. And so, you know, based on those facts, that we are only selecting this

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top tier, if you selected them all, I bet you 1 wouldn't be at two-and-a-half percent. 2 3 So there's a really fundamental 4 reason here why the Subcommittee, I think, 5 could come back and say it's not realistic, beyond the fact that it's an awful lot of work, 6 7 it's just not realistic because they're not two-and-a-half percent in the range we want to 8 9 look at. 10 CHAIRMAN MELIUS: Yeah, but another 11 take on that would be that because they're, you 12 know, over 40 percent, they probably are more 13 difficult to review. And since you were already prioritized, we're already selected to 14 15 try to address the problems, but it also makes 16 the work load bigger. 17 MR. KATZ: Right. 18 MR. HINNEFELD: Yeah. 19 CHAIRMAN MELIUS: Yeah, yeah. If it 20 was a random sample, I think we would probably 21 have a lot more done because it would be a lot 22 quicker to do. And that's not to fault the

selection criteria, but, again, I think we need

to sort of look at what comes out of the audit and what our past experience has been beside this. Are there other approaches we can use? Yes, David.

MEMBER RICHARDSON: Well, I mean, I=m going to play the devil's advocate. I mean, I can't go back 12 years, but I can go back a number of years when we talked about that logic of sampling. And we recognized we were doing stratified sampling, over-sampling certain types of cases, but nonetheless we had our sites set at at least sampling a couple percent of the cases in order to get some coherent picture of the information. And we're well below that target.

I mean, so that does force us to kind of reflect about how we can characterize the whole process based on, let's say, one-half of a percent or a one percent sample.

The second part was some of the problems we find would be, and these are more difficult cases in a sense, but some of the problems we find seem to be still sporadic,

episodic, quality control, sort of, you know, 1 quality assurance issues that I'm not sure, you 2 know, would be uninformative if we would sample 3 other parts. And we just don't know right now. 4 So there seem to be some of those QA/QC issues. 5 And the third one is we had set our 6 7 sites at this couple percent sampling issue, but since the ten year review, NIOSH has taken 8 9 on sampling some cases as well. And I don't 10 remember what your goal was or how many blind reviews or basically reconstructions NIOSH was 11 12 going to run through, but in a sense there's a 13 shadow program running and also doing this. I've forgotten now, 14 MR. HINNEFELD: 15 but it's maybe one percent. It's not a very big 16 percent. 17 MEMBER RICHARDSON: But it's in some 18 sense we have now, we have two one percent 19 programs going on and that may help us. 20 CHAIRMAN MELIUS: And it might be 21 helpful, when we're ready, is to get a report on the NIOSH review process also when we're 22

ready to be talking about the audit and what

1	should we be doing in terms of methodology. So
2	you could be able to summarize, you know, your
3	experience, Stu, with doing these reviews and
4	so forth. Okay.
5	MEMBER KOTELCHUCK: I mean, I'm
6	counting on the review for the Secretary as
7	informing us. I mean, I'm looking forward to
8	that, to learn, not just that I have to report
9	to somebody else above me but that it will give
10	us a picture, finally, of where things are
11	slowing down or can be changed to speed things
12	up.
13	CHAIRMAN MELIUS: Yes. Okay. Any
14	other comments on that issue? Okay. Do we
15	have any other Board
16	MEMBER KOTELCHUCK: Yes, two more.
17	Do you want something on Ames?
18	MR. KATZ: Ames.
19	CHAIRMAN MELIUS: Ames?
20	MR. KATZ: You skipped it.
21	CHAIRMAN MELIUS: Did I skip Ames?
22	MEMBER KOTELCHUCK: We have an Ames
23	Working Group. You skipped us.

1	MR. KATZ: I was going to catch you.
2	It's all right. It wasn't on the website, I,
3	guess.
4	MEMBER KOTELCHUCK: Right. I'll be
5	glad to report it.
6	CHAIRMAN MELIUS: Ames is missing
7	from the website. No respect. Unless it's
8	been renamed.
9	MEMBER KOTELCHUCK: Right. Well, it
10	has not even had its first meeting yet. But it
11	is getting organized and
12	CHAIRMAN MELIUS: Well, that's why
13	the website hasn't been alerted yet.
14	MEMBER KOTELCHUCK: Right. For the
15	record, then, let me say that the Ames Work
16	Group is Dr. Roessler, Loretta Valerio, Brad
17	Clawson and myself.
18	I've been in touch with Tom Tomes. He
19	has sent us background files which many folks
20	have been looking at and I've been in
21	discussions with him trying to basically get a
22	better orientation as to quite where we are.
23	He envisions that we need four more

1	White Papers, which he is talking about doing
2	next summer, of 2015. We have not met as a
3	group and we will be having an organizing
4	conference call sometime either in December or
5	January. That's when I'd like to have it.
6	And then we'll proceed in discussions
7	with Tom. I'll share the discussions that I
8	had with him with the rest of the Working Group
9	and we'll see.
10	Obviously, if it will take him until
11	next summer, then we're not going to be able to
12	do very much between now and then, regrettably.
13	CHAIRMAN MELIUS: My recollection is
14	that there was a Site Profile review from SC&A.
15	And so are these White Papers in response to
16	that? That's what I'm trying to
17	MEMBER KOTELCHUCK: Yes.
18	MR. STIVER: Yeah, I was going to say
19	that we turned in or delivered our review in
20	August of 2013, our 22 findings on that. So I'm
21	assuming is what is our response. This is
22	non-SEC.
23	CHAIRMAN MELIUS: Yes, yes. They're

1	Site Profile issues that
2	MR. HINNEFELD: These are for things
3	that are for this is in SEC for much of its
4	period and so many of these issues are related
5	to the non-SEC cancer claim dose
6	reconstructions.
7	CHAIRMAN MELIUS: And one of the
8	reasons we held up to do the Site Profile review
9	was so NIOSH could then focus some resources on
10	being able to address these Site Profile
11	issues. So that I think the schedule makes
12	sense in those terms, so. I just wanted to get
13	that on the record.
14	MR. KATZ: Okay.
15	CHAIRMAN MELIUS: Did I miss anybody
16	else? Did the website have any other failures?
17	MR. KATZ: No, that's good.
18	CHAIRMAN MELIUS: Yes.
19	MR. KATZ: Blockson, there's no
20	activity right now.
21	CHAIRMAN MELIUS: Yeah. I'm not
22	sure if Blockson needs activity does it?
23	MR. KATZ: There'll probably be a PER

1	at some point, which is why I'm not making it
2	inactive.
3	CHAIRMAN MELIUS: Okay. We'll wait.
4	Good. Any other things Board needs to do?
5	MR. KATZ: No, I think that does it.
6	CHAIRMAN MELIUS: Okay. Then we
7	will break until 4 o'clock.
8	I'm reluctant to try to move up the
9	Santa Susana because we have people scheduled
10	to come in and I don't want to though some
11	people are here, I'm concerned that other
12	people might come in around 4 o'clock given
13	MEMBER SCHOFIELD: And some are going
14	to call in later.
15	CHAIRMAN MELIUS: Yes. We have
16	people later, so, yes. Because I think our
17	presentations, I think, will go on a little
18	longer, at least a half-hour, but just a guess.
19	So we will start back up here promptly at 4
20	o'clock.
21	(Whereupon, the above-entitled
22	matter went off the record at 3:00 p.m. and
23	resumed at 4:02 p.m.)

1 CHAIRMAN MELIUS: Okay. We'll reconvene now. And my schedule here. 2 Phil, do you have anything you want 3 to say to start or should we just go into -- I 4 could use his --5 MEMBER SCHOFIELD: I think we should 6 7 just go ahead and get started. CHAIRMAN MELIUS: Okav. 8 9 MEMBER SCHOFIELD: The only think I 10 do have is just one brief thing. I really 11 appreciate DOE and all the people at the 12 facility yesterday for arranging the tour. 13 think that was very educational for us. CHAIRMAN MELIUS: So what we'll do is 14 15 we'll start. Dr. Hughes will do a presentation Then we'll hear from SC&A, John 16 from NIOSH. Stiver, we'll from 17 and then hear the 18 petitioner. 19 And at some points there may be after 20 presentations, some questions from the Board 21 Members, but then we will, after that, after hearing from the petitioner, we will go into the 22

public comment period. So let's do that.

1	we'll start with you, Lara. Welcome. Haven't
2	seen you for a while, so
3	DR. HUGHES: Yes.
4	CHAIRMAN MELIUS: good to see you.
5	DR. HUGHES: Thank you, Dr. Melius.
6	CHAIRMAN MELIUS: Yes.
7	DR. HUGHES: This is the Santa Susana
8	Special Exposure Cohort Site Profile update.
9	Okay. When we say Santa Susana in the context
10	of this program, we really talk about four
11	separate covered sites.
12	We kind of treat them as one thing
12	We kind of treat them as one thing because all the sites share the same operator
13	because all the sites share the same operator
13 14	because all the sites share the same operator over the course of time and also share the
13 14 15	because all the sites share the same operator over the course of time and also share the workforce and most importantly for our program,
13 14 15 16	because all the sites share the same operator over the course of time and also share the workforce and most importantly for our program, their dosimetry program, the issues affecting
13 14 15 16 17	because all the sites share the same operator over the course of time and also share the workforce and most importantly for our program, their dosimetry program, the issues affecting the dosimetry program are shared between all
13 14 15 16 17 18	because all the sites share the same operator over the course of time and also share the workforce and most importantly for our program, their dosimetry program, the issues affecting the dosimetry program are shared between all these sites.
13 14 15 16 17 18 19	because all the sites share the same operator over the course of time and also share the workforce and most importantly for our program, their dosimetry program, the issues affecting the dosimetry program are shared between all these sites. The first site, the largest, is Area

from 1955 through 1964, the Canoga Avenue

Facility which is covered 1955 through '60, the 1 entire period as an SEC Class. 2 The DeSoto Avenue facility has a 3 covered period of 1959 through 1995 as well as 4 5 1998. And it is currently in SEC Class from 1959 through 1964. 6 And the Downey Facility has a covered 7 period of 1948 through 1955 and the entire 8 9 covered period is an SEC Class currently. 10 Now, those of us who went on the tour 11 yesterday got a nice detailed history of the 12 The contractor history is somewhat 13 complicated and I don't want to go into it. All of these sites were non-weapons 14 facilities that did research into nuclear 15 reactors and other nuclear materials. 16 So they 17 were reactor operations. 18 There were about ten different. 19 reactors that were built and tested as well as a number of critical test facilities which is 20 21 kind of, it was a small reactor. And there was a number of nuclear 22 23 support operations. There was a hot lab facility. There was a small accelerator facility. There was radioactive material handling facility. There were waste handling facilities and so forth.

