# U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES CENTERS FOR DISEASE CONTROL NATIONAL INSTITUTE FOR OCCUPATIONAL SAFETY AND HEALTH

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## ADVISORY BOARD ON RADIATION AND WORKER HEALTH

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SUBCOMMITTEE ON DOSE RECONSTRUCTION REVIEWS

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TUESDAY APRIL 1, 2014

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The Subcommittee convened telephonically, at 10:30 a.m., Eastern Daylight Time, David Kotelchuck, Chairman, presiding.

#### PRESENT:

DAVID KOTELCHUCK, Chairman BRADLEY P. CLAWSON, Member WANDA I. MUNN, Member DAVID B. RICHARDSON, Member

#### ALSO PRESENT:

TED KATZ, Designated Federal Official BOB BARTON, SC&A HANS BEHLING, SC&A KATHY BEHLING, SC&A RON BUCHANAN, SC&A ZAIDA BURGOS, NIOSH GRADY CALHOUN, DCAS DOUG FARVER, SC&A ROSE GOGLIOTTI, SC&A DeKEELY HARTSFIELD, HHS STEVE MARSCHKE, SC&A BETH ROLFES, DCAS MUTTY SHARFI, ORAU Team SCOTT SIEBERT, ORAU Team MATT SMITH, ORAU Team JOHN STIVER, SC&A

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

2 | 10:36 a.m.

MR. KATZ: So let's begin with roll call first. And because it's complicated with so many sites with dose reconstructions, I'm just going to run through myself the recusals for Board Members that are on the line. And then if Mark joins us, I'll cover his at that point, too, rather than you having to remember your recusals.

So just roll call. I know who's on the line now. I'll just run down your names.

(Roll call.)

MR. KATZ: So that covers that.

Let me just also note that some of the materials related today that can be posted or are posted on the web site under today's meeting date of the Subcommittee. All of the Board Members and staff should have all of the materials, I believe, that we're discussing today because they've been distributed by multiple parties in many cases, myself sort of duplicating other

1	people's distributions to some extent.
2	And that's it for me. The agenda is
3	also posted on the web site. Dave, it's your
4	agenda, and off we go.
5	CHAIRMAN KOTELCHUCK: Well, okay.
6	Welcome, folks. This is, as I noted before, an
7	unusual two-day meeting because we had to
8	postpone our last meeting for lack of a quorum.
9	So thank you all very much for being on the line
10	and for being here for the two days. I hope
11	this is a rare back-to-back meeting.
12	So the first item on the agenda is
12 13	So the first item on the agenda is the dose reconstruction method for
13	the dose reconstruction method for
13 14	the dose reconstruction method for Westinghouse Nuclear Fuels. I don't know who
13 14 15	the dose reconstruction method for Westinghouse Nuclear Fuels. I don't know who wants to start on that. John, perhaps you?
13 14 15 16	the dose reconstruction method for Westinghouse Nuclear Fuels. I don't know who wants to start on that. John, perhaps you?  MR. STIVER: Yes. Hans had
13 14 15 16 17	the dose reconstruction method for Westinghouse Nuclear Fuels. I don't know who wants to start on that. John, perhaps you?  MR. STIVER: Yes. Hans had prepared the memo to Drs. Lemen and Field after
13 14 15 16 17 18	the dose reconstruction method for Westinghouse Nuclear Fuels. I don't know who wants to start on that. John, perhaps you?  MR. STIVER: Yes. Hans had prepared the memo to Drs. Lemen and Field after the one-on-one when this case came up. And I
13 14 15 16 17 18 19	the dose reconstruction method for Westinghouse Nuclear Fuels. I don't know who wants to start on that. John, perhaps you?  MR. STIVER: Yes. Hans had prepared the memo to Drs. Lemen and Field after the one-on-one when this case came up. And I believe Hans is on the line now.

1	memo, and if you would like to
2	DR. H. BEHLING: Yes. Just a quick
3	- I'm not sure if Ted Katz should have asked for
4	a roll call because no one has been asked to
5	identify themselves.
6	MR. KATZ: Oh, I'm sorry.
7	DR. H. BEHLING: We didn't do the
8	roll call.
9	MR. KATZ: I did only the Board
10	Members, and we jumped right into it. That's
11	my fault. Sorry.
12	(Roll call.)
13	MR. KATZ: Okay. Very good.
14	Sorry for having omitted this, Jim. You don'=t
15	have to transcribe this attendance, but on we
16	go again. Thank you.
17	MR. STIVER: Hans?
18	DR. H. BEHLING: Okay. Just to
19	recap what has already been stated by John
20	Stiver, this whole issue centered around a dose
21	reconstruction case that I reviewed involving
22	a person who was working for the Westinghouse

Nuclear Fuel Division and is really part of an 18th set. So I assume we're not going to be talking about this particular dose reconstruction, other than the issue that prompted this memo.

As part of a one-on-one that was done back in January, this past January here, we identified a couple of findings that made it to a document that was never identified to SC&A, and it's really not a Site Profile. But it was included in my assessment of this particular dose as part of Appendix B. And it is a entitled document that's "A Dose Reconstruction Methodology for Westinghouse Nuclear Fuels Division, Cheswick, Pennsylvania". It's a very short document but identifies certain it aspects for reconstruction involving people who may have been working at the Westinghouse facility during the time of operation of '71 - '72 and for the residual period, that is several years.

And part of my review, as always, we

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look at the dose reconstruction, and the first thing we do is try to duplicate any number that is identified for either external exposure or for internal exposure. We agree that the number is almost immaterial, but our first effort is always to simply try to reproduce the numbers that NIOSH has introduced in the dose reconstruction report for that individual.

And one of the things that I did was to look at really the internal exposure values that were cited on behalf of this particular case as just as a way of trying to get everyone on board here. And I'm going to ask John Stiver to perhaps identify a page of the report that might also be helpful for the people who are online here. John, are you in a position to provide us with that?

MR. STIVER: Hans, you're kind of breaking up there. I didn't quite hear you.

DR. H. BEHLING: I was hoping that you can identify a couple of pages that will help me explain what this issue was that came

1	to light as a result of a one-on-one. And what
2	I was hoping that you can do is to identify pages
3	on that particular report that I had enclosed
4	as part of my dose reconstruction that was part
5	of Appendix B and is in the "A Dose
6	Reconstruction Methodology for Westinghouse
7	Nuclear Fuel Division, Cheswick,
8	Pennsylvania." Can you pull up that document?
9	I believe it starts on page 139.
10	MR. STIVER: Okay. Let me see if I
11	can get that here.
12	DR. H. BEHLING: If you can't, I
13	will try it would just make it so much simpler
14	to identify a few things.
15	MR. STIVER: Okay. I've got it.
16	Let me go ahead and share it here.
17	DR. H. BEHLING: Okay. Can you
18	perhaps - let's see here pull up page
19	MR. STIVER: Okay. Do you see it
20	now?
21	DR. H. BEHLING: Yes. Can you
22	perhaps pull up page 141?

MR. STIVER: Is this what you need right here?

DR. H. BEHLING: Yes. And just as a summary for review here, the people in that facility were potentially exposed to three different types of radionuclides. And on page 141, you will see one of three options that would be permissible in a reconstruction of internal dose for an individual during that time period.

The first one is on page 141, a 2 percent enrichment cycle uranium ratio. And you'll see on the far left side the radionuclide mix that is part of that 2 percent ratio, 2 percent enriched uranium ratio. And on the next page, page 142, you see two others: the 12 percent ten-year-old fuel-grade plutonium ratios and the last, the third one is the natural thorium series. And using those three options, I guess the dose reconstruction makes the decision as to which one will give you the highest potential exposure and the highest PoC.

And in this particular case, the choice was the second one, the 12 percent ten-year-old fuel grade plutonium ratios.

And one of the things that -- this is just a sideline issue -- in looking at that table, you'll realize there are a total of four different radionuclides that are identified. The first one is Pu-238, the second is Pu-239, Pu-241, and americium-241. One of the first things I did sort of have to question is what point of identifying is the plutonium-241 as an alpha emitter? Because it's And, of course, it's very important not. because it turns out that has the highest value here. When you look at the right-hand side, it's 14.201.

And it's very difficult to really understand what's going on here, so it took me quite a long time. And what it really comes down to is something that I will explain. If you go to page 143, the next page, John, that particular page shows three tables, and that is

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really based on dose, I mean on air sampling data that were accumulated, general sampling data that would have corresponded to intakes of gross alpha emitters. And if you were, for instance, an operator, a general laborer, you would have obviously been eligible the consideration of the for inhalation quantity that you see on the top of the page that involves, during the operational years of '71 - '72, an intake of 965.121 dpm per day.

If you were, on the other hand, someone who was less likely to be exposed to such values, it would qualify for the supervisor. And in this case, the claimant was identified as likely to have been in that category.

And the important number here is to identify the 482.561 dpm per day, and that is gross alpha. Now, obviously, since you have three options for radionuclide mixes, and these are strictly the values, the 482.561 as alpha intakes per day, and now you have to assume what

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1	is the mix of those alpha emitters based on
2	which of the three. In this case, as I said,
3	it's the 12 percent ten-year-old fuel-grade
4	plutonium that was identified in the previous
5	page, on page
6	MEMBER MUNN: Hans, can you wait
7	just a moment, please?
8	DR. H. BEHLING: Okay.
9	MEMBER MUNN: I'm hearing Hans in
10	sound waves that are very difficult to
11	understand. Is it my telephone system?
12	CHAIRMAN KOTELCHUCK: You know,
12 13	CHAIRMAN KOTELCHUCK: You know, I'm hearing the same thing. I was thinking,
13	I'm hearing the same thing. I was thinking,
13 14	I'm hearing the same thing. I was thinking, although I'm chairing, I'm thinking let's see
13 14 15	I'm hearing the same thing. I was thinking, although I'm chairing, I'm thinking let's see if we can just switch off while he's talking.
13 14 15 16	I'm hearing the same thing. I was thinking, although I'm chairing, I'm thinking let's see if we can just switch off while he's talking.  DR. H. BEHLING: Let me do one thing
13 14 15 16 17	I'm hearing the same thing. I was thinking, although I'm chairing, I'm thinking let's see if we can just switch off while he's talking.  DR. H. BEHLING: Let me do one thing more. I just disconnected my headphone, which
13 14 15 16 17 18	I'm hearing the same thing. I was thinking, although I'm chairing, I'm thinking let's see if we can just switch off while he's talking.  DR. H. BEHLING: Let me do one thing more. I just disconnected my headphone, which may be a problem.
13 14 15 16 17 18	I'm hearing the same thing. I was thinking, although I'm chairing, I'm thinking let's see if we can just switch off while he's talking.  DR. H. BEHLING: Let me do one thing more. I just disconnected my headphone, which may be a problem.  MR. KATZ: Actually, your voice

just mute your phone because there's a lot of background noise. So \*6 to mute your phone if you don't have a mute button. There you go.

MEMBER CLAWSON: Hey, Ted, just to let you know, too, when it comes to questions, I still don't have my computer back. So I'm not able to see most of this stuff, so if there's some oddball questions that come from me, besides my normal ones, it's because I don't have my computer. So I may need a little bit more information, okay?

MR. KATZ: Okay. Thanks, Brad.

DR. H. BEHLING: Okay. For those who may not have access to the computer, as Brad does [not], just bear with me because, in the end, what we're going to do is come full circle, so this issue will be resolved. I just wanted to explain what the genesis of this was and how I came to that conclusion as to the fact that it might be a generic mistake that may not only impact this particular dose reconstruction but all others that involve the Westinghouse

facility. And I think this is what prompted this whole issue becoming an issue that was obviously put in the forefront for discussion.

Anyway, let's go back here. I just said before, when you start out with the assumption that this claimant was a supervisor, he was exposed to 482 dpm per day of alpha, dose Then you have to now decide what alpha. contributions were the different radionuclides that were part of the particular mix that we'=re talking about, the 12-percent ten-year-old fuel-grade plutonium. And as I said, there are only three alpha emitters: Pu-238, 239, and And now you have to decide how americium. you'=re going to separate those out in terms of what fraction of the 482 dpm per day was contributed by each of those three alpha emitters, Pu-238, 239, and [americium-]241. And you can obviously do that by a quick calculation by multiplying the 0.117, which is Pu-238, times 482 dpm per day, and understand what fraction of that total gross alpha was

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contributed by each of those.