The Work Group for Santa Susana was

The Work Group for Santa Susana was established in 2008 in response to the Board contractor doing a review of the NIOSH TBD on the sites. That was done in 2008. There were two SEC Evaluation Reports for Area IV that were delivered to the Board in 2009 and 2010.

The first SEC Evaluation Report on Area IV was also reviewed by the Board contractor and the TBD issues and the SEC issues resulting from those reviews were the discussion points in front of the Work Group.

The Work Group has met in 2008 and 2009 and 2010. And just recently, in 2014, the very last meeting last month was mainly to touch base and kind of reestablish the Work Group.

So the issues that were discussed in the past in front of the Work Group included things like the site definition and operations timeline of all four sites, incidents, internal

monitoring issues, internal coworker model feasibility and necessity.

I forgot to mention with all of the SECs that were established for these sites, were based on defining of internal infeasibility. So the internal data was a big discussion issue.

There was also issues with the external monitoring data, the neutron data, the environmental approach that is outlined in the TBD as well as how we deal with the tritium plumes that are on site and potential work exposures.

Currently, the Work Group is discussing the neutron/photon ratio White Paper that was sent to the Work Group in 2010, but that was kind of on hold pending some other issues.

What needs to be discussed is the internal coworker model that has been completed by NIOSH in March of 2014. And since I prepared this presentation, the Board contractor has also issued their review of the neutron/photon

ratio White Paper. So that's also in front of the Work Group to be discussed soon.

There are several NIOSH draft documents that are ready to release. The external coworker model is in a pending status pending the resolution of issues regarding neutrons. And there's several TBD revisions that are pending resolution of issues and they will be released as soon as those are resolved.

So since 2010, NIOSH has done considerable work on the site, although the Work Group has not met. In the last Work Group meeting in 2010 there was discussion on the internal and external coworker model. So ever since then NIOSH has continued the discussion and issue resolution affecting the internal data.

Back in 2010 we have been working with a database that was received from Santa Susana from Boeing that we attempted to work into an internal coworker model, however, there were numerous problems.

The reason being that this database

not developed for the purposes NIOSH 1 needed. Ιt developed 2 was for an epidemiological study, so there was some issues 3 with that. 4 5 So NIOSH has been negotiating with Boeing trying to obtain scanned 6 worker 7 dosimetry records so we could do the data entry ourselves. 8 9 Also in 2010, NIOSH prepared the 10 neutron/photon White Paper and revised the internal/external and environmental 11 TBDs 12 mainly to include the SEC language. 13 In 2011, the negotiations with Boeing about the worker records continued. 14 And we also, at the same time, tried to resolve issues 15 with the, what we call the Boice database 16 17 because it developed was for the 18 epidemiological study by Dr. Boice. 19 We found some issues that we decided 20 that we cannot use it for NIOSH purposes of developing internal coworker model. 21 And also in 2011 another iteration of 22

TBD revisions were completed, but they remain

pending resolution of issues with the coworker 1 studies. 2 finally Boeing decided 2012, 3 release the scanned worker records to NIOSH. 4 5 Those were received in March of 2012 and it consisted of 14,000 files that had to be sifted 6 through and classified. 7 So the data entry from those lasted 8 9 from May 2012 to January 2013. And the internal and external coworker models as a 10 result of those records were completed and 11 12 reviewed in 2013. 13 The external model is currently pending and waiting approval once we decided 14 15 how to approach the issues on neutrons which is something we've been working on in the past few 16 17 months. 18 So this year the internal coworker 19 model was approved and released in March. once we decided that the internal model was 20 21 feasible, we picked up the issue resolution on 22 the external because that was kind of on a

holding pattern pending to see if the internal

was going to happen.

So the internal coworker model will be known as OTIB-80. It's a plutonium model, uranium model and a gross beta model starting in 1965 which is the end of the SEC Class up until the modern day period.

The external coworker model will be known as OTIB-77, currently in draft status. It is a result of an analysis of about 175,000 data points and it will consist of a site wide model addressing photon, electron and neutron doses. It will also have a separate model for Area IV and a separate model for the DeSoto facility.

The neutron/photon ratio White Paper was developed to provide a bounding approach for unmonitored neutrons for the time span of 1956 to 1987.

To do this, we analyzed over 1,000 paired neutron and photon measurements around the reactor facilities. There was a lognormal fit data involved which resulted in a neutron/photon ratio of 1.73.

The neutron approach that's to be done with the neutron/photon ratio is that it's to be used for a worker that was either employed site wide or had an unknown work location.

We've also found that the accelerator, there was a small accelerator facility operated at site only for a short period of time and in cases where a worker could be placed at this facility, the NTA film with a correction factor would actually be useable. But it's not anticipated to be a large part of the neutron approach.

As indicated in the NIOSH White Paper, the reactor exposure is to use an N/P ratio and some additional research has indicated that for a situation such as fuel handling or other nuclear procedures that were done at the site that the reactor N/P ratio is most likely bounding.

So our current path forward is to issue the external coworker model to incorporate the neutron guidance and revise TBDs to continue issue resolution with SC&A

1	regarding the coworker models and the neutron
2	approach and address remaining SEC issues and
3	remaining Site Profile issues which regarding
4	to the SEC I think that was mentioned earlier
5	in the SEC updates.
6	There's a question of the year 1965
7	because that was a year that initially
8	qualified for evaluation, however, the Class
9	was only recommended through 1964.
10	So we still owe the Board some kind of
11	decision on what's going to happen with 1965.
12	And with that my presentation is complete. So
13	if you have any questions?
14	CHAIRMAN MELIUS: Okay. Questions?
15	Yes, Gen?
16	MEMBER ROESSLER: You mentioned not
17	being able to use the Boice database and I can
18	understand why NIOSH data that's put together
19	for dose reconstruction cannot be used for
20	epidemiology. But remind me why it can't be
21	done the other way.
22	DR. HUGHES: Well, we received the
23	Boice database from Boeing and as far as I'm

remembering correctly, we were assured that 1 this database was a complete database of all 2 workers, of all internal dosimetry data that 3 4 there was. 5 However, we also have in our database from the 1960s what's called annual exposure 6 7 It was reports written by the site reports. that indicated how many workers were monitored, 8 9 how many urine bioassays they did, how many were 10 positive. And we actually did some kind of 11 12 quality assurance and we found some 13 discrepancies that we were not able to resolve. So at this point we were saying that we do not 14 15 have enough confidence that the Boice database is actually complete. 16 17 Now, I'm not in the position to judge 18 the epidemiological study that Dr. Boice did, 19 it's just for our purposes, we found some issues 20 that we just said, well, we cannot use this 21 because we're not confident that it's complete. 22 CHAIRMAN MELIUS: Okay.

Other -- yes. Henry, you had --

1	MEMBER ANDERSON: No, no.
2	CHAIRMAN MELIUS: Oh, okay. Other
3	questions?
4	MEMBER RICHARDSON: And now that you,
5	if I understand correctly, you received a file
6	which you described as scanned records. And
7	you contracted ORAU to keypunch those records?
8	DR. HUGHES: Well, they were entered
9	into a database format. Each worker at the
10	site had a paper folder that contained the
11	dosimetry records, the batch readings, the
12	urine bioassay reports from the labs that did
13	the analyses.
14	And during the course of the Boice
15	study, actually, this was all scanned and
16	digitized and it was available in electronic
17	format. So what we got was the scanned raw
18	records of each single worker and that's what
19	we used to extract the internal/external data.
20	It's very much the same thing that we
21	receive for an individual worker during dose
22	reconstruction from the site, except that we

receive the entirety of the monitored workforce

1	at Santa Susana.
2	MEMBER RICHARDSON: And when you
3	described it as 14,000 files, is it one file per
4	worker or what makes it 14,000
5	DR. HUGHES: Yes.
6	MEMBER RICHARDSON: files?
7	DR. HUGHES: Yes. Now, not all of
8	these workers were actually monitored, but
9	there were that many files. So we had to pull
10	out. Some workers were not actually
11	monitored, but they might have been employed.
12	I'm not exactly sure. There are that many
13	files and each represents a worker, but not all
14	of these did actually contain monitoring
15	records.
16	MEMBER RICHARDSON: Some files were
17	empty?
18	DR. HUGHES: That's correct.
19	MEMBER RICHARDSON: So the
20	contention is that the workforce of radiation
21	monitored workers at the site is 14,000 people?
22	DR. HUGHES: No, it's more on the
23	order of 5,000, I believe. I don't want to say

1	anything wrong. I would have to check my
2	numbers.
3	MEMBER RICHARDSON: And then when you
4	compared that to the annual reports summarizing
5	the number of people who are bioassayed and were
6	radiation monitored in this case now, with the
7	ORAU data, does it correspond to the
8	expectation?
9	DR. HUGHES: I actually have not done
10	that comparison. However, we do know this is
11	all the site has.
12	MEMBER RICHARDSON: But it's, I mean,
13	I
14	CHAIRMAN MELIUS: But has it
15	completed?
16	DR. HUGHES: Yes.
17	MEMBER RICHARDSON: Because
18	basically you went back to the scanned files
19	that had been used for the epidemiological
20	analysis which
21	DR. HUGHES: Yes.
22	MEMBER RICHARDSON: your
23	records

1	DR. HUGHES: Correct.
2	MEMBER RICHARDSON: to be
3	incomplete and you reentered them and I guess,
4	it leaves the question now, are they complete?
5	DR. HUGHES: We hope so, yes. Well,
6	we actually have not gone back and done the
7	comparison. However, what we've done, we've
8	done a comparison with the NOCTS claim files
9	that we received and have
10	MEMBER RICHARDSON: I mean, it's
11	been I mean if some were even
12	DR. HUGHES: Which is a subset
13	MEMBER RICHARDSON: passed. One
14	is
15	DR. HUGHES: of it, but there was
16	no discrepancy with that, so.
17	MEMBER RICHARDSON: Yes. I mean,
18	one of them is, it's interesting to know whether
19	the effort paid off. Another one that=s
20	DR. HUGHES: Yes.
21	MEMBER RICHARDSON: just
22	interesting. I mean, there have been examples
23	in the past where we thought we had complete

1	data and then we realized that there were gaps
2	and we said there appear to be gaps. And then
3	further effort went into locating files and
4	there was more, so. I would just, it might be
5	worth
6	DR. HUGHES: Okay. Yes, the
7	MEMBER RICHARDSON: Wherever
8	possible trying to reconcile the
9	DR. HUGHES: Yes, that can certainly
10	be done.
11	CHAIRMAN MELIUS: Okay. Any other
12	questions?
13	MEMBER RICHARDSON: I had a
14	CHAIRMAN MELIUS: Oh, go ahead.
15	MEMBER RICHARDSON: another
16	question. Because the external file had
17	175,000 data points and the neutron to photon
18	ratio was derived from 1,180 paired
19	measurements. So is my understanding that of
20	the 175,000
21	DR. HUGHES: No, sorry. That's two
22	completely different things. The 175,000 is
23	

1	of thing. The 1,100 neutron/photon, this is
2	survey data, hand held survey meter data
3	MEMBER RICHARDSON: Oh.
4	DR. HUGHES: that was
5	MEMBER RICHARDSON: Oh
6	DR. HUGHES: collected
7	MEMBER RICHARDSON: these are
8	DR. HUGHES: around the reactors.
9	MEMBER RICHARDSON: area
10	monitors. Oh
11	DR. HUGHES: Yes.
12	MEMBER RICHARDSON: then those
13	aren't monitors. Okay. I got you.
14	DR. HUGHES: Yes. It's not
15	personnel dosimetry, it's area data.
16	MEMBER RICHARDSON: Okay. Now,
17	within the file, the monitoring file there's
18	for a subject in a badging period, they have
19	potentially estimates of their photon dose and
20	their neutron dose as well. Is that right?
21	DR. HUGHES: That's correct. If
22	they were wearing the NTA film badge and if it
23	had a reading, it would be reported in their

1	file. That is correct.
2	MEMBER RICHARDSON: Okay.
3	CHAIRMAN MELIUS: You set? Okay. I
4	couldn't tell if you were writing something
5	down earlier or had another question. That's
6	why I interrupted anyway. Any other Board
7	Members with questioning? Board Members on
8	the phone?
9	If not, I have one comment. First of
10	all, thank you for a very good succinct summary
11	of a long period of time. So it was helpful.
12	One thing I'd say, since, as you know,
13	we're as a Board and NIOSH are dealing with the
14	evaluation of coworker models, and I think it
15	would be helpful as the Work Group and SC&A and
16	NIOSH address these coworker models here, at
17	least keep in mind the kind of implementation
18	guidelines we have.
19	So I just don't want to have to
20	backtrack on this. So, again, I don't want to
21	sort of hold you to criteria that you haven't
22	seen yet.

DR. HUGHES: I have been warned, so.

1	CHAIRMAN MELIUS: Okay. Okay.
2	Okay. Well, not as much a warning as a request.
3	DR. HUGHES: Yes.
4	CHAIRMAN MELIUS: Do the best that
5	you can without having to I don't think it
6	affects it, it's as much as I think what you
7	present going forward and so forth. So thank
8	you. Okay. Now, we'll hear from John Stiver
9	from SC&A.
10	MR. STIVER: Thanks. Good
11	afternoon, everybody. I'm John Stiver from
12	SC&A and I'm going to be giving our and the
13	Board's perspective on the various activities
14	that have taken place since the initiation of
15	the TBDs and the reviews and the SEC petitions
16	and so forth since 2006.
17	The first half of the slides really
18	deal with kind of a timeline of the activities
19	that have happened. It's a very convoluted set
20	of findings and activities, as anybody who's
21	tried to wade through the transcripts from the
22	meetings would attest to.

And then after that, I'm going to go

ahead and just kind of give you an idea and let 1 you know where we are and how we got there. 2 Kind of give you a thumbnail sketch of what we 3 really believe the issues to be and then where 4 5 we're headed from there. Lara had given you some of this 6 2006 and 2007, the Technical 7 information. Basis Documents were first issued. In January 8 9 through April of 2006 there were a series of five worker outreach meetings. 10 Those have been incorporated into our Site Profile review. 11 12 And then in June of 2007, SEC Petition 93, which was a 83.13, I believe the initial 13 period was from 1955 through the present, 14 15 basically, including the post-1987 remediation 16 period. 17 The report was sent to the advisory 18 Board in February of 2008, discussed by the 19 Board in April and June of 2008 and then in August of 2008, we submitted our review of the 20 21 Santa Susana Site Profile.

that the Site Profile review was conducted in

And, so, this is somewhat unique in

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the midst of ongoing SEC deliberations in the 1 petition process. 2 Our review uncovered a total of 39 3 4 findings. You can see how they're parsed out 5 in the slides based on the different TBDs with most of the findings associated with the 6 internal dose TBD, Number 5. 7 On August 26th of 2008, ten days after 8 we released our review we had the first Santa 9 10 Susana Work Group meeting. And it's important to note that all the Site Profile findings were 11 12 discussed, but none were officially closed. 13 Obviously, within a ten day period, the petitioners hadn't had time to review and 14 comment on our review. There was the issue of 15 ongoing SEC deliberations. 16 17 And so the findings really discussed 18 in terms, the potential SEC impact. And those that were determined to be Site Profile issues 19 20 at that meeting were pended until the SEC issues 21 could be resolved. Excuse me, I've got a 22 little problem with my voice here.

The Work Group agreed that SC&A should

go ahead and combine some of the closely related 1 findings. This was kind of situation where 2 NIOSH was getting hit from several different 3 4 angles on one given issue. And so we went ahead 5 and prepared a new SEC issues matrix and condensed everything down into a total of 13 6 7 findings. really mixture 8 These are а 9 petitioner concerns, issues that were 10 identified by NIOSH and then some of the combined SC&A findings. 11 12 And on April 17th of 2009, we had the 13 second Work Group meeting. Oh by the way, this is the issues matrix, it's the 13 findings are 14 the ones that are still in the issues matrix 15 that we're working from today. 16 NIOSH was tasked with several action 17 18 items for the next meeting which was held in 19 April of 2010. And during that intervening 20 period there was a great deal of SEC activity 21 that took place.

Class be added to the SEC based on Petition 93

Basically, the Board recommended a

22

which was from January of '55 through December 1 of '58. 2 then, in November of 2009, 3 And Petition 156, which was an 83.14, which was from 4 5 NIOSH was kind of fast-tracked. It was, obviously, qualified very quickly. 6 January 7 15th, 2010, sent to the advisory Board. It was discussed at the February 9th, 2010, Board 8 9 meeting that I believe was here at Manhattan 10 Beach. then in March 11 And the Board 12 recommended another second class be added to 13 the SEC which was from January 1st, '59 through December 31st, 1964. 14 15 So the combined SEC's basically cover the period of January 1st, '55 through December 16 17 31st of 1964. And, obviously, the outstanding 18 SEC issues may impact that 1964 end date. 19 2010, April а third Work Group 20 meeting, the 13 issues were discussed 21 context of NIOSH's actions from 2009. Also we presented review of the Rev 0, OTIB 77, which 22 23 was the external dose coworker data set and coworker model.

We found five main issues all related to the appropriateness of the Boice database for coworker modeling. NIOSH had some tasking, obviously, to complete the external and internal dose coworker models and to provide updates to the environmental TBDs. And we were to review those products as they were made available.

And in June of 2014, as Lara's mentioned, NIOSH released their White Paper on neutron dosimetry. We began reviewing that and in October of 2010 we were pretty close to completion.

However, that was never finished up because of competing priorities regarding some of the other sites that we're dealing with.

And let me get to the next -- here we go. Hang on, I think we missed one.

Okay. Basically, although NIOSH was busy, you know, collecting data and putting together these coworker models, there was really no SC&A or Work Group activity since 2010

until 2014.

And in March, NIOSH released OTIB 80 which was the internal dosimetry data set. And this used the Boeing database and abandoned the Boice data set. We commenced our review in July of this year and we're not getting to the point where we're close to completing that review.

There were some brief discussions at the April and July 2014 Board meetings and then the fourth Work Group meeting was a teleconference meeting on October 16th, 2014, in which we kind of tried to jump start the SEC finding resolution process, get reacquainted with everything and NIOSH provided some of their updates on the forthcoming work products, OTIB 77 and the environmental TBD revisions.

Where do we stand, the issue matrix, the September 24th update of that that was used at meetings posted on the DCAS website at the URL that's listed here.

Only one finding was closed. This was Number 9, which is the question of which

areas, whether it was going to be Area IV, Canoga Park, DeSoto and Downey, how we should be considered in the SEC and what were the start dates for the SEC.

And our research, Boeing is up here 2005, made it pretty clear that before 1955 there really were no radiological activities taking place. It was mostly design and construction.

And, obviously, the petition was for Santa Susana, Area IV, so the Work Group felt that we could go ahead and close this one out.

There's one open combined finding that depends on the internal coworker model. This is Number 10. The adequacy of the internal monitoring program really subsumes five sub-issues related to the completeness of the bioassay data set and how well it correlates to specific radionuclides.

The issue of missing radionuclides, those are really the big ones, and then the fact that there was really no internal coworker model. You've got to keep in mind that some of

these findings really date back from 2009, so. 1 There were four open findings that 2 depended on the review of external coworker 3 This is Number 1 and the sodium models. 4 reactor experience incident in 1959. This 5 issue of maybe a technical shortfall in the 6 external radiation monitoring badges. 7 Number 7, identification of workers 8 9 with blank radiation exposure records. This 10 is a NIOSH generated issue. Those previous 2 11 were petitioner issues. 12 Number 13, this was a combined SC&A 13 finding and this is all related to the external dose coworker model. Mainly, it subsumes 14 three areas of concern, one that there was no 15 coworker model developed at this point. 16 There was the issue of low energy in 17 18 thermal neutrons and that was really below 19 about the 500 keV cut off the NTA film. 20 questions regarding there were some the 21 dosimetry response to low energy photons. 22 There are four findings that are open

that depend on revisions to both the internal

and the external coworker models. Now, this is 1 the petitioner issue of uranium fires and how 2 those are going to be treated. 3 The sodium burn pit, this is another 4 5 petitioner issue, whether the facility was adequately monitored and the records are 6 7 missing. Number 8 was a NIOSH issue. This was 8 9 about monitoring of firemen who might have been involved with fires or been around the sodium 10 burn pit when activities were going on there. 11 12 then Number 11 is kind 13 broad-based finding, another combined SC&A finding about incidents in 14 general. So 15 there's going to be some overlap here with the sodium reactor experiment and the sodium burn 16 17 pit. 18 Three findings dependent the 19 environmental TBDs. This is the issue Number 20 3 of tritium plumes. The workers may have been 21 exposed to contaminated drinking water on site. 22 The petitioner issue of insufficient 23 air monitoring, and another combined SC&A

finding, Number 12, lack of information on environmental exposures in general.

This just kind of subsumes two issues,

one being the back extrapolation of stack emission data for the years 1971 to 1999 when the measurements were taken to earlier periods from, in this case, would be '64 through 1970.

And then also the contaminated ground water impacting onsite drinking water, which is kind of overlapping a bit with petitioner issue Number 3.

Where do we stand now? Our review of the June 2010 White Paper on neutron/photon ratios was submitted on October 23rd. So I think NIOSH has had some time to start looking at that.

Now, we're just about done with our review of TIB-80. We'll have that probably in DOE review within another week or two. We're still awaiting the release of TIB-77 and the environmental TBDs. And so, obviously, the open SEC issues are contingent upon complete reviews of all three of these documents.