And I did all those things. And when I tried to reproduce the number, I came up with a value that was significantly lower than what NIOSH had introduced into the CADW data, which is also included in my review of this case.

And as it turns out, my assessment turned out to be considerably lower. So, again, this is an issue here that would probably not cause a major heartburn because we're overestimating. This error turned out to be an overestimate, as opposed to an under. We're always more concerned when we underestimate a claimant's intake of dose.

Anyway, so what it comes down to, I came up with the conclusion that for the two years, '71 - '72 as an example, the intake would have corresponded to 2010 becquerels per year.

NIOSH, in looking at the CADW data, had estimated 2932 becquerels per year. And I tried to reconcile the difference, and I

realized, only after the fact, that that difference is due to the fact that NIOSH had assigned that daily intake not for 260 days out of the year but for 365 days a year. turns out to be an error because, if you look at page -- and I'm going to ask John to turn to And on that page, the paragraph page 142. starting with air monitoring results are reporting both units of microcuries milliliter of air and dpm per cubic meter. So if you go further down, there is a sentence that starts out with: A"A daily weighted average was established based on the breathing rate of 9.6 cubic meters per day for 250 working days per year."

And on that basis, I realized that the error involved that I identified here was the use of 365 days instead of 250 days. And only this morning, minutes before we went on air, I received NIOSH's response to that particular finding, and I'll just read it to you if you don=t have it.

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NIOSH's response states that -- and I'll just skip the first couple of -- well, I'll read the whole NIOSH response. believes the values given in the methodology are correct for intake for workdays for a 250-day work year. However, when these values were entered into the CADW, the calculation applied these intake values for 365 calendar The assignment of the intake days per year. rate is based on 250 workdays, as a 365 calendar workday-based intake did result in an overestimate of the correct exposure." So I think we've resolved -- NIOSH has accepted the fact that there was some error introduced into CADW. The numbers that I just cited are slightly higher than they should be. And based on what I gather is that this will be corrected. So as far as I'm concerned, this

And if I have any other comment, I would like to at least draw attention to the fact that, since, again, we're on page 142, when

issue has been resolved.

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you have, for instance, the third option of natural uranium, I'm sorry, natural thorium ratios, again, there are two radionuclides in that table: actinium-228 and radium-228. And for each of those, there is the issue of alpha. They're not alpha emitters.

The only three alpha emitters on that particular table are thorium-228, 232, and americium. And so they all have to add up to 100 percent, and, as you see here, the constant ratios of 0.333 would add to the value of 1.66, which is obviously incorrect. We really want to apportion the intake as defined in that table -- in this case, the 482 dpm per day -three equally-divided ratios contributed by thorium-228, 232, and radium-224. And so that should be also corrected in the revision to that particular document. just leads Ιt to misunderstanding, and it's just an easy fix.

MEMBER MUNN: This is Wanda. I misunderstood what you just said about that last table we're looking at. I at first

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thought you said the two radionuclides of 1 thorium and actinium and then you instead, when 2 3 you were speaking last, you said the two radionuclides were the two thorium isotopes and 4 radium-224. 5 Yes, those are the 6 DR. H. BEHLING: 7 three alpha emitters that have to be used to estimate the quantity of the - we're dealing 8 with a supervisor. Four hundred eight-two dpm 9 10 per day would have to be split into three equal 11 I don't have my calculator, but you can parts. 12 obviously --13 MEMBER MUNN: Oh, yes --And the other 14 DR. H. BEHLING: three are not alpha emitters. 15 16 MEMBER MUNN: Okay. 17 DR. H. BEHLING: Actinium-228 and radium-228 are not alpha emitters, and they 18 should be included in the dose reconstruction 19 but they should not be part of the gross alpha 20 numbers that we're trying to use in separating 21

which alpha emitters contributes what to that

1	gross alpha intake.
2	MEMBER MUNN: Alright, alright.
3	MR. SHARFI: This is Mutty Sharfi.
4	I can explain the table for you if you'd like
5	and why I felt that way.
6	DR. H. BEHLING: Okay.
7	MR. SHARFI: Since we're applying
8	the ratios to a gross alpha result, you're
9	correct that three of them are alphas and two
10	of them are beta emitters, but we're just using
11	the ratio to determine intakes. So there are
12	five radionuclides that you're going to end up
13	assigning. So when you sum the three alphas,
14	you do get 100 percent. But because there are
15	five radionuclides, the total fraction will end
16	up being, when you talk about total intake to
17	the gross alpha intake, the total intake is
18	actually 166 percent since all the
19	radionuclides are in equilibrium.
20	DR. H. BEHLING: Yes, I understand
21	
22	MR. SHARFI: So we are so it's

1	really an application issue. The table wasn't
2	to imply that they're alpha emitters, just how
3	the ratio gets applied to the gross alpha
4	result.
5	DR. H. BEHLING: Yes. In this
6	case, and I totally agree with you, the third
7	table would have been used. When I looked at
8	the ten-year-old fuel grade and I saw the large
9	value for plutonium-241 as an alpha emitter, it
10	kind of just threw me a curve ball, that's all.
11	MR. SHARFI: Yes, yes. All the
12	tables, yes, were all tagged to the gross alpha
13	result because that's how, yes, that's how the
14	information, that's how the intake is
15	originally designed.
16	DR. H. BEHLING: Yes. It's just a
17	nit. I'm not saying that this is an issue here
18	that needs to be belabored, but it's just very
19	confusing when you first try to get into NIOSH's
20	head in saying how did they do this, how did they
21	come up with those numbers, and

MR. SHARFI: Yes. I think the

confusion is because, as an AWE, we deal more with gross alpha samples. And in most cases, when we talk about DR, we're talking about DOE sites that we're tagging to, like plutonium bioassay. So it's not a tag to a gross alpha sample but a plutonium alpha or a thorium alpha, you know, specific. And since these are not radionuclide-specific intakes that we're tagging them to, it's probably unusual compared to other claims that we usually look at.

DR. H. BEHLING: Okay. The only last question I think you should answer for the people on the line here, was this issue of applying the intake values for 365 instead of 250 days, was that something that we can reasonably assume also applied to the other cases that were done, or was this unique to this particular DR, this error of applying 365 days as opposed 250 days a year?

MR. SHARFI: I would have to generically go look at individual claims to probably answer that because, generically, I

1	mean, when we do TBDs, you develop
2	methodologies for particular claims. And I'll
3	let Grady talk more about that.
4	But I can't specifically say
5	whether or not we have gone and corrected to be
6	more specific in these tables. So if this
7	information is used for a future claim, so that
8	it now says per calendar day or per workday, so
9	that if that confusion, it's made sure that we
10	don't have that again.
11	DR. H. BEHLING: Yes, because
12	that's really what triggered this concern that
13	says, if this is not the only case where this
14	error occurred, then this is why it was an issue
15	that prompted this memo. If it is the only
16	case, then it's obviously a minor problem.
17	CHAIRMAN KOTELCHUCK: Except for
18	that case.
19	DR. H. BEHLING: Yes. Again
20	MR. SHARFI: Again, this does
21	result, as Hans cited, to an overestimate, not
22	an underestimate.

1 DR. H. BEHLING: Yes.

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MS. K. BEHLING: This is Kathy Behling. I believe this also, though, brings up a bigger or more generic issue in my mind. The fact that we were really not aware of this dose reconstruction methodology, and I realize that there are these documents out there and they are now showing up in the claim files, I think it's important again, when the --Subcommittee selects cases, there may be these smaller sites out there that don't have Site Profiles where these dose reconstruction methodology documents are in the files, and we never get to see them unless there is a case that we're assigned.

So I don't know how many of these types of documents exist out there, but maybe NIOSH could shed some light on that because it's not something that we will typically review or see unless it's a case that's selected by the Subcommittee.

MR. CALHOUN: This is Grady.

We've done that intentionally because, as you know, we have hundreds of covered sites. And the thought is that those sites without an approved TBD, the reconstruction itself needs to be detailed enough to judge on its own. It needs to be able to stand alone.

So there are sites with these what we call methodologies out there. We have no intent of making those approved TBD documents. In this case in particular, you think we've had less than a hundred cases in the 13 years we've been in existence. And there's a lot of other sites that we've had fewer than that, so those do exist. But the intent is that the DR is detailed enough, the dose reconstruction is detailed enough it can stand alone.

So that's what we need to be looking at is the actual dose reconstruction. And if it's not detailed enough to verify that the answers are okay, then we need to revise how we do the dose reconstruction, rather than making a full-blown approved TBD for every site we do

_	a dobe recomberded for
2	MS. K. BEHLING: Okay, fine.
3	Thank you.
4	DR. H. BEHLING: And just to recap
5	for everyone, including Dave Kotelchuck, I
6	don't think anything really extravagant needs
7	to be done here, even if this is not an isolated
8	case, simply because if this even applies to
9	other previous cases that have been
10	reconstructed then those estimates would be
11	higher than would normally result if you used
12	the correct 250 days. And if I recall, we never
13	correct an error that is on a claimant's side
14	or favor, so, at this point, I think we can put
15	this to rest.
16	CHAIRMAN KOTELCHUCK: Okay. And,
17	NIOSH folks, do you agree?
18	MR. CALHOUN: Yes, I'm good with
19	it. How about Scott and Mike?
20	MR. SHARFI: I agree that the
21	agreed-upon correction would only reduce the
22	dose, not increase it.

a dose reconstruction for.

1	CHAIRMAN KOTELCHUCK: Right,
2	right.
3	MR. SIEBERT: And this is Scott.
4	The other thing to note is this will come up
5	again when we deal with the next set because
6	this is SC&A 434. And at that point, we'll
7	actually respond to all actually, we already
8	did respond to all the findings in that document
9	that was sent out this morning, but I believe
10	we'll cover that when we hit that in the next
11	groupings that we deal with the 14th through
12	18th set.
13	CHAIRMAN KOTELCHUCK: Very good.
14	Subcommittee Members, anybody have any further
15	comment or concern?
16	MEMBER MUNN: This is Wanda. Not
17	here. No, I think Hans' explanation was well
18	received here.
19	CHAIRMAN KOTELCHUCK: Okay. And I
20	don't have any further comments. So I think
21	that this issue is resolved, and we're ready to
22	move on to our case reviews.

1	Okay. Now, actually, let us go
2	back to the single case that's sitting out in
3	set 9, where we were with 270 no.
4	MR. FARVER: This is Doug. It's
5	185
6	CHAIRMAN KOTELCHUCK: 185.7,
7	right? We needed a report, or there was some
8	can we put that on the screen? And then,
9	hopefully, it will be resolved today.
10	MR. FARVER: Dave, this is Doug.
11	Probably the best thing we could do is put up,
12	we have a matrix of the Huntington Pilot Plant
13	issues, and it contains the NIOSH responses and
14	what our reply to those are.
15	CHAIRMAN KOTELCHUCK: Alright.
16	MR. FARVER: I think John Stiver is
17	going to get that on the screen. Just to recap,
18	we had that one outstanding finding from the
19	Huntington Pilot Plant, and it all stemmed back
20	to a report that SC&A wrote reviewing the
21	revised Site Profile.