1	Our next Work Group meeting is going
2	to be happening, I believe, back to back with
3	the Kansas City plant meeting in January of
4	2015. And that really completes my update.
5	If there are questions, I'll entertain those.
6	CHAIRMAN MELIUS: Board Member
7	questions? Oh, Paul.
8	MEMBER ZIEMER: Sorry. John, has
9	SC&A, to your knowledge, examined any data that
10	tells us where the tritium plume is located and
11	where it's migrated to or has NIOSH?
12	MR. STIVER: That's pretty well
13	MEMBER ZIEMER: Well, actually
14	MR. STIVER: established.
15	MEMBER ZIEMER: it was in the
16	report, but I don't recall seeing it.
17	MR. STIVER: Yes, it's in the TBDs.
18	It's been pretty well-established where the
19	pluming has it's really a matter of were
20	workers monitored for that and who
21	MEMBER ZIEMER: Well, you
22	mentioned
23	MR. STIVER: might expect

1	MEMBER ZIEMER: the drinking
2	water. I'm really asking what evidence there
3	is that it may or may not have?
4	MR. STIVER: Well, we were
5	MEMBER ZIEMER: Where does the
6	drinking come from on the
7	MR. STIVER: Well, there's
8	MEMBER ZIEMER: site and where's
9	the plume?
10	MR. STIVER: To my understanding,
11	there was some of the monitoring wells showed
12	contamination.
13	The ones that were kind of downgraded
14	from the reactors or, basically, a neutron
15	activation with water would produce the tritium
16	and then that would, you know, flow down grade.
17	However, the drinking water wells, to
18	my knowledge, never really showed any
19	contamination. NIOSH, and you guys can
20	correct me if I'm wrong, but I believe this was
21	sort of a kind of a high side approximation to
22	account for the possibility that there was some
23	intermixing with the aquifers that could have

contaminated the drinking water wells. 1 And so there was a great deal of 2 discussion about this, I believe, in 2009 in a 3 4 Work Group meeting. And we were kind of in 5 agreement with them at the time that the well, I think it was Number 34, was a pretty solid 6 estimate to be used in this. 7 But, you know, once again, until we 8 see the revised TBDs, we really can't comment 9 10 on the adequacy of that. 11 CHAIRMAN MELIUS: Thanks. Any other 12 Board Member questions? David. 13 MEMBER RICHARDSON: I just had a few clarifying questions. On one of your slides 14 you said one of the open issues that was related 15 to internal coworker models was insufficient 16 17 correlation of bioassay data to specific 18 radionuclides. What did that mean? MR. STIVER: Well, and I think that 19 20 was mainly regarding the gross beta and how that 21 would then be correlated to the fission 22 products. This is an issue, I believe, that

was raised back in the Site Profile review.

1	But there was that and I think the
2	bigger issue there was the missing
3	radionuclides. You know, the new model
4	basically looks at plutonium, uranium and then,
5	you know, basically mixed fission products.
6	And so that still leaves some others that really
7	are not accounted for
8	MEMBER RICHARDSON: Okay.
9	MR. STIVER: so.
10	MEMBER RICHARDSON: And then on a
11	subsequent slide, there were four open findings
12	about the external coworker model and one of
13	them was a Tiger Team report indicating
14	inadequate radiation badges. What do you mean
15	by inadequate?
16	MR. STIVER: I think this was related
17	to, I think, it was post-1987 when there was
18	DOELAP accreditation for the film badge
19	dosimetry programs.
20	And I believe this is a petitioner
21	issue that they were concerned that that might
22	have rendered some of the dosimetry
23	questionable.

1	And we were kind of under the
2	impression at the time that we discussed this
3	that at least, you know, during the previous
4	period that probably wouldn't be an issue
5	really because, you know, it was really more
6	about DOELAP accreditation as opposed to
7	whether the badges had some sort of a
8	technological shortfall.
9	MEMBER RICHARDSON: Okay.
10	MR. STIVER: But once again, until
11	the
12	MEMBER RICHARDSON: So this
13	MR. STIVER: TBD's released, we
14	can't really close this out.
15	MEMBER RICHARDSON: Okay. So it
16	wasn't a judgment about adequacy or inadequacy
17	of coverage of the badging program
18	MR. STIVER: Right.
19	MEMBER RICHARDSON: but it was a
20	question about the performance of the badges.
21	MR. STIVER: It was more of a
22	performance issue, you know, with the badges.
23	MEMBER RICHARDSON: Okay.

1	CHAIRMAN MELIUS: Okay. Any other
2	Board Members with questions? I have one.
3	I'm not sure if it's for John or for NIOSH, but
4	I just want to make sure I understand the issue
5	of the 1965 SEC, sort of how that fits into this
6	schedule. It's sort of in the background here.
7	MR. RUTHERFORD: Yes.
8	CHAIRMAN MELIUS: I can't quite tell.
9	MR. RUTHERFORD: Okay. Well, I
10	wanted to, actually, make a clarification
11	anyway. John said that the qualified period
12	actually extended much farther than that. The
13	actual qualified period for the petition ended
14	in 1965. And so, that's why the remaining year
15	that's open is 1965.
16	CHAIRMAN MELIUS: Yes.
17	MR. RUTHERFORD: Okay. And right
18	now, we don't have an SEC issue that would move
19	that forward to extend that Class or we haven't
20	identified one as of yet.
21	And, so, our position was at the time
22	when we added the Class was up through 1964, we
23	made the recommendation to add the Class, but

1	we'd seen nothing at that time beyond 1964.
	we a seen nothing at that time beyond 1904.
2	CHAIRMAN MELIUS: It's sort of
3	bureaucratic, but did we actually close out for
4	'65 and
5	MR. RUTHERFORD: No, that's, again,
6	another one of the one's that
7	CHAIRMAN MELIUS: Okay.
8	MR. RUTHERFORD: Yes.
9	CHAIRMAN MELIUS: Okay.
10	MR. RUTHERFORD: All right.
11	CHAIRMAN MELIUS: Okay. But we're
12	expecting that the work that's ongoing
13	MR. STIVER: Yes.
14	CHAIRMAN MELIUS: with the
15	coworker models, I suspect would address that
16	'65 and I think also if we found issues that
17	would extend beyond '65 as part of the Site
18	Profile review, there are other ways of
19	addressing that through the SEC process, 83.14s
20	and so forth. So it wouldn't be ignored, but
21	we do have to address '65 at some point. Okay.
22	Good. Any other questions? Yes, David.
23	Yes.

Can I? And this 1 MEMBER RICHARDSON: is, again, just to help me wrap my head around 2 the -- there were roughly 14,000 electronic 3 images of paper files with radiation dosimetry 4 5 data provided and that would be either external or internal dosimetry information. 6 Of those there were five or 6,000 of 7 them which were not blank folders, if I'm 8 9 understanding this correctly. And this pertains to the radiation 10 dosimetry information for workers at all four 11 12 sites of which the Boice report says there's 13 maybe five or 6,000 people radiation monitored and maybe 42,000 people also employed at those 14 four sites who are not radiation monitored? 15 16 DR. HUGHES: I'm sorry. I'm having 17 trouble following. Yes, 14,000 files is what 18 we received, each file representing a worker. 19 Not every worker was monitored at all. 20 Some only monitored for were 21 external, and I think the 5,000 might be the internal number. 22 Some were monitored for

23

external and internal --

1	MEMBER RICHARDSON: Yes.
2	DR. HUGHES: exposure. So
3	depending on the worker, there were a lot of
4	workers where you just would have a card with
5	somebody's name in the file, but no dosimetry
6	entries.
7	And I would have to go back check my
8	numbers because there's so many numbers, I
9	simply just don't
10	MEMBER RICHARDSON: But then
11	DR. HUGHES: remember.
12	MEMBER RICHARDSON: the total
13	workforce population
14	DR. HUGHES: The
15	MEMBER RICHARDSON: at these four
16	sites
17	DR. HUGHES: 40,000
18	MEMBER RICHARDSON: is
19	DR. HUGHES: I think is the
20	Rocketdyne. Yes, that's the
21	MEMBER RICHARDSON: Yes.
22	DR. HUGHES: entirety of the four
23	sites and the

1	MEMBER RICHARDSON: So the
2	importance of the coworker models as they've
3	been highlighted as issue here, partly relates
4	to the fact that radiation dosimetry
5	information maybe is available for 5,000 out of
6	48,000 or
7	DR. HUGHES: Well, not all of these
8	40,000 would be covered under this program.
9	This is the entire workforce that was looked at
10	by Boice who did the entire Rocketdyne,
11	Rockwell International workforce at the time,
12	so
13	MEMBER RICHARDSON: So that might
14	DR. HUGHES: this is one of the
15	issues that currently only Area IV is covered,
16	but there's also Area I, II and III which, you
17	know, the workforce was in all of these areas.
18	But, however, currently, what's
19	covered under this program is only Area IV, so
20	you would only look at the workers from Area IV
21	follow-up program. So the 40,000 encompasses
22	all workers at all sites, so.
23	MEMBER RICHARDSON: But going back

1	you had described covered periods, I think, at
2	all four.
3	DR. HUGHES: That is correct.
4	MEMBER RICHARDSON: And so it's not
5	possible that you would need to reconstruct
6	doses for somebody who had a non-covered cancer
7	at any of those
8	DR. HUGHES: Yes.
9	MEMBER RICHARDSON: four?
10	DR. HUGHES: Yes.
11	MEMBER RICHARDSON: Okay. I'm
12	sorry. You know, it's
13	DR. HUGHES: It's
14	MEMBER RICHARDSON: not easy and
15	I'm trying to catch
16	DR. HUGHES: a complicated
17	MEMBER RICHARDSON: myself up
18	DR. HUGHES: site
19	MEMBER RICHARDSON: on it.
20	DR. HUGHES: and with the
21	different areas, it gets very complicated.
22	CHAIRMAN MELIUS: Okay. Phil, then
23	Brad.

1 MEMBER SCHOFIELD: You know, if you read some of the interviews and stuff, people 2 talk about coming up for Canoga, DeSoto and 3 going into Area IV and back and forth between 4 5 the different facilities. question is what kind of 6 Мγ 7 documentation? Do they have a guard gate with a very good documentation system to know who was 8 9 coming into Area IV, who wasn't, where they were 10 coming from or in some cases even where they 11 were going? 12 Because you have these people going to these different facilities, including Area IV, 13 but they might have been stationed out of Area 14 15 II, DeSoto, Canoga. We don't know how some of these people went back and forth, but according 16 17 to their interviews, they did go back and forth. 18 DR. HUGHES: Yes, we're aware of that 19 and as far as a I know the dose reconstruction, 20 if there is such a situation, tries to address. But, however, mainly we go by the dosimetry 21 records that are available for the worker. 22

don't

assign

Since

we

23

area

monitoring, we assign the dose based 1 internal/external monitoring or in cases where 2 that's not available, using the coworker 3 models. 4 5 CHAIRMAN MELIUS: Okay. Brad. I'm 6 MEMBER CLAWSON: trying to 7 understand, a little bit like Phil, the area there because yesterday 8 as we saw how 9 everything kind of runs downhill and I'm pretty sure that the contamination that from some of 10 these went elsewhere. 11 12 How are we able to just look at Area 13 They had other DOE things going on in Area I'm trying to figure out how we're able to 14 15 just put our hands around Area IV. What's --16 MR. HINNEFELD: Well, I can just 17 offer that Area IV, if you're talking about 18 Santa Susana Field Laboratory, Area IV is the 19 covered facility. And we didn't make that 20 decision. And so we reconstruct doses that 21 occur on the covered facility. That's what the

Reconstruct doses that occurred on

statute says.