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And

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report, we had

identified six findings. Two of the findings 1 were closed out with the 185.6 discussion, 2 3 which was a finding before this one. That left four findings that NIOSH was going to go back 4 5 and read the report and respond to. 6 So that we could capture all this, 7 we just put this in a matrix. And that way, we have all the information in one piece. 8 CHAIRMAN KOTELCHUCK: 9 Okay. 10 MR. FARVER: They're on the screen. CHAIRMAN KOTELCHUCK: It is on the 11 12 screen, findings one and two. 13 MR. FARVER: Okay. These were issues that Steve Marschke identified based on 14 a review of the Site Profile. So the first one 15 deals with contaminants that are in the TBD, and 16 17 NIOSH's response was that they're revising the TBD and they will include the contaminants that 18 we're concerned about. it's a 19 So TBD modification, and, as long as they modify the 20 TBD, we really don't have a big concern about 21

just

including

this because they're

information that we would like to have them 1 include. 2 3 CHAIRMAN KOTELCHUCK: Good. Finding two is very 4 MR. FARVER: similar. with 5 Ιt has to do the 6 uranium-specific activity, and there were some rounding issues where they rounded the two 7 significant figures. And all we're asking --8 and they said they're going to modify the TBD. 9 10 We just want to make sure the TBD is specific 11 and states that those results are rounded to 12 significant figures, and it also quotes the DOE standard, STD-1136 as a reference. 13 So we really don't have any concerns 14 as long as they modify the TBD to be specific 15 to say that they rounded it. You know, pretty 16 17 much put in the TBD what they put in their response is what it comes down to. So then we'd 18 have no concerns, and we can close that issue. 19 20 That's the short version. 21 CHAIRMAN KOTELCHUCK: Okay. 22 MR. FARVER: And the third finding

1	on the next page, we had identified a unit
2	conversion error. There really wasn't a
3	conversion error, but the labeling of the
4	columns one of the columns was
5	incorrect. So they're going to correct it in
6	the TBD revision, so we have no concerns with
7	that. I mean, we agree that's good.
8	MR. CALHOUN: Doug, this is Grady.
9	I'm assuming here that you don't have a copy of
10	the revised TBD. Is that a true statement?
11	MR. FARVER: Not [one] that it's
12	been revised to include these issues.
13	MR. CALHOUN: On 2/21, the 2/21
14	2014 revision?
15	MR. FARVER: Oh, probably not.
16	MR. CALHOUN: Okay. I'm going to
17	shoot that off to you. I don't expect you to
18	do anything with it right this moment, but I
19	should have sent you that.
20	MR. FARVER: Grady, have all these
21	changes been incorporated?
22	MR. CALHOUN: That's what I

I believe that is the case. 1 believe. MR. FARVER: Okay. 2 3 CALHOUN: But I understand you've got to take a look at it so --4 5 MR. FARVER: Okay. Thank you. 6 And, finally, finding four, which is on page 7 four, has to do with the energy spectrum or energy fractions. And they identified that 8 the data that they used was from the National 9 10 Nuclear Center, which is а little Data 11 different than what we were assuming. 12 have no problems with their response. And really this was just to clear up 13 We had talked about them before, 14 some issues. but we wanted to get everything down in writing 15 and all in one place. These were the four 16 outstanding issues. And then, if you want to 17 go on, you can see that the findings five and 18 six have already been addressed at our August 19 meeting of last year. So as long as those TBD 20 changes are made, that should wrap up the 21

Huntington Pilot Plant issues and that should

1	wrap up our 9th set matrix.
2	MEMBER RICHARDSON: Doug, this is
3	David Richardson. Could you describe number
4	four?
5	MR. FARVER: Number four.
6	MEMBER RICHARDSON: It had to do
7	with the dose distributions, assumption of
8	energies.
9	MR. FARVER: Okay. What I'm going
10	to do is look at Steve's report and get a more
11	detailed answer. It looks like we were just
12	trying to identify what gamma spectrum they
13	used. It was not identified in the document
14	where the data came from.
15	MEMBER RICHARDSON: Right.
16	Because this was I mean, there are several
17	issues here, but one of the issues is I wasn't
18	aware before of there being parameters in a dose
19	reconstruction which are dynamic. That is, if
20	today I were to go and try and figure out those
21	parameters, they would not be the ones which

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1	reconstruction. And unless somebody is
2	archiving the history of those changes that
3	they evolve in this data center, there's not a
4	clear reference table. It would have to be
5	accessing a website on a given date. Is that
6	how this is? I've just not seen parameters like
7	that before.
8	MR. FARVER: I really don't have a
9	good answer for you, David.
10	MEMBER RICHARDSON: My
11	understanding, you used a library, a, quote,
12	frozen library. That's a lookup table which
13	you were going to use as a reference that might
14	be in a Technical Basis Document, and the
15	response was we checked a different library
16	which is dynamic and is online.
17	MR. FARVER: I understand your
18	point. I don't have a good answer. Grady, do
19	you have any input?
20	MR. CALHOUN: Basically, what we do
21	when we determine what the energies are that
22	we're going to use, it's based on what materials

were present during that time frame. I don't know exactly what, I don't know exactly what that one is. It's not that they change, and I don't know if this is a library like they're talking about whole body count, a software library. That's what it sounds like. I'll have to go back and look and see the particulars on this case.

But they don't vary from day to day like you're saying. It's just something based on, if it's a nuclear library that's included in the software, that's what is used to detect photons and whole body count. I don't know if that's the case in this one, but in other of these, when we're assigned photon dose, you base the energies on the materials that are present because there is a breakdown of the different energy spectra that we see, and we have to bend those to put them into IREP.

MEMBER RICHARDSON: Those are sort of physical properties, which I would imagine are invariant, and so it's difficult for me to

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MR. CALHOUN: That's the fact, and that's how we'll use it. However, if this is a nuclear library, and I don't know that, but based on this comment, when a whole body count exists in the setup or any kind of gamma spectroscopy is set up, there's a library that's in that software that's used to detect certain energy level photons. I just don't know, based on this information here right in front of me, if that's the case or not.

MR. FARVER: David, a little bit more information. It looks like we were trying MicroShield to check the use calculations that NIOSH performed and were explained in Appendix A of the Site Profile. why having trouble That's we're with MicroShield because it's a different library. We're having difficulty matching the dose were assuming a different rates, and we spectrum and we still couldn't quite match.

MR. CALHOUN: How far off were

1	these? Is it 50/50 to 70/30?
2	MR. FARVER: I don't know.
3	MR. CALHOUN: So, basically, what
4	this comes down to is you guys used a little bit
5	different library than we used.
6	MR. FARVER: Correct.
7	MR. CALHOUN: Oh, okay.
8	MEMBER RICHARDSON: Okay. I think
9	I'm seeing the issue. And if you have the
10	library and it's assumed to be fixed, in
11	principle, one could go back and find it, and
12	it's not the issue that there was one which was
13	frozen and one which is dynamic. There are two
14	different libraries based on two different
15	software systems.
16	MR. FARVER: I believe that is the
17	case.
18	MR. CALHOUN: That sounds like my
19	understanding, too, now from this. And unless
20	somehow ours is wrong, then it's just a
21	difference that we picked that they picked
22	a different library to evaluate what we had done

already. 1 But if, for CHAIRMAN KOTELCHUCK: 2 3 any reason, anybody came back years from now and wanted to take a look at that calculation, you 4 5 would have the information in what you did about 6 which library you used? We would for sure. 7 MR. CALHOUN: And so really the issue is whether or not the, 8 quote, library that we used was incorrect, and 9 obviously we don't believe it is. 10 CHAIRMAN 11 KOTELCHUCK: Right, 12 right. Okay. But yours, which is to say NIOSH's calculations do say what library was 13 used, and that can be looked at in the future, 14 and if, for any reason, somebody ever were to 15 think it was wrong it could be changed or 16 So it doesn't matter whether SC&A... 17 modified. I mean it matters whether your documentation is 18 not so much reproducible but is discoverable, 19 and it is apparently, right? 20 MR. CALHOUN: I believe that's the 21

case, yes.

1	CHAIRMAN KOTELCHUCK: So, David
2	Richardson, that should resolve it, shouldn't
3	it?
4	MEMBER RICHARDSON: Yes, that's
5	clearer to me. Thank you.
6	CHAIRMAN KOTELCHUCK: Yes, okay.
7	Thank you for asking.
8	MEMBER MUNN: Well, I'm glad it's
9	more clear to David than it is to me. This is
10	Wanda. And what the puzzler, from my point of
11	view, is why do two perfectly reliable,
12	supposedly, libraries have different values
13	for the same spectra? It doesn't seem to
14	follow that that would be expected, does it?
15	Am I missing something?
16	MR. CALHOUN: No, you're not.
17	This is Grady. I don't think you're missing
18	anything at all. I think that we need more
19	information here to look into what assumptions
20	were made with the different percentages of
21	what radionuclides were assumed on our side

versus your side because you're absolutely

right: the energy coming off of a given radionuclide is pretty much constant, for the most part.

So I think that we'd have to look at what the assumptions were with both cases if we need to pursue this any further. And I guess it really depends on how wrong the SC&A guys think we are.

MEMBER MUNN: Well, based only on the information that we have for a person like me who knows absolutely nothing about even the existence of the libraries, much less their content, it does create a puzzler because one would assume that you could expect the same information about spectra from one reliable library to another.

MEMBER RICHARDSON: Yes. Wanda, so was I -- that part of the issue, kind of one of them has to do with process and being able to document and reproduce what was done. The other part of the content of these two software packages and why there's a kind of disagreement

1	between this, that isn't obvious. My
2	experience in the past was if we were to hold
3	this finding until we had resolved that I
4	mean, I don't know. In some cases, it would be
5	to try and reconcile two different software
6	programs. It could be potentially a lot of
7	effort.
8	MEMBER MUNN: Oh, I'm not concerned
9	with the programs itself. I'm concerned, as I
10	think I said, with the libraries.
11	MEMBER RICHARDSON: Right, right.
12	MEMBER MUNN: It's strange to me
13	that using a different library would give you
14	a different result.
15	MR. FARVER: Well, I think it's a
16	matter of just using different assumptions.
17	CHAIRMAN KOTELCHUCK: Different
18	assumptions about what?
19	MEMBER MUNN: This is the level of
20	detail which is probably beyond our purview,
21	but it is, you can understand why it raises the
22	question.

1 MR. FARVER: Oh, I understand. looks like we're trying to model a 20-gallon 2 3 drum and the dose rates that are coming off --4 MEMBER MUNN: Yes, Ι objection to closing this item. 5 6 CHAIRMAN KOTELCHUCK: Yes. Well, this is 7 MEMBER CLAWSON: I understand where we're going, but, Brad. 8 Doug and everybody, I thought one of the things 9 we'd be able to, 20 years from now or whatever 10 11 else, be able to come back and figure out how 12 these things were done, and we can't even figure 13 that out for sure today. How can we say that 14 that is correct? 15 MR. CALHOUN: That's not true, This is Grady. We certainly can show 16 Brad. how we did ours. The question is they used a 17 know, when they 18 different, you qo modeling, someone else goes in to try to model 19 a different situation. You know, there are so 20 many parameters that come into effect that 21

could affect the energy distribution.

22

You

know, if you're talking about a drum, you're talking about the density of the drum itself, you're talking about the density of the material in the drum, you're talking about the type and quantity of the material in the drum.

So I think, basically, we have our justification and it's readily reproducible. The issue is that when an evaluation of our approach was done, a different tool was used. So it's not really an issue with our process being unreproducible, because I believe ours is.

MR. STIVER: This is John Stiver.

Maybe I can jump in. I said something similar to this in the back and forth that went on with GSI when a lot of the results and the spectral differences depend on, as Grady said, on the different assumptions about the density of material and the types of material and so forth.

And what we did to kind of help resolve that was just have NIOSH send us the input file that they used and we could check the assumptions. So at

least that way, you know, we had to be on the same page, as opposed to kind of running off and doing a separate analysis with our own assumptions and coming up and trying to resolve all this. Some of those things, you know, maybe some communication up front, at the front end, might help too, it would save us a lot of work later on.

MR. FARVER: Rather than trying to beat this to death right now, let's try to get Steve Marschke on the line -- maybe after lunch -- since he's the one that reviewed the profile and he's the one that reviewed the responses and he's the one that actually wrote the report when he did the model. And he didn't have a concern about it, but we'll try and see if he can give us a quick explanation for that.

CHAIRMAN KOTELCHUCK: That sounds reasonable. So if you can have him come on the line after lunch, let's do that. I'm satisfied that the result is reproducible and could be reproduced at some later time. The fact that

1	data might change, and spectra measurements may
2	result in, I hope, small changes, that does not
3	bother me, as long as the result that is agreed
4	upon is reproducible in the future by other
5	parties.
6	So if we want to, let's go on and
7	let's come back to that finding with Steve
8	later. Let's go on to finding five.
9	MR. FARVER: Finding five we
10	already addressed in a previous meeting and
11	closed that issue.
12	CHAIRMAN KOTELCHUCK: Okay.
13	MR. FARVER: Same with finding six.
14	CHAIRMAN KOTELCHUCK: Good. And
15	that's the last finding?
16	MR. FARVER: Yes. That's the last
17	one for that report, yes.
18	CHAIRMAN KOTELCHUCK: Okay. So we
19	have that one, finding four, which we will come
20	back to later with Steve.
21	MR. FARVER: We won't close out the
22	9th set yet. We'll let it hang on a while.