22

1	the covered facility. If that covered
2	facility affected a neighbor, the way the
3	statute stands now, that neighbor has no remedy
4	under our program.
5	MEMBER CLAWSON: Okay. But if you
6	were in Area I doing work for DOE, then that
7	tells me that you've got a problem with the
8	covered area. So how do we remedy that?
9	MR. HINNEFELD: That's another
10	agency's decision.
11	MEMBER CLAWSON: Okay. So that's
12	the Department of Labor?
13	MR. HINNEFELD: Yes, I guess. I
14	think that's Labor.
15	MR. RUTHERFORD: Let me. And
16	recognize, if we have information that there
17	was DOE work going on, I mean, real information
18	documents that there's DOE work going on, we
19	would provide that to the Department of Labor
20	and Department of Labor would make that
21	decision. We've done that I don't know how
22	many times in the past
23	MEMBER CLAWSON: Okay.

1 MR. RUTHERFORD: -- so. MEMBER CLAWSON: 2 Right. I'm just 3 wanting to understand how we can parcel that up. Thanks. 4 5 CHAIRMAN MELIUS: Wanda and then I'm going to close comments because we need to move 6 7 along here. Wanda, do you have a question or comment? 8 9 MEMBER MUNN: My comment has to do 10 with how easy it is to confuse what we're talking about when we speak about this site. 11 12 When people talk about there being four areas, 13 in my mind I'm thinking four areas are Area IV, DeSoto, Downey, et cetera, Canoga and other 14 people who are talking about four areas are 15 talking about Area I, II, III and IV. 16 17 And I would hope that we'd be very 18 careful in distinguishing that in the way we 19 talk about these things because it's very easy, 20 I think, to find yourself talking about an area 21 on top of a mountain when someone else is thinking you're talking about an area down in 22

the flats at Downey or someplace.

1	CHAIRMAN MELIUS: Thank you. Now,
2	we'd like to hear from the petitioner, Bonnie
3	Klea.
4	MS. KLEA: Is there anyone here who
5	did not go on the site tour?
6	CHAIRMAN MELIUS: Oh
7	MS. KLEA: Okay.
8	CHAIRMAN MELIUS: lots of us.
9	MS. KLEA: Anyway, my name is Bonnie
10	Klea and I've met some of you before. And I so
11	appreciate you taking another look at Santa
12	Susana. We have so many workers that have not
13	been paid, a lot of families without fathers,
14	a lot of widows.
15	Anyway, I wanted to tell you a little
16	story. I've been working with EPA for 20 years
17	since I was diagnosed with cancer in 1995.
18	I worked at the SRE and I worked in the
19	nuclear area. Wasn't told what they were doing
20	up there, didn't know anything about nuclear
21	because I was so young. But I was diagnosed
22	with bladder cancer 25 to 30 years after I was

up there. And that's second only to lung

Lung is number one, bladder is number 1 2 two. But I just wanted to tell you a little 3 story from the EPA's historical site review. 4 5 Anyway, this is the reservoir. Here's Area IV. The reservoir was built in 1919 to 6 capture the water from Area IV, 1919. 7 was a burrow, flat -- what am I trying to say, 8 There's a fault that runs from here to 9 fault. fill the reservoir and it was built in 1919. 10 In 1954, we found a memo written by the 11 12 company saying well, they'll build the reactors there and there's no problem if there's an 13 accident, they'll just divert the water. 14 That's what they did. 15 We had, of course, you know, the partial meltdown in 1959. 16 17 By 1962 we have records of monitors put in the 18 reservoir with high levels of radiation. 19 have those records. 20 By 1968, they built big concrete diversion paths for all the drainage from Santa 21 22 Susana to go directly into the river instead of

into the reservoir.

By 1969 the reservoir was closed.

And that reservoir was water served to everyone in the San Fernando Valley. I don't know how many millions of people there were.

We have a little community right over here called Hidden Lake. In the earthquake of '94, the lake was smelly and dirty and they called in a remediation company to clean it. And they said we can't clean it because your sediment is all full of TCE. So we know stuff got off site.

And in the 1959 sodium reactor experiment accident, the workers in Area I all had to have their cars repainted. And we have historical interviews with the workers that EPA did and many comments in there about having their cars repainted.

And we just had a meeting, just a few weeks ago, with John Pace, who was one of the operators trying to shut down the reactor. And he said they knew the wind data and they knew the releases were going to go all over the San Fernando Valley and Eastern Simi Valley.

And they had to release the gases or blow up the reactor. And so they released the gases for two to three weeks before they could shut it down.

But, anyway, I'm grateful that you're

But, anyway, I'm grateful that you're here taking another look at the situation. I'm the one that turned in the drinking water data, went to the Health Department and then to our County and found their records that, indeed, Ventura County and Rocketdyne knowingly at that time, gave us water that was contaminated and they knew it, from the wells.

And we were in a drought and also during that drought they started recapturing the Area IV water run-off. This is all the nuclear area. And they piped it up to these tanks up here and used it for every rocket engine test.

So not only was that water contaminated, it was storm water run-off, it was used to cool down the rocket test and so whatever was in it was airborne again.

Many of the workers are sick, who

worked in other areas. And this is Area IV, 1 across the street was Area III. We had all the 2 maintenance workers over here who would come 3 over and clean up spills and fires. 4 5 In our audience today we have one of the auxiliary fireman's widow who is here and 6 she'll be talking about how her husband was 7 called into the SRE during the accident. 8 fire 9 And he went. in with 10 extinguisher to put out spot fires, didn't know and he had to decontaminate and bury his 11 12 clothes. And he's died of cancer. 13 So we have that worker who is not 14 getting compensated because he was an engineer 15 over in Area I. And we had DOE operations 16 everywhere. 17 Over here in Area I we had an oil rig 18 that was a drop tower to test the fuel rods so 19 they could practice dropping them into the 20 We had DOE workers everywhere. reactor. 21 And I worked up here. I don't 22 remember going through a gate. Once I came in

through the main entrance I don't remember any

other place where we had to show ID to get into 1 Area IV. 2 And during the SRE accident the 3 workers were all sent home. 4 They were heavily 5 contaminated and so they pulled in workers from all the other areas to come in and to help them 6 shut down the reactor. 7 And they threw all the debris from the 8 SRE out in the back lot. And John Pace should 9 be on the phone today to talk about that. 10 So I can't think of anything else that 11 12 I'm missing. It's a very toxic site. 13 found in one of the canyons, somebody used it as a dump and they had barrels that look to 14 similar to what I've seen at Santa Susana where 15 they took barrels of waste and got behind a 16 17 board and blew them up. 18 They did that in the burn pit. 19 found 55-gallon drums in a canyon right off site 20 that all had high powered rifle holes in them. 21 So there's a lot of history and we're still 22 trying to find out everything.

anyway, thank you again for

1	coming. I can't think of anything else I'm
2	forgetting. And like I say the workers came
3	from everywhere
4	CHAIRMAN MELIUS: Yes.
5	MS. KLEA: the maintenance
6	workers, the fire department was over here, the
7	cafeteria was here. I have a lady whose mother
8	worked for one of the cafeteria companies who
9	provided food. And so the cafeteria was right
10	next to Area IV.
11	And that company's out of business, so
12	her mom who died of lung cancer can't get
13	compensated for that because they can't find
14	the subcontractor. So I'm hoping that you'll
15	get that SEC moved forward. I'm hoping through
16	the whole DOE period, I think it was '89, not
17	'87.
18	I even know clean-up workers who are
19	sick just from doing the remediation up there.
20	So thank you and I'm hoping and I'm praying
21	CHAIRMAN MELIUS: Okay, ma'am.
22	MS. KLEA: to help some of the
23	widows. Thank you.

1	CHAIRMAN MELIUS: Thank you. Okay.
2	MS. KLEA: Do you have any questions?
3	No?
4	CHAIRMAN MELIUS: I don't think
5	right
6	MS. KLEA: Okay.
7	CHAIRMAN MELIUS: now. Thanks.
8	Okay. Now, we go into a formal public comment
9	period and so I will have Ted tell you the rules.
10	MR. KATZ: Yes, just
11	CHAIRMAN MELIUS: These are pretty
12	simple, so.
13	MR. KATZ: Yes, very simple. It's
14	just for those of you who have comment, your
15	comments all of the proceedings of these
16	meetings, including this today, are
17	transcribed and published in a transcript on
18	the NIOSH website.
19	So everything you say, verbatim, will
20	be repeated there on the NIOSH website, so if
21	you have private things you talk about about
22	yourself or about another party, all of those
23	will be captured.

The material you give about yourself will be published just as you say it without any kind of redaction and without any editing.

If you talk about another person, though, that's not here speaking for him or herself, the things you say about another person may be edited, redacted to the extent they need to be to protect that person's privacy because they're not here to state that they actually want this information released to the public. So just to let you know, that's how the transcript for this will be handled.

add, if there's some private personal information or otherwise that you, or you prefer not to talk about certain issues in front of the group, there are people from NIOSH and from SC&A here who would be glad to talk to you or follow-up and so forth.

We're not limited to just what people say in the public comment period. And what we get later can be as valuable and is considered as important in terms of what applies to helping

address the issues at that site as what's said in the public comment period. So it's not weighed any differently or treated any differently onto that.

How we do this is we do go through and I have a listing and I'm going to go in sort of an initial order of what people did. We will first deal with people that are in the room and then we will go on. We take comments over the phone also and we have some people that are signed up to do that. But first I'll start with people in the room and the first person I have listed is Charlene Roesch.

MS. ROESCH: My husband wrote his occupational history before he passed away.

My name is Charlene. I'm the widow of James Roesch who worked at Rocketdyne for Santa Susana for over ten years. He died in 1998.

The details of his employment, he did many things up there as are listed in his occupational thing that he did. There was a tab that shows where they have the nuclear contamination and so let me just kind of

paraphrase or read a little bit. It's kind of hard, so bear with me.

In approximately 1957 he was assigned and trained as an auxiliary fireman. His badge number's 219 which I have with me. They had training sessions monthly and he really felt that he benefitted from that.

And moving on to 1959, you all know about the partial meltdown of the nuclear reactor. He was called in as an auxiliary fireman. He was given a hard hat and a fire extinguisher and told to go extinguish spot fires.

He went in the building. He described it as being smoky and twisted and the fire was basically out. He saw a little room with a closed door inside the reactor building and went inside. And there were, he called them like tech guys in lab coats and they were doing something feverishly around this area.

He thought so maybe later on, maybe something with the fuel rods, he wasn't sure. But when he went in they yelled at him and said

what are you doing in here? And he said, well, 1 I'm looking for spot fires. So they said get 2 out of here, there's no fire here. 3 And he left and he continued his job. 4 5 And then after the fire was all done, they took the boys to the firehouse and had them shower 6 7 and shampoo. He said it was creepy because they watched him do it, gave him coveralls to 8 9 come home in and told us to wash his clothes, which I did. 10 diagnosed 11 Не with multiple was 12 myeloma in 1996. I remember him asking the 13 doctor how do you get this, and they said well, one of the ways you get it is nuclear radiation. 14 And he went oh. And that is one of the covered 15 illnesses that we know about. 16 17 He had tumors all up and down his 18 spine. It was really advanced by the time they 19 caught it on him. And the X-rays, there were 20 a few vertebrae on his neck, especially, that 21 were just shadows on the X-ray. And they didn't know why he wasn't 22

dead or paralyzed at that point. But he went

through major chemo, radiation and then a drug called Aredia, which helped to rebuild bones.

And for a while he was basically kind of cancer free for a little bit, but they don't call it really remission. And then it came back with a vengeance and he died in February of '98.

And, basically, been denied compensation because he didn't normally work in Area IV. He was a mechanic in Area I, if I remember. And so every time it would come up again, they'd say oh, you have a wonderful case, you know, no problem, you'll get compensation. And then because he's not an Area IV, it was denied.

So I thank you for listening. If anybody has any questions. He wrote his history down and he did a lot of things up there. Then the last page shows an article that was done years ago with me holding his auxiliary fireman's badge which I brought if you want to see it. But it's not contaminated. He didn't have it on that day.

1	And so thank you for your attention to
2	this matter. And I certainly hope that things
3	happen, not only from me, but I've heard a lot
4	of sad stories from other fireman families as
5	we've gone to many of these meetings and it's
6	hard every time we do it.
7	But if people get compensated or at
8	least get recognized for what they did, it would
9	be nice. So I thank you. Any questions?
10	CHAIRMAN MELIUS: Okay. Thank you
11	for
12	MS. ROESCH: Thank you.
13	CHAIRMAN MELIUS: sharing. And
14	the next person I have listed is Lorraine
15	Kurowsky, is it?
16	MS. KUROWSKY: Kurowsky.
17	CHAIRMAN MELIUS: Yes.
18	MS. KUROWSKY: I have a similar
19	story. My husband started to work in the
20	CHAIRMAN MELIUS: Well
21	MS. KUROWSKY: area quite
22	CHAIRMAN MELIUS: can you get up to
23	the mic and then start over again. I'm

My husband started 1 MS. KUROWSKY: working at Canoga Park 12/11/78, and they 2 refused my application for recognition for what 3 And they said, well, he didn't work 4 he did. 5 with radioactive material enough. He was only 22.78 percent. 6 And then they said that he wasn't 7 really working with radioactive material, but 8 9 yet he told me two stories that stayed in my 10 mind. that 11 One was there was a deer 12 struggling walking up in that area in IV at 13 Santa Susana. And he says it was struggling and it was collapsing and it would try to get 14 And he said after a while it just 15 back up. 16 expired. 17 And someone said, hey, you guys in 18 this area go clean it up. And my husband was 19 a blue collar worker and he did crating and 20 packing. So he says go clean it up and let's 21 pack it up and get rid of it, whatever get rid

Then also they were dispersing a truck

of meant. I don't know.

22

and my husband worked with this little guy. 1 Just my husband, he's a big guy, 6'5" and he 2 worked with this little guy who was, what, about 3 5-foot. 4 And my husband was going on the back 5 of the truck lifting these, what they called 6 7 pigs to take them off of the truck, and making sure that it wasn't too heavy for his coworker. 8 9 So he says, hey, he says you can handle 10 some of these, but I'll stay up here. And he 11 was handling these pigs that I don't even know 12 what they were. But anyway, he would hand them to him. 13 And when he handed it to his co-worker who was 14 this little guy, he handed it to him and it just 15 went off. 16 And then grabbed this Mr. Waco and 17 18 took him off the site and washed him down and 19 took his clothes and gave him coveralls and sent him home. 20 21 And he did get cancer and die of 22 cancer. But they never touched my husband. 23 They never said anything to him. He says, you

know, just go on your business and that was it. 1 And yet they said because he had 22.78 2 he wasn't qualified. Well, he died in '01 -- in 3 '03, I'm sorry. He had '01 in cancer in his 4 5 pancreas and they said it wasn't part of what the cancer they were looking for, but then 6 again, they said, well, it could have been, but 7 it was always not here not there. 8 9 And as the other lady that stepped up, we feel sorry for the people that weren't 10 compensated because we did lose our husbands. 11 12 My children lost their father. 13 And it's just hard. And just thinking we should have been recognized in some 14 Thank you. 15 way. 16 CHAIRMAN MELIUS: Okay. Yes. 17 Again, just if you want to talk to some of the 18 people here from NIOSH and may be able to do some 19 follow-up on the individual case if there was. 20 I'm not sure how much information they had when 21 they were doing the dose reconstruction. 22 may be helpful. But thank you. D'Lanie

23

Blaze, is it?

1	MS. BLAZE: D'Lanie.
2	CHAIRMAN MELIUS: D'Lanie, I'm
3	sorry.
4	MS. BLAZE: Yes.
5	CHAIRMAN MELIUS: It's not your
6	writing, it's my eyes.
7	MS. BLAZE: The acknowledgment of
8	Santa Susana Field Laboratories complete site
9	history on behalf of Area I, II and III
10	personnel has been a passion of mine since 2007.
11	I'm very happy that the Presidential
12	Advisory Board on Radiation and Worker Health
13	has come back to Santa Susana Field Lab with an
14	interest in touring Areas I, II and III and
15	that, at last, an extension to the Area IV SEC
16	is moving forward.
17	We are grateful for your presence and
18	your efforts on behalf of Santa Susana Field Lab
19	personnel. Worker advocates and SEC
20	petitioner and former worker and a site
21	historian were not permitted to go on
22	yesterday's tour of the facility.
23	In lieu of our absence, it was

requested that we provide a list of things that we'd have liked to point out to the Advisory Board if we had been on the tour.

Each of you were provided with a copy to your guide to sightseeing hot spots of the Santa Susana Field Lab which features a detailed fold-out map of Atomic Energy Commission and Department of Energy activities throughout Areas I, II and III.

This guidebook is based on historical documents that were published by Department of Energy, its contractors and other agencies.

All information is cited. All resources are included in their entirety on an accompanying disk. And over 300 worker interviews that were conducted by Department of Energy and the Environmental Protection Agency in 2009 are also included.

Upon your review, we are confident that you will find that Areas I, II, and III satisfy legislative criteria that is used to determine a Department of Energy facility under the statute.

Currently, Santa Susana Field Lab claimants are denied EEOICPA compensation at a rate of nearly 90 percent, often based solely on an Area I, II or III work location.

However, Area I, II and III workers, employees of a DOE contractor, were rotated to Canoga and DeSoto facility regularly.

Those are both SEC facilities wherein the SEC covers all workers. It is therefore illogical to exclude Area I, II and III personnel from EEOICPA.

Further, as illustrated the quidebooks, Atomic Energy Commission Department of Energy's engagement in nuclear laser and coal gasification research, waste, storage and disposal of Area IV material, personnel who were monitored for radiation exposure, accidents and spills involving DOE waste and construction, modification integration of new and existing facilities for use by or on behalf of Department of Energy throughout Areas I, II and III of SSFL is very well documented.

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Again, this documentation has been 1 provided in DOE's own words, from their own 2 published documents. There is no denying any 3 documented history by this agency and its 4 5 predecessors. And Department of Energy may have put 6 it best in their statement that, "Historically, 7 great benefits have been obtained by separating 8 9 growing and diverse programs and test facilities at the Santa Susana Field Lab." 10 It is my sincere hope that you will 11 12 ardently support and argue for the inclusion of 13 Area I, II and III personnel to EEOICPA in 14 accordance with the legislation and in the 15 spirit of the Act as it was intended by 16 Congress. I wanted to address the question on 17 18 the tritium plume that was raised after SC&A's 19 The tritium plume would be presentation. underneath the SNAP Area IV Number 59. 20 21 And Department of Energy acknowledged 22 transport of contamination and tritium to the

site wide reclaimed water system and stated in

the guidebooks, that you guys have, that the drainage from SNAP Building 59 reached the Silvernail pond in Area III which was the entry point to the site wide water reclaim system.

Now tritium has also been discovered in Areas I and II. And there are two documents, specifically, on the disk that you were provided yesterday that speak to the tritium plume, one by the EPA, the Rocketdyne Santa Susana Field Lab sample analysis report from 1989 and the document, Radiation Protection and Health Physics Services, tritium production and release to groundwater at Santa Susana Field Lab. That's on your disks.

I thank you very much for coming to Santa Susana and for the opportunity to comment.

CHAIRMAN MELIUS: Okay. Thank you. Okay. Is there anybody else in the audience here, who wishes to speak to the Santa Susana site before I go to the phone? Okay. Good. Is there anybody on the phone who wishes to speak to the Santa Susana site or offer

1	comments?
2	MR. PACE: This is John Pace. Can
3	you hear me?
4	CHAIRMAN MELIUS: Yes, we can.
5	Thank you, Mr. Pace. I had you
6	MR. PACE: Okay.
7	CHAIRMAN MELIUS: on the list
8	here. Go ahead.
9	MR. PACE: Okay. Well, I would like
10	to help Mrs. Roesch out there. She spoke, one
11	of the first speakers. Her husband was a
12	fireman and I was, when she told the story, I
13	was there at the time and at the SRE reactor when
14	her husband through the door to help put out the
15	fire that we had.
16	We had a fire and explosion in the high
17	bay area at the time, pulling out the sodium
18	pump which caused the reactor to go down in
19	1959.
20	And we were trying to replace it. As
21	we was gotten it all lose and a man come out of
22	the down the floor where it was at, came up.
23	And we had a kind of a tent we had covered over

to keep the oxygen out.

And then when he came through, somehow it allow oxygen in the area where that sodium was at with the sodium pump and it had a big explosion and a fire on top of that.

And that's when Mr. Roesch came with the, you know, fire department to see if he could help us out. And he come through a small room that we had there and into the high bay area.