1	CHAIRMAN KOTELCHUCK: Well, we are
2	going to do our very best to finish up 9 and move
3	on to 10 and 13 and try our best to get those
4	finished.
5	So let us go on to sets 10 through
6	13. Doug, you sent out the matrices, and we had
7	the Portsmouth and Paducah. There was one open
8	case that I saw, and then we had Oak Ridge,
9	Hanford, and remaining sites.
10	Do we want to start off with the
11	Portsmouth/Paducah, the gas diffusion plants?
12	MR. FARVER: We'll go ahead and do
13	that. That's in the Paducah/Portsmouth
14	district, and it's finding 273.2.
15	CHAIRMAN KOTELCHUCK: That's
16	right.
17	MR. FARVER: And this talks about
18	the dosimeter correction factor was being used
19	for missed proton dose.
20	CHAIRMAN KOTELCHUCK: We don't
21	have anything up on the screen. You'll put it
22	up on the screen? Okay.

1	MR. FARVER: And the action was
2	NIOSH was going to write a White Paper about,
3	you know, do they or do they not need it.
4	MR. STIVER: Hey, Doug, what was
5	the finding number?
6	MR. FARVER: 273.2 on page two.
7	MR. STIVER: Okay, alright. Here
8	we go.
9	CHAIRMAN KOTELCHUCK: Here we go.
10	MR. FARVER: If you recall,
11	Portsmouth was the only place that they used a
12	dosimeter correction factor for missed dose.
13	Normally, it's just applied to the dosimeter
14	dose and not the missed dose, and this is why
15	we questioned it in our findings.
16	And I believe for this it's not
17	really what you would call a dosimeter
18	correction factor. It's a how do you say?
19	It's a correction factor, but it's not
20	correcting like you would for the other site.
21	I'm trying to find NIOSH's
22	response, but I'm having some difficulty.

1	Scott, can you come in and do you have any
2	insight on this?
3	MR. SMITH: This is Matt Smith. I
4	can talk to the response.
5	MR. FARVER: Thanks, Matt.
6	MR. SIEBERT: Do you guys have the
7	response? This is Scott. I'm just verifying
8	you have the response and the White Paper.
9	MR. FARVER: Yes.
10	MR. SIEBERT: Okay, thanks.
11	MR. SMITH: Yes, we did a White
12	Paper. That came in a January time frame. As
13	I took a look at this, it looks like the TBD
14	author was doing some referencing to the
15	Savannah River Site Profile, which was one of
16	the first Site Profiles put together on this
17	project. And what this factor is is basically
18	an approach to taking the dose that was measured
19	and converting it to what we would call modern
20	Hp10 dose.
21	In all reality, that kind of
22	approach does not need to be taken because we

have the DCF, the dose conversion factors, available out of DCAS-IG-001, and that lets us either use an exposure DCF for the error where a film dosimeter was used and calibrated but calibrated not on a phantom as you would see in a modern era with a DOELAP type of program. When we're in that era, then we would use the Hp10 DCFs when phantom is used as part of the calibration routine.

The White Paper kind of describes how we took a look at it and we can see where the author is pulling some of the information and data from the SRS TBD. The bottom line is the correction does not really need to be applied, and you can even say the same thing for the Savannah River TBD, which is I know under revision right now as well.

In retrospect, all that need be done really is to apply the proper DCF, again the exposure DCF for an era where film dosimetry is used and then the Hp10 DCF for an era where you have a TLD type of dosimeter that's being

calibrated on a phantom, so the backscatter is being taken into account.

The recommendation of the White Paper was basically to update the Portsmouth TBD to simply go with the recommendation that's already in there regarding the proper DCF values to use and to take out the references to the Savannah River approach.

MS. K. BEHLING: And, Doug, this is Kathy Behling. This was one of the issues that I brought up because I was questioning if perhaps this correction factor should be applied at other sites.

And I did read through this White Paper and I agree with what they have written in here. I didn't realize what the correction factor was, how that was being applied, or what the reason for the correction factor was, if it was an under-response of the TLD or the film badge. But as, Matt had just explained, it is a conversion from a calibration that was done free in the air to now trying to determine the

Hp10 dose. And so I agree with what was written 1 in the White Paper, and I feel comfortable that 2 3 this is not a problem that exists at other facilities or in other TBDs, and I agree that 4 5 this should be changed in the Portsmouth TBD. 6 CHAIRMAN KOTELCHUCK: Sounds like 7 subcommittee agreement. Any comments by members? 8 9 MEMBER RICHARDSON: This is David 10 Richardson. follow-up, there So to 11 different corrections. I quess maybe 12 issue is how the correction factor is being used At Savannah River and other sites, there 13 here. 14 issues about under-response were or over-response of different types of dosimeters 15 to different geometries of exposure, different 16 17 energies of exposure. And on top of that, there was talking about under-response 18 estimation of 19 over-response to the what 20 And one of those quantities was quantity. Hp10. 21

The discussion that's happening

here is you're just talking about the distinction between estimates of dose in air to Hp10 but not about the performance of the dosimeters. That's being wrapped up in a different DCF that is being taken into account or is [it] not?

MR. SMITH: The former is a true statement, and what Kathy said based off of what I said is what the White Paper is putting forth. With respect to the dosimeter performance in the Portsmouth environment, the TBD itself does state that the Portsmouth two element film would overall be favorable to claimants and no corrections are needed for the response of the dosimetry to their radiation work environment.

With respect to Portsmouth, what we see with that two element film dosimeter, you know, basically it's the similar design [to] that we found at X-10 with Oak Ridge, and it's described in that TBD, as well. As we get down to around 100 keV, you even start to see some over-response of that dosimeter.

The bottom line is only the DCFs 1 that help us convert from exposure free-air or 2 3 in the sense when the dosimeter was calibrated on a phantom, which is an Hp10 situation, 4 5 convert those to organ dose. In the case of 6 Portsmouth, no other correction factor was 7 warranted based on any kind of energy response. When you see the paragraph in the 8 Portsmouth TBD talking about under-response in 9 terms of calibration, what's being discussed 10 there is the response to backscatter from a 11 12 phantom that may or may not have been used. it's also discussed in the Savannah River 13 14 document as well. CHAIRMAN KOTELCHUCK: 15 Comments? MEMBER CLAWSON: This is Brad. 16 Τ 17 don't have any. This is Doug. 18 MR. FARVER: Just so 19 you know, I've captured this into the matrix [with] some comment or some of the quotes from 20 the conclusion of their White Paper to explain 21

the situation, so it's better explained in that

1	matrix.
2	CHAIRMAN KOTELCHUCK: Sounds like
3	we're ready to close this.
4	MEMBER MUNN: Sounds like it to me.
5	CHAIRMAN KOTELCHUCK: Okay. Then
6	I believe this one is closed, and I think that
7	is the only one that was open in
8	Portsmouth/Paducah; is that correct?
9	MR. FARVER: I believe so. So that
10	will close out that
11	CHAIRMAN KOTELCHUCK: The gaseous
12	diffusion plants. Okay.
13	MEMBER MUNN: That does this set,
14	right?
15	MR. FARVER: Yes.
16	CHAIRMAN KOTELCHUCK: Yes.
17	MEMBER MUNN: Great.
18	CHAIRMAN KOTELCHUCK: Well, we
19	have Oak Ridge and Hanford now, and do we want
20	to start with Oak Ridge for no special reason?
21	They're both about the same
22	MR. SIEBERT: This is Scott. I'd

1	just like to point out Hanford actually only
2	has, the Fernald/Hanford only has one
3	outstanding finding, so that might be wise to
4	get that one out of the way.
5	CHAIRMAN KOTELCHUCK: Okay, fine.
6	Good, good. Actually, I have some questions
7	about some of them, but that's fine. Let's go
8	to Hanford. Was it 242.1? Is that the one
9	that
10	MR. FARVER: That is correct.
11	CHAIRMAN KOTELCHUCK: is
12	outstanding?
13	MR. FARVER: John, that's page 17,
14	I believe. The action was that NIOSH will
15	investigate the extent of the workbook issues.
16	This goes back to some where we believed it was
17	summing up a column, and the dose that was
18	omitted was outside the range of the sum.
19	CHAIRMAN KOTELCHUCK: Right.
20	MR. FARVER: And so we asked them to
21	check the workbooks and then see if that is an
22	issue.

MR. SIEBERT: Okay. And this is Scott. I'll go ahead and address that. We have been working diligently on that since the last meeting because it involves going through a lot of workbooks. I'm going to be writing up the report for NIOSH to review probably in the next couple of weeks, so we probably won't get this closed out today but I want to let you know where we are in the process.

When we went back and looked at the tool in question, as Doug said, there were two columns where the summation did not necessarily go far enough to include all the information in it. Fortunately, those summations were only dealing with the dosimeter error that we need for full best-estimate calculations, so it would be a subset of the number of times we had to run the tool that would actually be affected by it. If we use under- or over-estimates in the tool, it does not use that portion of the tool, so any error in that area would not impact the claim.

So that was helpful to figure out.

CHAIRMAN KOTELCHUCK: Okay.

MR. SIEBERT: So what we did is we looked back and it really affects the years prior to 1958 only. There are times where, up until approximately 1950, the site used weekly badging, and the tool was set up to look for weekly badging. But if a person had multiple facilities, they may have had more than 52 badges, and that's what happened in this case.

So we've looked back at that. So pre-1950, that can be an issue. From about '51 to '57, they went to bi-weekly, so we would expect 26. The tool was still set up to look for 52, so, in most cases, that would still be okay because people generally weren't getting twice as many dosimeters. But we included looking at that as well in this review.

And then once we hit mid 1957 is when they went to monthly badging. So if a person has monthly badging, we were still looking for at least 52 rows. In some cases, we were

looking for 200 and some rows.

So once we hit the monthly badging period, we were very comfortable that we were summing all the doses that were needed for that error calculation.

So we went back and we looked at all the claims that were done prior to, that had employment prior to 1958 and looked at conversion of the tool that updated to reflect enough rows to ensure that we're catching everything. We had that updated in 2010 when we went to the full Vose calculations instead of the old Crystal Ball [calculation]. So we looked at all of the tools that we used prior to that point.

I won't get into all the specifics.

I know I've already gone a little long, and it's a lot of stuff. But I'm going to be verifying the numbers that we have, so this is not gospel. But we are looking. I think we found six claims where this error affected the total of the error calculation. All but one were already

compensable, so it had no impact. The one that was not compensable, it looks like it missed a difference of approximately 30 millirem. And when we added additional dose into it, it did not impact that claim.

Like I said, I'm going to write all this up and give a lot more verification to ensure it is right before I give it to Grady to go over to the Subcommittee. But I wanted you to know where we were and kind of get an idea that the scope of the issue is not as wide as we had feared earlier on.

CHAIRMAN KOTELCHUCK: Okay. So do we, does the Subcommittee need to look at this when you finish? It's correct that you're going through, you're verifying all the workbooks, which is excellent. Do we need to -- I guess we need to leave it on our matrix just to make sure it was done, although it sounds like you will make corrections, when you find problems you will make corrections, and that is being done.