And we yelled at him to get out of the building or out of the deal just exactly like his wife just had told you. And because we was worried because with the explosion and all the radiation that came out of the pump area, it was only three feet from the reactor where we was working.

And we was worrying about having other fires here, of more radiation coming up out of the reactor core. So what she tells you is actually true and I'm a witness to that.

And I was there on that. That was an incident that I've talked about many times and

NIOSH knows about it. It's on the record with them. But I haven't had anything for evidence to prove to them that this occurred.

And I got exposed to a lot of radiation myself on that. Me and three other men, we got blown clear across the room on our tippy toes there trying to keep from falling down.

With that explosion we ended up all having about four showers apiece trying to get that radiation and contamination off us when it happened.

And then most of them went home in the coveralls, we call redlines, what we wear for protection. And I happened to be lucky and have clothes in my locker I was able to wear home.

But I just wanted to be of verification to Mrs. Roesch that I was there and I'll be glad to help her on anything. And I would like to make one, just real quick point, is the radiation that came out of the SRE reactor, and I was there, I was there at the time of the nuclear accident and afterwards.

I was there and my crew had started the 1 reactor up and ran the reactor for the two 2 That's all everybody talks about. 3 weeks. 4 They ran a broken reactor, a reactor that wasn't 5 suitable for running, but we was told to start it back up again. 6 7 during that time, а lot radiation leaked out of that reactor through 8 9 the fuel elements, the seals around there. 10 reactor had gotten so hot before we was able to get it shut down on the 13th that it damaged that 11 12 it was leaking out into the high bay area. So there's lots there that NIOSH needs 13 14 to learn about and I've already spoke on this I don't want to do a lot of talk on it 15 before. because there's records of it already, but key 16 17 thing was just for Mrs. Roesch there. 18 I was there and I'll be glad to work 19 with her and help her out on that incident there 20 with her husband. So if there's any questions 21 anybody would like to ask me that would be fine. 22 And anybody have a question at all or B-

CHAIRMAN MELIUS:

23

I don't think right

1	now. But thank you very much, Mr. Pace, for
2	MR. PACE: Okay. And thank you.
3	You
4	CHAIRMAN MELIUS: Yes.
5	MR. PACE: And bye, now.
6	CHAIRMAN MELIUS: Sir. Anybody else
7	on the phone that would like to make comments
8	on Santa Susana? Okay. If not, I'm going to
9	go I have at least three people that have
10	wanted to make comments, I believe, on other
11	sites.
12	The first one is Terrie Barrie.
13	Terrie, are you on the line? Okay. Terrie, do
14	you maybe have a phone on mute or are we off?
15	MS. BARRIE: All right. Can you hear
16	me?
17	CHAIRMAN MELIUS: Oh, now we can.
18	There you are. Okay.
19	MS. BARRIE: Okay. Do you hear me
20	now?
21	CHAIRMAN MELIUS: Yes, we can. Yes.
22	MS. BARRIE: Okay. I'll start all
23	over again. Good evening, Dr. Melius and

Members of the Board. This is Terrie Barrie of the Alliance of Nuclear Worker Advocacy Groups and I thank you for allowing me to call in my comments.

Part of my comment does concern Santa Susana. So I was happy to hear that some of the Board Members and members of the community questioned why Areas I, II and III are not covered under this program.

From the oral history I've heard from the workers and their advocates and apparently attested to tonight, it was not uncommon for the workers from Santa Susana to be detailed from one area and assigned to another.

It seems common practice with all DOE workers. And as you know, workers assigned to Areas I, II and III have been denied classification as a DOE worker and are not covered under this program.

A month or so ago, I found a Department of Labor final decision which spells out the criteria necessary for which DOL will designate a site as a covered DOE facility.

The test needed to overcome is that the claimant or advocate must prove that Department of Energy had use of or controlled the site. I need to emphasize the word or. It appears that Department of Labor has a tendency to ignore that word in the statute.

The law states the DOE must have proprietary interest in the facility or have a certain type of contract with that facility. The law does not state DOE must have proprietary interest and a service contract.

The final decision states quite clearly what is needed to prove proprietary interest. And I quote, "The evidence must establish that the MED, Manhattan Engineering District (Department of Energy), had rights of ownership, use or control of the buildings in which the employee worked."

D'Lanie Blaze just prepared an excellent guidebook for the Board which gives many examples of the Department of Energy's use of Areas I, II and III.

It is my hope that the Board and the

affected claimants petition the Department of Labor to designate Areas I, II and III as a covered DOE facility. And I'd be happy to supply the link to that final decision.

I'd like to turn now to the issues with the Rocky Flats SEC petition. I want to thank LaVon Rutherford for his offer to discuss the issues and I'm looking forward to our call next week.

For those of you who are not familiar with this, here's a brief summary. A man who was assisting a family member with cancer who worked at the Rocky Flats facility after 1993, which is the end of the current SEC Class, alerted me to a recent release of NIOSH's occupational internal dose document for the Rocky Flats facility.

The document states, and I quote, "Because of data issues and limitations, no specific methods to bound doses from 233 uranium and 232 uranium had been determined. Therefore, doses to unmonitored RFP workers from neptunium, thorium and 233 uranium and its

associated 232 uranium and 228 thorium 1 contaminates cannot be reconstructed." 2 One would think that means since NIOSH 3 cannot reconstruct dose for these elements, 4 that the SEC Class should be extended. 5 Please note there is no qualifying 6 7 statement that limits the years NIOSH cannot reconstruct those in that document. But. 8 9 apparently, this statement is not enough for NIOSH to recommend to the Board to withstand the 10 SEC Class for Rocky Flats. 11 12 It is my understanding that for some 13 reason NIOSH needs to release the White Paper on neptunium, and I might be wrong about this, 14 15 before making any kind of recommendation to the 16 Board. But if they already know they can't 17 18 reconstruct dose for these elements, why is it 19 necessary to wait for the release of this paper. 20 I must remind everyone that time is 21 something many sick workers do not have an 22 abundance of. A friend of mine passed away He did not fall within the 23 this summer.

current SEC Class.

And as for the worker I mentioned earlier tonight, even if the Board recommends tomorrow to expand the Class for the Rocky Flats facility, outside of a miracle that worker will not live to receive the deserved compensation because of the aggressive nature of the cancer.

Please keep the deteriorating health of the workers in the forefront of this process. Thank you again, for allowing me to call in these comments and I look forward to my conversation with LaVon.

CHAIRMAN MELIUS: Okay. Thank you.

For those of you that weren't part of the emails, there was -- Terrie Barrie and LaVon scheduled a conference call for next week sometime. I can't remember the date, but -- MR. RUTHERFORD: Next Wednesday at 1:00.

CHAIRMAN MELIUS: Next Wednesday to try to clarify that particular issue. Anyway, thank you. The next person I have to be listed is Jeff Schultz. Jeff, are you on the line?

1	MR. SCHULTZ: Yes, I'm here.
2	CHAIRMAN MELIUS: Oh, okay. Go
3	ahead.
4	MR. SCHULTZ: Thank you for the
5	opportunity, everybody, to speak tonight. My
6	name is Jeff Schultz and I'm in Westminster,
7	Colorado.
8	I'd like to comment tonight on the SEC
9	Petition 192 regarding neptunium-237 exposure
10	at the Rocky Flats plant. My document's
11	recently been made available on the internet.
12	The title of the document is
13	Validation of Rocky Flats Plant Radionuclide
14	Inventory and the Historic Data Using the SWEPP
15	Assay Data and it's dated August 2004.
16	The abstract document states that,
17	"This report presents the results of a
18	descriptive statistical analysis of isotopic
19	characteristics of radioactive waste stored at
20	the Idaho National Engineering and
21	Environmental Laboratories Radioactive Waste
22	Management Complex."
23	In the body of the document there's a

1	section 4.1.4 neptunium. And the document
2	states, "At least some of the SWEPP waste drums
3	contained neptunium-237. However, neptunium
4	quantities are not measured or calculated by
5	the PAN/gamma system. Neptunium data are only
6	available when the SGRS absolute system is
7	used.
8	Of the SWEPP graphite waste drums,
9	only four were assayed using the absolute gamma
10	system, hence, data on neptunium is very
11	limited."
12	Section 4.2.4 states that,
13	"Measurable quantities of neptunium-237 were
14	found in all but four waste drums for which
15	there was SGRS data."
16	Section 4.3.4 states, "Since there
17	were only neptunium data for 14 mixed metal
18	waste drums, plotting of the histograms is not
19	useful."
20	Section 4.5.4 states no neptunium
21	data were available for organic setup waste.
22	Section 4.6.4, "no neptunium data were
23	available for special setups weight."

From this document it's clear that 1 waste shipments to INL were being scrutinized 2 for the content of neptunium-237 in this 2004 3 document. 4 5 Further, they using were а statistical approach to guess what amount of 6 neptunium-237 was used from a fraction of data 7 sampled from the drums that were surveyed with 8 9 the SGRS system. A coworker of ours on a crew in 10 11 Building 371, who was assigned to survey drums 12 with а Canberra SGS system in 2002, 13 [identifying information redacted] task was to survey legacy drums which had been in storage 14 15 for many years in the plant. [identifying information 16 Ву redacted] account some of the drums had no 17 18 labels, other had labels that deteriorated 19 making them illegible. Some drums were re-labeled with information as to what they 20 21 thought was in those drums at that time.

with this SGS system, they did a step where they

In the process of surveying the drums

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actually opened the lid of the drum and took a 1 gas sample from inside the drum. 2 During 3 one of those tests, [identifying information redacted]. 4 5 The fumes from the drum had caused [identifying information redacted]. The 6 7 fumes were probably generated from radiolytic decay of the plastic bags, the chemicals and the 8 radiation all doing their work in the waste drum 9 10 over that period of time. [Identifying information redacted]. 11 12 But [identifying information redacted] does remember operating the SGA 13 system and finding neptunium-237 in some of the 14 drums that [identifying information redacted] 15 [identifying 16 surveyed. And information 17 redacted] has one document in [identifying 18 information redacted] possession that shows 19 the survey of a drum and neptunium-237 was 20 present in that drum in that 2002 period. 21 [Identifying information redacted] 22 also commented that some of the drums [identifying information redacted] surveyed 23

and repacked contained waste from the 1969 Mother's Day fire that occurred in Building 776 and Building 777.

And this is a testament that waste was store in these drums at the Rocky Flats Plant for a very long time. 1969 waste was still sitting in drums.

Our workers contend that detection of neptunium at the site was largely missed since this is very weak gamma and it's merely lumped in with the other gamma signatures and it shows up in the total plutonium count.

And that they weren't really tasked with looking for neptunium. There was no reason to look for it. Only the real modern SGS equipment that was brought to the site around the year 2000 could distinguish between plutonium and neptunium. Reliable neptunium detection in the old days would have required the use of a spectrometer.