1	MR. SIEBERT: Well, the tool in
2	question was already updated in 2010. That
3	issue was solved already back at that time
4	frame, so it's not an issue anymore.
5	CHAIRMAN KOTELCHUCK: Yes, right.
6	But you have to check back through previous
7	calculations or reconstructions.
8	MR. SIEBERT: Right. I'm
9	verifying all the work we've done to this point,
10	correct.
11	CHAIRMAN KOTELCHUCK: Sure, okay.
12	MEMBER RICHARDSON: So this is
13	David Richardson. So when you have, when you
14	said the tool was corrected, you mean that you
15	maybe I'm not understanding the relationship
16	of the tool to the dosimetry information. The
17	correction is that you've updated the
18	period-specific assumptions about the number
19	of badgings in different time periods?
20	MR. SIEBERT: Indirectly, yes.
21	What we corrected was the issue was it
22	would only be looking for 52 lines of data when

1	there was weekly badging. If an individual had
2	an additional badge if they were in multiple
3	locations and there was a 53rd row, the tool
4	would not be summing that into the error
5	calculation. Now, it would sum it into the
6	dose total, but it would not add it into the
7	error portion, the calculation, which is what
8	we ran into in this claim.
9	So when we updated the tool in 2010
10	sorry. I keep getting a little bit of echo
11	and it's bugging me When we updated that,
12	it actually is looking for over 200 rows of
13	data. So when you run into that additional
14	badging, it is included in the summation for
15	that error calculation.
16	MEMBER RICHARDSON: Okay. So in a
17	calendar year, it's looking for up to 200
18	records for a worker. Is that right?
19	MR. SIEBERT: Correct, correct. I
20	think it's 260, something like that.
21	MEMBER RICHARDSON: And it's doing
22	that in all calendar years, from '43 forward,

1	or does it stop at some point?
2	MR. SIEBERT: No, it's always
3	looking for that.
4	CHAIRMAN KOTELCHUCK: It sounds
5	like you're pretty close to concluding your
6	work.
7	MR. SIEBERT: Correct. It's just
8	verifying what we've got and writing it up. So
9	however the Subcommittee wants to handle it,
10	obviously it's up to you.
11	CHAIRMAN KOTELCHUCK: Right.
12	Well, to the extent that we want to eventually
13	move on to writing our report, we could probably
14	consider this done. But we'll leave it open
14 15	consider this done. But we'll leave it open until, I assume we'll leave it what do others
15	until, I assume we'll leave it what do others
15 16	until, I assume we'll leave it what do others think in the Subcommittee? My feeling is we
15 16 17	until, I assume we'll leave it what do others think in the Subcommittee? My feeling is we have to leave it open just simply as a signal,
15 16 17 18	until, I assume we'll leave it what do others think in the Subcommittee? My feeling is we have to leave it open just simply as a signal, a reminder to us to come back and make sure
15 16 17 18 19	until, I assume we'll leave it what do others think in the Subcommittee? My feeling is we have to leave it open just simply as a signal, a reminder to us to come back and make sure MEMBER CLAWSON: This is Brad. I

1	what was said was done was done and that it's
2	working properly.
3	CHAIRMAN KOTELCHUCK: Okay. So I
4	think we can move on. I believe we can move on.
5	We'll leave it marked, though. We'll leave it
6	marked to come back to. But this is
7	essentially done. Now, do others agree? No
8	problem?
9	I had some questions on Hanford.
10	The number, the cases 319.1 to .4, I was a little
11	unclear what that was all about. It was just,
12	these were a number that were marked right at
13	the end of the Hanford set that you gave us.
14	Could we look at them? Could somebody tell me
15	what those
16	MR. FARVER: I think we already
17	addressed them when we did the Group A findings.
18	And I meant to add the responses in here because
19	they've all been closed out.
20	CHAIRMAN KOTELCHUCK: So we've
21	gone over these before?
22	MR. FARVER: We've gone over these.

1	I was just putting them in here for
2	completeness.
3	CHAIRMAN KOTELCHUCK: Okay.
4	MR. FARVER: They are contained in
5	a Group A matrix, which I have to track down.
6	CHAIRMAN KOTELCHUCK: Right.
7	Now, this is on the last page I believe, the last
8	page of this matrix.
9	MR. FARVER: Right.
LO	CHAIRMAN KOTELCHUCK: Could we
L1	take a look? There we are. Okay. So what
L2	you're saying is we've done them, and I see the
L3	markings for what the problems were. So you're
L4	just going to fill in the rest of the matrix?
L5	MR. FARVER: I'm going to fill in
L6	from the Group A matrix.
L7	CHAIRMAN KOTELCHUCK: Okay.
L8	MR. FARVER: I mean, that's my
L9	thought.
20	CHAIRMAN KOTELCHUCK: Okay. So
21	there's nothing that we have to consider
22	MR. FARVER: No.

1	CHAIRMAN KOTELCHUCK: as a
2	Subcommittee; is that correct?
3	MR. FARVER: That is correct. And
4	when I send this out after this meeting, all
5	that will be filled in.
6	CHAIRMAN KOTELCHUCK: Good.
7	MR. FARVER: I hope.
8	CHAIRMAN KOTELCHUCK: Right. And
9	the suggested action will be closed on all of
10	those four cases.
11	MR. FARVER: Correct.
12	CHAIRMAN KOTELCHUCK: Okay. Then
13	that would be it for Hanford. Any other
14	Hanford issue? Good.
15	It is 11:50 on the East Coast.
16	There was the thought, since we started at
17	10:30, that we might go on until 12:30 and then
18	take a lunch break. I'm open to that. How do
19	other people feel? That is to say to continue
20	on for about a half an hour more?
21	MEMBER CLAWSON: This is Brad.
22	I'm fine with that.

1	MEMBER RICHARDSON: Could we
2	possibly take a five-minute break?
3	CHAIRMAN KOTELCHUCK: Yes.
4	MEMBER CLAWSON: No, David.
5	You've got to sit there and just smile.
6	CHAIRMAN KOTELCHUCK: Okay. We
7	will take a five-minute break. And it's 11:53.
8	Hey, we'll make it seven minutes. We'll be
9	back at noon, and we'll start with Oak Ridge.
10	(Whereupon, the foregoing matter
11	went off the record at 11:53 a.m.
12	and went back on the record at
13	12:00 p.m.)
14	CHAIRMAN KOTELCHUCK: Hello,
15	folks.
16	MEMBER CLAWSON: Hi, Dave. This
17	is Brad.
18	MR. KATZ: Okay. Are we all back?
19	CHAIRMAN KOTELCHUCK: It's noon,
20	and we're ready to restart.
21	MR. KATZ: Super. Take off.
22	CHAIRMAN KOTELCHUCK: Oak Ridge is

1	on the screen, and the first one, I believe, is
2	246.2. I believe that's the first open one. A
3	little bit more. We need to flip up one more
4	page, I think.
5	MR. FARVER: Well, we haven't
6	discussed any of these yet. This is the first
7	time you're looking at these.
8	CHAIRMAN KOTELCHUCK: Okay.
9	MR. FARVER: I know NIOSH sent out
10	some responses yesterday, but I'm not sure
11	which file you're looking at.
12	CHAIRMAN KOTELCHUCK: Well, right
13	now I happen to be disconnected. I assume
14	that, it seems to be trying to
15	MR. STIVER: Doug, the one I've got
16	up is entitled "10th to 13th Oak Ridge Sites
17	SC&A 2014." That's the one you sent over that
18	was posted on the O: drive. You're telling me
19	there's a newer one that we should be looking
20	at?
21	MR. FARVER: Well, they sent one
22	yesterday morning, or yesterday sometime I saw

1	it.
2	MR. STIVER: Well, let's see if I
3	can pull that up.
4	CHAIRMAN KOTELCHUCK: Okay.
5	MR. STIVER: Bear with me one
6	moment here.
7	MR. SIEBERT: This is Scott. It
8	has the same name with an extension of dash
9	NIOSH March 2014 on it.
10	MR. STIVER: Okay. Hang on just a
11	minute.
12	MEMBER MUNN: Oh, there it is.
13	CHAIRMAN KOTELCHUCK: There we
14	are.
15	MEMBER MUNN: Did Board Members get
16	that transmission, also? Should I have that
17	somewhere other than on the screen here?
18	CHAIRMAN KOTELCHUCK: No. I don't
19	recall, I did not see it.
20	MEMBER MUNN: Okay. Just checking
21	to make sure I wasn't missing something.
22	MR. FARVER: Okay. Are we on the

screen?

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CHAIRMAN KOTELCHUCK: We are.

MR. FARVER: 229.1. This is from the K-25 plant. The person worked there from '45 through '53, and it looks like he had a couple of skin cancers.

The first finding has to do with not all occupational medical exams the were accounted for, and it looks like three of the that were in the record were exams considered, and that's what prompted finding. We couldn't figure out why because the medical dose calculations were a little confusing because the ones that were provided to us were not the ones in the final IREP. it's hard to determine why the three exams were And NIOSH says they were omitted and omitted. they should have been evaluated for the dose assessment, but, you know, I'm still concerned why were they omitted to begin with? So I'll come back to NIOSH with why were they omitted, and how will we prevent this?

MR. SIEBERT: This is Scott. This claim was done in 2006 by somebody who's no longer on the project, so I couldn't dig out exactly why they did not do it. I can just state that they did miss assigning those three x-rays.

CHAIRMAN KOTELCHUCK: Okay. And that's just charged up as an error, category C, QA. Well, it is now corrected or in the process of being corrected?

MR. SIEBERT: Well, this is Scott. Even though there wasn't specifically an initial question in SC&A's response, we did look back at the claim, along with the rest of the responses for the rest of this claim, and we did reassess it using the information that We actually had to go back and request we had. actual x-rays from the site to verify whether We did that, and we they were PFGs or PAs. included information, that along with everything else in this claim. And we actually discussed the final outcome in the next

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finding, if you can hold your suspense until the end of the next finding.

CHAIRMAN KOTELCHUCK: Certainly.

MR. FARVER: Okay. Move on to the next finding right now?

CHAIRMAN KOTELCHUCK: Yes.

MR. FARVER: Okay. The next one has to do with the adjustment factor for the employment period. Well, we thought it was too small, and there was no calculation to show how it was calculated or how it was determined in the record. Once again, we just believe the case, you know - that's a QA concern again because it's not something you go back and reproduce. No one is ever going to know why that number was used, and I believe, like I said, they can't really go back and contact the dose reconstructor to figure out why, but it's just one of those things that's just not clear. We feel things like that should get caught when it's reviewed, but I'm not sure there's much you can do about it at the moment.

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1	MR. SIEBERT: And we agree
2	wholeheartedly that it should have been caught
3	in the review. This is the one where I said
4	that we included this change, as well as the
5	x-ray and as well as the other things that are
6	further down that we're discussing. We did
7	re-run it, and the final PoC changed from 46.79
8	to 48.93. So the compensability decision was
9	still the same.
10	CHAIRMAN KOTELCHUCK: Okay.
11	MR. FARVER: Are we going to move to
12	close that issue?
13	CHAIRMAN KOTELCHUCK: Yes.
14	MR. FARVER: Okay. Next was just
15	an observation. It looks like there was
16	another error, a number was transposed.
17	Instead of 0.622 it was 0.662. You know, it's
18	one of those things where we couldn't even
19	determine how this type of error occurred
20	because the workbooks in the employee's file
21	did not contain the same direct input as the

other, as the final IREP table.

MR. SIEBERT: And that one, when I dug a little bit further into this, this one I can answer specifically why it did happen. That time frame, these doses had to be prorated, rather than for a full year of exposure for a partial year of exposure. The values that were coming out of the workbook initially are full years, so we had to do the prorating off to the And it looks like what happened is the side. dose reconstructor did the prorating separately, and when they were entering the information they just mis-keyed one of the entries to be 662 instead of 622.

So what I want to point out on this is we've updated our external workbooks now that the dose reconstructor doesn't have to do that prorating off to the side by hand anymore. You can add in a fraction value, and it will apply that automatically for coworker and ambient doses directly. And we've mitigated that issue. It will not happen from a dose reconstructor point of view anymore.

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MEMBER RICHARDSON: As a question,
The second of th
the change to the workbook and the ability to
have fractions, was that motivated by this
finding or was that independently
MR. SIEBERT: No, we did that years
ago.
MR. FARVER: Okay. The next,
observation number two, assigning the recycled
uranium component doses for '45 through '53
appears to be overly claimant-favorable.
Apparently, the tool has now been corrected to
reproduce specific intake rates and time
periods for coworker uranium, recycled
uranium, and technetium. So I guess this is
one of the early on issues to be corrected or
modified.
That takes care of case 229. 235
CHAIRMAN KOTELCHUCK: Wait a
minute. I thought I was holding my breath for
l I

MR. FARVER: Oh.

1	CHAIRMAN KOTELCHUCK: On 229.
2	Scott, you said I was
3	MR. SIEBERT: Yes, that was the
4	fact that we did run the claim again with
5	everything that we found here, the recycled
6	uranium, the x-rays, and so on. We ran it to
7	see if there was any impact and
8	CHAIRMAN KOTELCHUCK: Very good.
9	Okay. So the 48.79 was the resolution?
10	MR. SIEBERT: Right. Forty-eight
11	
12	CHAIRMAN KOTELCHUCK: The
12	CHAIRMAN KOTELCHUCK: The correction on the errors.
13	correction on the errors.
13 14	correction on the errors.  MR. SIEBERT: Correct.
13 14 15	correction on the errors.  MR. SIEBERT: Correct.  CHAIRMAN KOTELCHUCK: Okay. Then
13 14 15 16	correction on the errors.  MR. SIEBERT: Correct.  CHAIRMAN KOTELCHUCK: Okay. Then that should close it.
13 14 15 16 17	correction on the errors.  MR. SIEBERT: Correct.  CHAIRMAN KOTELCHUCK: Okay. Then that should close it.  MEMBER MUNN: Well, we need to
13 14 15 16 17 18	correction on the errors.  MR. SIEBERT: Correct.  CHAIRMAN KOTELCHUCK: Okay. Then that should close it.  MEMBER MUNN: Well, we need to remind ourselves from time to time that
13 14 15 16 17 18 19	correction on the errors.  MR. SIEBERT: Correct.  CHAIRMAN KOTELCHUCK: Okay. Then that should close it.  MEMBER MUNN: Well, we need to remind ourselves from time to time that observations do not rise to the level of

observations and not specific findings. 1 I'm certainly content with the response that's 2 3 been given. KOTELCHUCK: Okay. 4 CHAIRMAN Brad, David? 5 MEMBER CLAWSON: This is Brad. 6 Ι 7 don't really have anything at this time. CHAIRMAN KOTELCHUCK: Okay. 8 David? 9 10 MEMBER RICHARDSON: I was looking 11 for whether there were lessons learned from 12 this, and it seems to be very much like other 13 things in the past. Some of them are 14 inexplicable. It's not clear why things weren't caught in review, and it's not clear if 15 anything has been learned from it. And then 16 17 revisions that have been made, it seems like we're chasing after a tail of a moving truck so 18 19 that there are we're laboring over 20 observations and responses. Well, that was a

long time ago and things have changed, and what

you're studying is not relevant to the practice

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today. So those are my observations.

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CHAIRMAN KOTELCHUCK: Yes. Well, your latter observation says that we are going to put in -- those may be old errors, but they are errors. And when we write our report to the Secretary, those will have to be included and, if you will, count against us. But I hope that what will be reflected in that report is that those were done a while ago, they were corrected, they were updated and corrected afterward, but that the errors were made at that time and I hope that the errors that are made on the more recent sets will be fewer. We believe so. I hope so.

Okay. Well, then let's move on, folks.

MR. FARVER: Okay. Tab 235, there were no findings. There were two observations. The first observation has to do with a reference that was used in the DR report, an incorrect reference, and it should have pointed to the Y-12 K-25 plant, rather than

PROC-61. So we've seen those type things before.

The second observation is similar to the previous one where the CAD tool now takes in the specific intake rates and time periods with recycled uranium. It's very similar to the previous observation.

CHAIRMAN KOTELCHUCK: Okay.

MR. FARVER: And that will take care of 235. 236, finding one, NIOSH did not account for all the reported dose. When we went through the record, we found 20 millirem that was in the dosimetry records but was not accounted for in the dose calculation.

When I went and started doing some digging, I found out that the lens of the eye doses were entered as deep doses and the deep dose was entered as lens of the eye dose. So it was entered incorrectly by the keypunch operator. Now, I don't know what kind of controls they have to prevent this, but it happened.

1	CHAIRMAN KOTELCHUCK: Yes. NIOSH
2	folks, I see your comment. Do you want to say
4	Torks, I see your comment. Do you want to say
3	anything?
4	MR. SIEBERT: Yes, the only
5	additional thing that we did is we have gone
б	back and we've updated the data entry files.
7	We've switched those numbers so that they are
8	correct.
9	CHAIRMAN KOTELCHUCK: Yes. Well,
10	that's all you can do when you have an error is
11	to correct it and then look at its impact.
12	Okay. You had observations on 236?
13	MEMBER RICHARDSON: Can I follow
14	up?
15	CHAIRMAN KOTELCHUCK: Yes, please.
16	MEMBER RICHARDSON: Because, I
17	mean, we've had this discussion before. The
18	other thing you can do with data entry errors
19	is do some sort of double entry, at least on a
20	fraction of them. I mean, the response has
21	been that it takes time and it costs money. But
22	to me, you know, starting off with data entry

errors is, you know, it is a problem that can be handled through -- the way that research has handled data entry problems.

So I believe nothing has changed. It's always been a hard thing to get a handle on. We review such a small fraction of cases. We're talking, you know, in proportional terms that when we're doing a survey sample, if we extrapolate up and we find keypunch errors, we have to say, well, in a one-percent sample, we find, you know, one, two, three data entry errors, that suggests that the data entry keypunch problem is on the order of magnitude of whatever it's going to be, one, two, three percent of cases have problems of mis-entry of basic information going into a claim.

CHAIRMAN KOTELCHUCK: That's a fair appraisal. At least to me, it seems worthy of further consideration. That is to say further consideration as to whether there should be some double-checking, double data checking of the second checking of the data for

some fraction of the samples, even of those that we=re doing, that NIOSH and SC&A are doing.

I hadn't been party to that earlier discussion about, you know, we can't double enter the data, we can't do double entry.

Does someone want to fill me in on

Does someone want to fill me in on what has gone on before or what --

MR. CALHOUN: Yes, this is Grady. And I think that David is right. In the past, we've said that it's just too costly for us to do that. I mean, we can always take a look back and see if there's something additional that we can do, but I'm not prepared to say that we're going to go forth and start doing that at this point.

CHAIRMAN KOTELCHUCK: Well, maybe this is something that we can hone in on in our report. I mean, the report is certainly on my mind as we're coming toward it. And if we're able to use the data that we've been reviewing, it may make sense to reopen that issue, and we'll do it in a way that will go to the

Secretary and move up the ladder a little bit because, obviously, it does cost money. But it may be, I'm not going to say worth it. It may be proper. So --

MEMBER RICHARDSON: Yes. I don't it's worth it know if in the sense financially or anything else. I mean, it's an issue which I still don't feel like I have a handle on. And there's a non-random selection of cases, so it's hard to understand where --I don=t understand the etiology of these data keypunch errors. If it was a research project, I would want to understand the etiology of it. I would want to know are they recurring from the same keypunch operator, or are there multiple keypunch operators who have done it?

There's been no description of, you know, who entered the data and why, who oversaw it, who was responsible for signing off and saying it was correct, and what's the magnitude of this? If we had to put confidence intervals around the uncertainty on keypunch errors at

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this point, it would be very, very wide because we have the time and resources to only review a very, very small fraction of the number of claims that have gone in and the number of records that have been keypunched. And so I have a huge amount of uncertainty about the fundamental issue of errors in the basic information that go into the calculations.

CHAIRMAN KOTELCHUCK: Yes. In the earlier case, I certainly took note of the comment that the person who made the error is no longer with us, with the team. And I did not know whether that meant that the person left on their own, I should say left on their own accord or whether the person had been found to have too many errors and something was done about it in employment terms.

And I don't want to know specifically about that case, but maybe, as we write our report, we can definitely go back and find, we'll have a record of the cases where there were keypunch errors and we can ask for

those cases. We can ask the NIOSH folks if they would give us data on how the errors were spread out.

MEMBER RICHARDSON: Right. Now, we asked to look at that, and that was not information, I believe, that was tracked at the time that we asked about it. And it's, you know, again, my comment then was I don't want to track it to be punitive in anyway, but, to me, it would be part of understanding how those problems arise and kind of where the target efforts to make sure that we get the highest quality data as input.

CHAIRMAN KOTELCHUCK: Right. No, I also agree. The issue is not punitive. I think the issue is finding out where the problem is, and we do have a way of dealing with that problem called double-entry data input. But that costs money, and there has to be a very good justification for it, mainly that there are too many, there are a moderate number of claims where this is a problem. It's hard to say too

1	many claims because one, you know, one claim
2	wrong is one too many. We're not perfect, and
3	we can never be perfect in any effort of this
4	sort. We can try as hard as we can, though, to
5	be.
6	Anyway, I think we can do this in the
7	future as we write up our report. So I don't
8	know that we need to talk further on this, or
9	I should say I don't know if we should belabor
10	this more, though I'm one of the belaborers.
11	Should we move on? It is 12:25, folks. So
12	maybe
13	MEMBER RICHARDSON: Sure. I just
14	didn't want to drop it because it's been a pet
15	project of mine.
16	CHAIRMAN KOTELCHUCK: Okay.
17	Well, it's appropriate that it should be a
18	project of the Subcommittee, as well, I
19	believe. Do we have anything more on 236? We
20	have an observation. Can we just finish up
21	236?

Sure.

MR. FARVER:

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Observation.

When we were reviewing the doses, what we tried to do was match the doses. And in this case, we could not match the doses. This would be the photon doses. And we were off by approximately 30 percent. That's where the 1.3 comes in. NIOSH's doses were 1.3 higher or 30 percent higher, I believe. Yes.

What it comes down to in this case, based on their explanation, is it was a Monte Carlo calculation. You know, I've seen them where they can be 10 percent, they can be 30 They could even be higher than that, percent. deviate from where you're just going with a straight, using a DCF instead of using Monte Carlo version for the DCF. It was an artifact of the Monte Carlo calculation. But, usually, if it's 30 percent more, we will probably write that up as a finding if we can't determine what really caused it. But we just wrote it up as observation because it an was claimant-favorable, in our opinion.

CHAIRMAN KOTELCHUCK: Okay. Does

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what. 1 anybody have а comment about confidence interval is or what the potential 2 3 deviation between Monte Carlo calculations Is 30 percent considered too high? 4 would be? We've talked about the fact that Monte Carlo 5 6 calculations vary, the results often vary. 30 percent variation, is somebody willing to 7 say that's really high? 8 MR. SMITH: This is Matt Smith with 9 10 ORAU team. It's really going to depend on 11 things like geometric standard deviation if 12 it's a log-normal distribution. If it's a 13 log-normal with a rather large GSD, then you can get a pretty wide spread in the data. 14 Conversely, if 15 it's а normal distribution and, you know, just a 10 percent 16 plus or minus spread, well, then things usually 17 stay pretty tight to that. We really see it in 18 the log-normal. And years from now when folks 19 are looking at CLL claims, they'll see things 20

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KOTELCHUCK:

with Weibull distribution embedded.

CHAIRMAN

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Yes.

1	That's helpful. Thank you. That's helpful,
2	to me at least. Okay. Then I think we're
3	finished with 236. 236 is closed then. And it
4	is just about 12:30, so can we come back at 1:30
5	and we'll start with 246.1?
6	(Whereupon, the foregoing matter
7	went off the record at 12:29 p.m.
8	and went back on the record at
9	1:33 p.m.)
10	CHAIRMAN KOTELCHUCK: Let's go.
11	Alright. Then, Doug, would you start? And I
12	believe it was on 246.
13	MR. FARVER: Yes, 246, Finding One.
14	These have to do with the x-ray doses, medical
15	doses. A little background: This employee
16	worked at K-25 for about 35 years, so there were
17	a lot of x-ray exams, not just chest x-rays but
18	there's injuries. When I went back to review
19	the exams, there were over 60, so it's just
20	numerous x-ray exams.
21	So when we're reviewing the
22	original dose reconstruction, we noticed that,

according to Procedure 61, there's certain exams that you do not include as part of the dose reconstruction. And when we reviewed it, at least some of those exams that were included fell into that category where they strictly should not have been included, and that was the basis for our finding.

Basically, what it comes down to is NIOSH was claimant-favorable. And it was very confusing. When you look at all these exams, it's just very difficult to tell what is illness related and what is just a regular chest x-ray. So it was very difficult to read some of those, especially the earlier ones.