The Rocky Flats plant had many barrel storage areas. Drums were constantly moved around by workers between 90-day satellite

storage areas.

A shell game was conducted, so that drums in these storage areas wouldn't violate the storage time limit of 90 days in these so-called temporary storage areas.

So when the 90 days was up drums would start being moved from one area to another.

Materials in suspect corroded drums were repacked into new drums over the years.

When a new requirement was instituted to install carbon filters on all the drums to relieve a possible problem of hydrogen build-up in the drums and possible explosions, a project was started and around 10,000 drums were re-lidded with new lids that had a threaded opening where they could install the carbon filter.

Many drums had to be surveyed and repacked over the years to comply with plutonium limits dictated by the waste sites to allow our shipments. So the drums were handled a lot.

As we ran out of space in the

designated storage areas and because of extended periods of waiting for the website to open and a period of time where shipments to the INL area were curtailed by the governor of Idaho new drum storage areas had to be created because the main drum storage areas were full.

So drum storage started being pushed into the process rooms where the people were. And as workers, we had to work around these hot drums and we had to be shined by these drums on a daily basis.

This caused a lot of exposure to employees. Some of these drums contained neptunium as proven by the fact that neptunium is showing up at the waste sites.

I'm in the process of trying to find another coworker who works at the Savannah River plant. He was quoted by another employee as saying that waste shipments received at the Savannah River plant from the Rocky Flats plant were surveyed at Savannah River and they were found to have neptunium in them.

And this occurred in the time period

of the early 2000s when Rocky Flats was closing and shipping their waste around. When the plant finally closed, the equipment used for the pre-1984 time period when neptunium work was done, that equipment was demolished and removed.

In the process of removing this equipment, ventilation duct work that went to this equipment and the work of the equipment itself exposed our workers to neptunium contamination that was left behind.

This equipment went into waste crates or into cargo containers ultimately shipped to the Nevada Test Site. And this material is really only surveyed for plutonium, uranium, and americium. They, again, did not count the neptunium present.

I just wanted to ask the Board Members to consider this evidence that Rocky Flats workers were exposed to neptunium-237 well into the 2000s when the plant was closed and demolished and consider extending our SEC further out to the closing date. And thank you

1	again for the opportunity to comment this
2	evening.
3	CHAIRMAN MELIUS: Okay. Thank you.
4	And we will follow-up on this and, obviously,
5	NIOSH is here in the audience, too, and they're
6	actively evaluating Rocky Flats. And also our
7	contractor is here, so this information will be
8	available, obviously, to them for follow-up
9	also. So again
10	MR. SCHULTZ: Thank you.
11	CHAIRMAN MELIUS: we appreciate
12	the thoroughness of your follow-up. That
13	was
14	MR. SCHULTZ: Thank you.
15	CHAIRMAN MELIUS: helpful. Good.
16	The other person I have listed who wanted to
17	make comments on the phone is Dr. Dan McKeel.
18	Dr. McKeel, are you on the line?
19	DR. MCKEEL: Yes, I am. Can you hear
20	me?
21	CHAIRMAN MELIUS: Yes, we can very
22	well.
23	DR. MCKEEL: All right. Well, good
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the Board Members and 1 afternoon to I'm Dan McKeel. I'm the General 2 assembled. Madison 3 Steel Industries and Dow SEC co-petitioner. 4 First off, I certainly will sincerely 5 thank the Board for tasking SC&A to review the 6 Dow Madison Appendix C Revision Number 1 that 7 was issued on April 3rd of 2014. 8 I'd ask that this be done twice and I 9 certainly appreciate that it now is going to be 10 As was stated this SC&A review is 11 done. 12 absolutely necessary, the first step toward 13 NIOSH issuing a Program Evaluation Report for Dow even though Director Hinnefeld tells me 14 that the PER for that site is being prepared. 15 I don't know how far along it is. 16 But my remarks tonight mainly address 17 18 processing of a revised GSI Site Profile, which 19 is Appendix BB, Rev 1 that Dr. Ziemer mentioned 20 in his TBD-6000 Work Group report earlier this 21 afternoon. 22 I would note that a GSI Program

Evaluation Report based on GSI Appendix BB, Rev

1, which was issued June 6, 2014, has been 1 stopped pending release of SC&A's review which 2 was just released a few days ago for Appendix 3 BB, Rev 1. 4 5 The dose reconstruction Subcommittee at its 10/29/14 meeting declined to make a 6 detailed review of four completed GSI dose 7 reconstruction cases based primarily on the 8 9 future availability which was said to tomorrow by Mr. Mauro of SC&A's review of 10 Appendix BB, Rev 1. 11 12 So that document was not available for 13 the Subcommittee Members to review. The SC&A memo we're talking about was drafted by Drs. 14 Robert Anigstein and John Mauro of SC&A. 15 The dose reconstruction Subcommittee 16 17 Members on October the 29th were assured by John 18 Mauro that the SC&A finds were minor and would 19 not require changes in dose reconstruction 20 practices for GSI. SC&A stated the main

Acting chair Wanda Munn, for the dose

changes involved language tweaking primarily.

And this was also suggested today by Dr. Ziemer.

21

22

reconstruction Subcommittee assured the Members that all dose reconstructions shortcomings and NIOSH four methods had been dealt with from the previous Rev 0 of Appendix BB which came out in June of 2007.

The Board DFO, Ted Katz, admitted that these older methods were now outmoded. And he declared, and I'm quoting him, "This committee is done with these cases."

Should this unfortunate decision stand, no GSI completed dose reconstruction will have been reviewed by the DRSC. All of this was really shocking to me because I'd been asking Ted Katz and DRSC chairs for years about these GSI case reviews with completed dose reconstructions without getting any satisfactory answers on why no cases had been reviewed, what the case numbers were et cetera.

Anyway, I want to now turn to the comment about my part in Appendix BB, Rev 1. The SC&A review of the revision of Appendix BB also address in part, an extensive critique dated July the 16th, 2014, that I had made of

the revised DCAS GSI Site Profile.

I received my copy of the Anigstein-Mauro SC&A memo on November the 3rd and then wrote an eight-point initial reaction memo to all Board and TBD-6000 Work Group Members the same day.

My concerns were initially, besides some of the content and the findings of the SC&A report that even though SC&A was reviewing my work as well as that of Dave Allen and NIOSH, my name was not actually mentioned in the report title, the body text or in the references of the SC&A memo.

However, they did quote findings of mine, but they refer to me as the GSI co-petitioner throughout.

So despite these allusions to my work, my White Paper, which was 87 pages, critiquing Appendix BB, Rev 1, which was posted on the DCAS website for three-and-a-half months before the SC&A review was released was not cited in the text of in the references in their 10/29 GSI Site Profiles review memo.

I observed in reading through the content of the body of the report that SC&A had cherry-picked and briefly mentioned a few of my many objections to the Allen DCAS 6/16/14 Appendix BB paper.

I equated this tactic to DCAS's Dave Allen's throw them a bone technique. Mr. Allen admitted using this strategy as demonstrated by email for the Hooker Electrochemical site that was obtained by the site petitioner through a FOIA request. DCAS director Hinnefeld later apologized to the Board for these actions on the part of his DCAS personnel.

Further review of the 10/29 Anigstein-Mauro Appendix BB, Rev 1 review memo, and this is the most important thing I'll say to you tonight, showed that there were eight new SC&A findings.

Several of which will require extensive new modeling and dose recalculation by NIOSH. An example, which I also pointed out in my Appendix BB, Rev 1 review was that GSI radiographers during the radium 226 area were

also exposed to the betatron beam and activated 1 high nickel steel castings with respect to 2 photons and betatron beam neutrons. 3 4 had not included those NIOSH 5 important doses from the betatron to GSI radiographers during the radium 226 era. 6 So SC&A noted they had to be added. 7 Also SC&A differed with NIOSH and Dave 8 Allen on skin beta doses from the betatron and 9 said that those doses needed to be resolved as 10 well, and so on. 11 The remedies and resolution of these 12 13 eight new SC&A findings on Appendix BB, Rev 1, 14 would likely occupy several more meetings and White Papers to be resolved completely. 15 Please recall the 13 SC&A findings on 16 17 Appendix BB, Rev 0 from 2007 took seven years 18 until January of 2014 to be considered to be 19 completely resolved by the TBD-6000 chair as 20 stated in his report to the Board earlier this 21 year. 22 For the record, the correct full

citation from my White Paper dealing with

1 Appendix BB, Rev 1 is as follows. The URL, the link it is 2 to http/www.cdc.gov/NIOSH/ocas/gsi.html. 3 And this is part of NIOSH docket 140 for the GSI AWE 4 Illinois site. 5 The full citation on the DCAS web page 6 7 is submission by Daniel W. McKeel, Jr., M.D., GSI SEC 1005 co-petitioner. And the title is 8 9 Critique of GSI, Appendix BB, Rev 1 by Dave 10 Allen, DCAS, 6/6/14 and it's a pdf two megabyte 11 87 page file. 12 In light of all this, I feel even more 13 strongly that the Board has been seriously misled by the TBD-6000 Work Group including the 14 15 SC&A and NIOSH members on the finality of resolution of the original 13 findings of 16 Appendix BB, Rev 0, both leading up to and 17 18 during the final nine to eight vote for GSI SEC 19 105 on 12/11/12. 20 And this misleading has continued 21 even afterwards. Even today it is abundantly Site 22 clear that all GSI Profile dose

reconstruction findings from the Rev 0 June

2007 version are not fully resolved.

NIOSH and DCAS have more details to work through. And for the same reasons, I believe the D.R. Subcommittee Members were misled on October 29th by SC&A and John Mauro. The new findings are not all minor and cosmetic, far from it. More dose calculations are required by NIOSH.

Finally, I note the SC&A memo was included as a discussion paper for this meeting. During the Work Group reports, Dr. Ziemer mentioned he had seen the SC&A Appendix BB, Rev 1 review.

He did not mention that SC&A was tasked by the Board and the DFO to also review my detailed White Paper on that same Rev 1, Appendix BB document.

Finally, I sincerely and humbly ask all Board Members to please read all three papers, Appendix BB, Rev 1 issued 6/6/14, the Dan McKeel critique of Appendix BB, Rev 1 issued on July the 16th, 2014 and the Anigstein-Mauro SC&A memo about Appendix BB, Rev 1 that was

1	issued on October the 29th, 2014.
2	Thank you very much and I appreciate
3	you're letting me address you. Thank you.
4	CHAIRMAN MELIUS: Thank you. Is
5	there anybody else on the line that wishes to
6	make public comments? Okay. If not, I
7	believe that concludes our public comment
8	session and concludes our meeting. So we will
9	talk to everybody again after the holidays.
LO	MR. KATZ: Thank you, everyone, for a
L1	productive meeting.
L2	(Whereupon, the above-entitled
L3	matter went off the record at 5:46 p.m.)
L4	
L5	
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L8	