So, in essence, they were claimant-favorable, adding some x-rays that strictly probably should not have been included. This was a difficult case for them. Let's cut them some slack. They did good, given the number of exams. But our finding was just based on strictly by the procedure, and there's some that should not have been

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1	included. However, it was
2	claimant-favorable, so we figured we might as
3	well just close this. It's okay.
4	CHAIRMAN KOTELCHUCK: Okay. Was
5	that because I don't have my screen up yet.
6	Was that an observation?
7	MR. FARVER: It was a finding.
8	CHAIRMAN KOTELCHUCK: Okay. Any
9	other comments?
10	MEMBER MUNN: No. I think we ought
11	to accept SC&A's recommendation to close.
12	CHAIRMAN KOTELCHUCK: Okay.
13	MR. SIEBERT: And this is Scott. I
14	do just want to point out that I wouldn't
15	necessarily say we didn't follow the procedure,
16	as we were probably more claimant-favorable
17	when the dose reconstructor wasn't sure if it
18	was an x-ray exam or for employment. So they
19	were claimant-favorable and let some in.
20	CHAIRMAN KOTELCHUCK: Good.
21	Okay. Well, I think we can close.
22	MR. FARVER: Alright. Oh, I did

1	ask Steve Marschke to phone in.
2	CHAIRMAN KOTELCHUCK: Oh, yes.
3	MR. MARSCHKE: I'm on the phone.
4	MR. FARVER: Okay. Do you want to
5	go back and let him explain about that finding?
6	CHAIRMAN KOTELCHUCK: Yes, please.
7	MR. FARVER: Steve, we were talking
8	about the Huntington Pilot Plant, that matrix
9	from the findings from your report.
10	MR. MARSCHKE: Okay.
11	MR. FARVER: And you sent me your
12	recommendations about modifying the TBD. And
13	during the meeting today, we found out that they
14	already revised the TBD, so we'll have to make
15	sure these changes were made.
16	There were some questions about
17	finding four. I believe it comes down to a
18	difference between MicroShield and what NIOSH
19	used to do their calculations.
20	MR. MARSCHKE: Exactly. I think
21	that's exactly right. I don't have, I didn't
22	have access to MCNP, so I did the calculation,

both calculations using MicroShield. But in the TBD, in Appendix A, Table A-2, NIOSH gives the photon profile, if you will, or photon spectrum that they used that were generated by, I think it was probably by MCNP.

and put it into MicroShield, and MicroShield calculated a dose and it got about the same results that NIOSH reported in the TBD. About 50 percent of the dose was due to photons with a 0.25 or less, MeV or less, and the other 50 percent was due to photons with 0.25 or more. And so that, you know, confirmed what they said.

But then when I ran MicroShield and put in and let MicroShield calculate the photon spectrum from U-238, by decaying U-238 and U-235, I got a different spectrum. I got the same, about the same dose when you look at the total integrated dose over all the energy spectrum. I got good agreement, but a little bit more of the dose came from the less than 0.25 MeV photons and a little bit -- oh, no, it's the

other way around. A little bit more of the dose 1 2 came from the less than 0.25 photons, and a 3 little bit less of the dose came from the greater than 0.25 photons. 4 And so, you know, NIOSH came back 5 6 and said, well, we used the latest and greatest 7 energy data, photon data, and, you know, it's hard to argue with that. So that was, you 8 know -- I still don't know exactly why the doses 9 or the breakdown was different, but it's, you 10 11 know, if they're using the latest and greatest 12 data, you know, again, like I say, that's the way I think you should go because I'm not 13 completely enamored with 14 the MicroShield energy nuclide library. 15 16 CHAIRMAN KOTELCHUCK: Okav. This 17 MEMBER MUNN: sounds like another one of those cases where it's an issue 18 of differing tools. 19 CHAIRMAN KOTELCHUCK: 20 Yes. Alright. I quess 21 MEMBER MUNN: 22 you can't use apples to check oranges.

1	MR. MARSCHKE: That's correct.
2	CHAIRMAN KOTELCHUCK: My basic
3	concern, which we discussed earlier, was that
4	20 years from now somebody could come back and
5	run it as NIOSH had run it and it would be
6	reproducible. And this is a matter of
7	different libraries, and that seems I'm not
8	too concerned about that, other than I don't
9	understand why.
10	MEMBER MUNN: I didn't hear you
11	very well. That broke up pretty badly.
12	CHAIRMAN KOTELCHUCK: Oh, I say I
13	don't understand why the two libraries differ,
14	but, as long as it's reproducible, to me that's
15	the critical issue.
16	MEMBER MUNN: I suspect we'd have
17	to get pretty deep in the weeds of the software
18	to understand why they differ.
19	CHAIRMAN KOTELCHUCK: Yes. And
20	that's not our responsibility.
21	MEMBER MUNN: Yes, I think that's
22	beyond our scope. The fact that two different

1	tools are being used to approach the same
2	problem could be applauded from one direction,
3	and from another we could say, well, that
4	obviously can't be used as a total baseline.
5	MEMBER CLAWSON: This is Brad.
6	We're all supposed to be using the same tool,
7	so tools are supposed to be set up so this is
8	going to be able to be represented and redone
9	and there won't be
10	CHAIRMAN KOTELCHUCK: But I've
11	just been reading email in the last week that
12	there's a real problem for SC&A being able to
13	get access to the tools that NIOSH uses for,
14	let's say
15	MEMBER MUNN: Yes, I
16	CHAIRMAN
17	KOTELCHUCK: bureaucratic reasons.
18	MEMBER MUNN: Yes, that seems to be
19	true in several occasions.
20	CHAIRMAN KOTELCHUCK: Yes. I
21	think that's a longstanding problem. But I'd
22	move to just close this now.

1	MR. FARVER: Okay. Now, if we
2	accept the HPT matrix and all that, that will
3	close our final finding, which was 185.7, from
4	the 9th set.
5	CHAIRMAN KOTELCHUCK: That's
6	correct.
7	MR. FARVER: Okay.
8	CHAIRMAN KOTELCHUCK: And that's
9	lovely. Okay. [Set] nine is closed. And
10	thank you, Steve.
11	MR. MARSCHKE: You're welcome.
12	CHAIRMAN KOTELCHUCK: And we
13	should now get back to Oak Ridge and 246.
14	MEMBER MUNN: Did I lose you or did
15	we lose somebody?
16	MR. FARVER: No, I'm making
17	modifications to the 9th set, and then I'll go
18	back to my
19	CHAIRMAN KOTELCHUCK: Oh, sure,
20	sure. We're fine.
21	MR. FARVER: Then I'll go back to
22	the Oak Ridge set. Okay. 246.2. The finding

1 reads that NIOSH did not appropriately assign external ambient dose. This finding came out 2 3 of a --CHAIRMAN KOTELCHUCK: 4 By the way, 5 we don't have anything on our screen as you're 6 reading, Doug. 7 MR. FARVER: Okay. CHAIRMAN KOTELCHUCK: But that's 8 9 okay, although if we're just moments away let's 10 get it on the screen. There we go. Nope. 11 There we go. 12 Great. Thanks, MEMBER MUNN: John. 13 The basis for this 14 MR. FARVER: finding comes from our one-on-one conference 15 calls that we had with the Board Members when 16 17 we were reviewing the draft. And one of the Board Members had a concern regarding the 18 assignment of external on-site ambient dose for 19 20 46 and 47 when the employee was a truck driver because the truck driver's duties involved many 21

different things, such as loading and unloading

radioactive cargo. And in the Board Member's opinion, the assignment of on-site external ambient dose does not account for the worker's external exposure during these years, and it would have been more claimant-favorable to assign unmonitored dose based on co-worker data. So that's the gist of the finding.

CHAIRMAN KOTELCHUCK: Okay.

MR. FARVER: We can see NIOSH's response then. I agree there is no information provided in the files that shows that the employee handled radioactive materials. Agree.

CHAIRMAN KOTELCHUCK: And security clearance in 1948, which is to say later, so that if the person were handling nuclear materials, they would have had to get security clearance earlier. I don't know how long security clearance takes, but it would quite reasonably be, in my mind, the case that the person was working in '47, already starting to load nuclear materials, in which case his

supervisor would say, wait a minute, you've got 1 to get security clearance, and that would take 2 3 some period of time. Ι assume months, And, therefore, possibly the person 4 whatever. 5 was handling in '47. 6 MR. CALHOUN: This is Grady here. 7 One thing we've got to make sure of is the idea of him loading nuclear materials is completely 8 hypothetical. We have no idea that's what he 9 10 was doing. 11 MEMBER MUNN: Would it -- how can we 12 make the assumption that that happened and, at the same time, fail to note that he was being 13 badged for such things? 14 Even in the early years, it was recognized that folks 15 actually handled radioactive material needed 16 badged, regardless 17 be of their description. 18 Right. 19 MR. CALHOUN: And it's my understanding, too, that in the earlier years, 20 you had to have a security clearance before you 21

were allowed to work in the areas where

1	radioactive material was.
2	MEMBER MUNN: Pretty much. The
3	whole thing was top secret for years after the
4	40s.
5	CHAIRMAN KOTELCHUCK: Yes.
6	MR. CALHOUN: So, basically, I
7	think what we're saying here, and Scott or Mutty
8	can jump in, is that kind of the weight of the
9	evidence here supports the use of ambient dose
10	here. And we do get dosimetry from this site
11	routinely, and this guy didn't have it, you
12	know, in that period. So I think that that and
13	some of the other information that we have, it
14	seems to support that decision.
15	MR. SIEBERT: Yes, this is Scott.
16	I would agree with Grady that that's what we
17	looked at. The preponderance of information
18	did not support him doing any radiological work
19	during that time.
20	CHAIRMAN KOTELCHUCK: I'm just
21	worried about sort of boundary value problems.
22	I would agree for '46, but, at some point, well,

at some point I would have just a concern, although there's not clear evidence. But I think it's, you know -- well, could I ask someone how long would it take to get security clearance back in those days if somebody decided this person needed security clearance? Weeks? Months?

MR. CALHOUN: I don't know that, but I think, David, one thing you're thinking is that he'd be allowed in that area without a clearance, and I'm not so sure that's true. And there's another thing in there, too, if you look. There's a sign-off on a supervisory industrial health exam, and that also coincides with right before we got our bioassay. So we've got a minimum of two things that kind of indicate that he wasn't working with rad material, along with the big thing that there's no indication that he was.

CHAIRMAN KOTELCHUCK: Well, that's evidence. That is evidence. Could somebody move the screen up a little bit so I can read

a little bit more of the NIOSH answer, the greens? Yes. Yes, okay. You present evidence here and admit that it's supposition, certainly on my part and maybe on the other reviewer's part. So I'm willing to accept that the person did not begin to get exposure until '48. What do others think, particularly Subcommittee Members?

MEMBER CLAWSON: This is Brad. You know, we're making a lot of assumptions here, and that's what we have to come down to. And we can all think that in the ideal situation that he didn't get there. You know, he may not have been able to work with S&M, but he also might have been able to work with waste because that didn't need a clearance. There's a lot of things that play into this, and, you know, we're trying to make a decision here of using the ambient dose or the missed dose. And myself, I personally think for that time, you ought to give him -- if he's in that area, if he's driving around in those areas and stuff like that, give

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him the missed dose or whatever. That's just feelings on it because, you know, you're painting one side of the picture, but we could actually paint the other side of the picture and say, yes, but he could have been in there. It's the whole thing, neither side has --

CHAIRMAN KOTELCHUCK: Well, yes, There's reality and there's that's right. evidence, and I think there may be lots of perceptions about reality, but if we don't have evidence I find it very hard to act on that. When we have evidence, I think we have to follow it, even though I'd agree with what you said. And I would also, I can't help but believe, that fellow was exposed in '47, I mean even though it shouldn't have been. But there's so many things that shouldn't have been that people do because you've got to get production done, you've got to get things delivered, you've got to -- this was, you know, this was, I wouldn't say a war-time setting but it was a setting of great tension and, you know, defense

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However, that's speculation. That is just speculation on my part, and there is no evidence to support it.

Well, actually, some MEMBER MUNN: of us actually remember the late 40s, and it was not a time of buildup. Au contraire. glorying at the end of a terrible war that had taken literally millions of our people by surprise, and we were able to prevail in that. And we were in the process of falling back from that as much as possible. The armed forces, obviously, were reducing their ranks by the We had guys out and everybody was thousands. breathing a great sigh of relief. No war on the horizon yet as far as the Korean Peninsula was concerned, and I wouldn't think, of any time during the entire history of the weapons complex, I would think this would be one of the times when there was least production pressure.

So I guess we're talking assumptions here, but we really have to, I

think, go on the bulk of obvious evidence. And the obvious evidence says to me that it's unlikely that this individual was handling radioactive materials at that time because we were very sensitive about that. I would imagine he would be badged as soon as he began to do that. I would consider it to be a requirement, as a matter of fact.

CHAIRMAN KOTELCHUCK: We'll talk about that sometime. You lived it on site there, so I will not.....it will be an interesting conversation. The Cold War had already begun, and these were important things.

However, I agree with you that you have to go by the evidence, and the evidence, to the extent that it's there, does not speak to exposure before '48. So as far as I'm concerned, my vote is that we have to just simply accept that it started in '48 and that the concern of the person who reviewed this was a good concern, but I don't think we can act on it.

MEMBER CLAWSON: But, Dave, I want to throw out one thing because Wanda threw out something really wonderful and I thought it was great. Let's look at all the DOE sites and all the facilities and let's figure out by weight of the evidence how many times there was a lot of problems there and what they said they didn't do, and it was a whole other issue. We keep forgetting that. That's part of the reason why we're even in this mess right here.

CHAIRMAN KOTELCHUCK: Yes.

MEMBER CLAWSON: So you talk about weight of the evidence, there's a lot of evidence there, too. But you know what? I agree. I don't have a problem with this one here. It's a little bit bigger picture. I think we're still making assumptions on this, and when we don't have the weight of the evidence I think we ought to go with the side of the claimant. That's my personal --

CHAIRMAN KOTELCHUCK: Right. And I agree with you that we go with the evidence,

1	despite whatever feelings we have about what
2	might have been. David Richardson, you're a
3	Subcommittee Member who hasn't said anything on
4	this. How do you feel?
5	MEMBER RICHARDSON: I'm fine with
6	kind of moving forward with this. I don't
7	think there's much more we can resolve.
8	CHAIRMAN KOTELCHUCK: I agree.
9	So, folks, I think we have pretty well come to
10	a conclusion to close this.
11	MR. FARVER: Okay. We'll close
12	that one. Now, that one is for the external.
13	The next finding, 246.3, deals with the
14	internal for '46 and '47. They assigned an
15	environmental dose, and the finding says that
16	they should have assigned a co-worker dose
17	based on OTIB-14, job category of a driver,
18	category two, which has jobs generally that
19	have some potential for workplace internal
20	exposures, depending on job specifics.
21	This kind of ties on to the first
22	one. But I would say that if you can't buy into

1	the external, then you can't buy into the
2	internal. So they assigned environmental,
3	which, if they're going to assign ambient for
4	external, it probably is appropriate to do
5	environmental for the internal for that same
6	time period. We're talking two years.
7	MEMBER MUNN: Can we move the
8	screen so that we can see the rest of the
9	response there? Even though it was just read,
10	it would be helpful. Thank you.
11	CHAIRMAN KOTELCHUCK: Yes.
12	MEMBER MUNN: So what are we
13	discussing?
14	CHAIRMAN KOTELCHUCK: 246.3. The
15	assignment of environmental dose.
16	MEMBER MUNN: And I'm still waiting
17	to see the bottom part of it. We've seen the
18	top of it here, right?
19	MR. STIVER: Wanda, basically,
20	it's the same responses for 246.2, so I just
21	went back up to that.

CHAIRMAN KOTELCHUCK:

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Right.

1	MEMBER MUNN: Right. That's what
2	I thought.
3	MR. FARVER: Yes, it's just the
4	internal component. I mean, it's pretty much
5	the same argument.
6	MEMBER MUNN: Yes.
7	CHAIRMAN KOTELCHUCK: Yes, I think
8	so.
9	MR. FARVER: So I would just use the
10	same exposure
11	MEMBER MUNN: Yes, yes.
12	CHAIRMAN KOTELCHUCK: I guess so.
13	MR. FARVER: Insufficient evidence
14	to make a change.
15	MEMBER MUNN: And it's unlikely.
16	I can't see that there's an issue that we
17	haven't already discussed.
18	MR. FARVER: The only thing that
19	makes this a little different is there is a
20	little basis in Attachment A of OTIB-14. But
21	like I said, if you don't buy into the external
22	side, then I'd have a hard time justifying the

1	internal side.
2	MEMBER MUNN: Yes, that doesn't
3	seem reasonable unless there is evidence of
4	internal injury that carried a deposition with
5	that.
6	CHAIRMAN KOTELCHUCK: Okay. Yes,
7	I think so. Should we close it, folks?
8	MEMBER MUNN: Yes, please.
9	CHAIRMAN KOTELCHUCK: Hearing no
10	objection and some support, let's go on. Let's
11	close it.
12	MR. FARVER: Okay. Now, 247.1.
13	Inappropriate exposure period. They prorated
14	the dose, but it was prorated incorrectly.
15	MR. KATZ: While you're doing that,
16	Doug, I have a note from someone asking if I
17	would just remind folks who are not speaking to
18	mute their phones. So anyone who's just
19	listening, please mute your phone. Press *6 to
20	mute your phone, and press *6 again to come off
21	of mute, or press the mute button if you have

Thanks.

a mute button.

1	MR. FARVER: Okay. The employee
2	worked for six months in 1956, and the dose
3	reconstructor used a value of 0.49 months
4	instead of 0.49 years, [that is] 0.04 years.
5	CHAIRMAN KOTELCHUCK: That's it.
6	MR. FARVER: Oh, okay. They used
7	0.04 years instead of 0.5 years. They prorated
8	it incorrectly is what it came down to.
9	MEMBER MUNN: Do I understand
10	correctly that it appears that review will more
11	than likely change the PoC?
12	MR. SIEBERT: Yes. Wanda, this is
13	Scott. We worked this one because, looking
14	forward in time, the X-10 co-worker values
15	changed dramatically and there's already the
16	assumption we will have a PER for that coming
17	along down the road. So we ran with those
18	numbers, and it will likely be compensable
19	based only on the plutonium itself, without
20	even dealing with everything else. So we
21	didn't go back and look at it.

MEMBER MUNN: Yes, yes, I imagine

1	that the PER will take care of the whole thing.
2	CHAIRMAN KOTELCHUCK: But it
3	hasn't yet? Question.
4	MEMBER MUNN: I don't think the PER
5	has been issued yet, has it?
6	MR. SIEBERT: That's correct.
7	There's still some other updates to the X-10 TBD
8	that we're working on. So once those all get
9	rolled out, then we'll do a PER all rolled
10	together. Grady can probably correct me if I'm
11	wrong, but that's my understanding of what
12	we're planning for X-10.
13	CHAIRMAN KOTELCHUCK: Okay.
14	MR. CALHOUN: That's correct.
15	Yes, that will flip because of a revision to the
16	co-worker data, or may flip, not because of the
17	mistake in prorating.
18	MEMBER MUNN: There's a lot going
19	on still.
20	CHAIRMAN KOTELCHUCK: Yes,
21	apparently. It seems like 247.1 is still going
22	to be open, right? It's open.

1	MR. CALHOUN: We may not get to that
2	PER for a year. So if we want to keep it open
3	for a year, that's okay, just so you guys know.
4	CHAIRMAN KOTELCHUCK: Well, I most
5	certainly do not want to keep it open for a year
6	but
7	MEMBER MUNN: Well, it's a process
8	issue we probably need to resolve in our own
9	minds because this is not going to be the only
10	time that will occur.
11	MR. CALHOUN: We have reviewed
12	hundreds and hundreds and hundreds of cases via
13	our PER system and the system we call PADs. And
14	whenever we come up with a case that flips or
15	has the potential of flipping, we make that
16	request to DOL to reinstate the case, and they
17	do and we rework it.
18	MR. FARVER: Now, keep in mind that
19	the finding had to do with a calculation error
20	in the prorating. This information about the
21	co-worker and the PER that's also going on, but

that did not really have to do with the finding.

MEMBER MUNN: That's a good point because what this boils down to is calculator error. This is a human error issue. They chose the wrong input, and the PER is not going to change that. It will change the end result, but it's not going to change what happens in this case.

CHAIRMAN KOTELCHUCK: I'm a little concerned now because we can correct the, we can take care of the calculational error now. That will not change the PoC, and I believe the person will get a rejection letter. We can then open it up in a year when the PER comes out, as will open up all the cases that were affected by that PER.

At a human level, I feel like to deny the person in anticipation that we probably will change it or maybe there could be something in whatever statement that's given to the person who filed that, there are other changes that may occur in the future and that we're anticipating some changes and that may have

some effect. I'm just worried about telling a 1 person, no, you're claim is denied when we know 2 3 at this point that probably it will not be. can't say it will not be or it will be accepted. 4 But it's clear in our own minds that there's a 5 6 pretty good chance it will be accepted. Well, this raises a 7 MEMBER MUNN: process issue as far as the Agency is concerned, 8 I was of the impression that once 9 in my mind. 10 an error had been identified that the Agency would, by their own process, recalculate that 11 12 PoC and proceed accordingly. Am I incorrect in 13 that? 14 MR. CALHOUN: You're not incorrect 15 in that. Typically, we come across those 16 issues through the PER or PAD process. in 17 MEMBER MUNN: Yes. But situations like this, and this is not the only 18 19 one, you've looked at this case now and we've found that there's an error and we think it's 20 21 likely compensable, will the Agency not, as a

matter of process in their own procedures,

1	undertake the reassessment of the case for the
2	client?
3	MR. CALHOUN: I'd have to look at
4	that case. It seems like we could.
5	MEMBER MUNN: It seems likely to
6	me. I thought that's what occurred. I didn't
7	think we had to wait for a PER in a case like
8	this. I guess I'm trying to identify whether
9	Dave's concern is justified. In my mind,
10	you've found an error and I thought the Agency
11	proceeded accordingly.
12	CHAIRMAN KOTELCHUCK: Which is to
13	say corrected it.
14	MEMBER MUNN: Yes.
15	CHAIRMAN KOTELCHUCK: That error,
16	yes.
17	MEMBER MUNN: Yes, it's my
18	understanding that this would automatically
19	trigger a recalculation of the case for
20	compensability.
21	MR. CALHOUN: Just leave that one
22	open then and let me look at it and make sure

that the TBD is, in fact, the co-worker study is, in fact, revised officially and we can get back to that one next week. And if it is and it will flip, and I don't think we've done the calculations officially through the PER, then we'll make that notification.

MEMBER MUNN: Yes, I think there was a question in Dave's mind as to whether or not we had to wait for a PER for this claim to be addressed and the error corrected. I thought you corrected errors when we found them.

CHAIRMAN KOTELCHUCK: Right. I'm thinking of farther along. But you know what? We've got to leave this open a little bit, and I'm open to leaving it open just for a bit.

MEMBER MUNN: Yes, I think that's fine because we can get a report back as to whether or not NIOSH is undertaking a redo of this claim, which is a logical thing. We've found an error, a human error but error nevertheless. It will be addressed.

1	CHAIRMAN KOTELCHUCK: Agree. And
2	resolved.
3	MEMBER MUNN: Yes.
4	CHAIRMAN KOTELCHUCK: Okay, folks.
5	This will remain open.
6	MEMBER CLAWSON: This is Brad.
7	That's fine.
8	CHAIRMAN KOTELCHUCK: Yes. Okay.
9	MR. FARVER: Okay. 247.2,
10	inappropriate method used to determine modeled
11	photon dose at Y-12. The basis for this
12	finding is in the file we came across this
13	co-worker data photons tool, worksheet,
14	whatever you want to call it, that we have not
15	seen before and not familiar with how it was
16	used. And this is their person who normally
17	does all the Oak Ridge cases.
18	So I had a concern about what the
19	basis is and is it working correctly and so
20	forth. And when he went back to the ORAUT
21	report 32 and looked at some numbers and tried
22	to do some calculations, they did not match up

1 to what this worksheet was saying. That's what 2 prompted this finding. 3 MR. KATZ: Doug, are you still there? 4 Yes, I'm still here. 5 MR. FARVER: 6 Really, what it comes down to is the new 7 worksheet that we had not seen, and we did not understand the basis for it. And the NIOSH 8 9 response is, basically, that they're putting 10 together files to help explain it to us. Well, 11 this is pending. 12 CHAIRMAN KOTELCHUCK: And, NIOSH folks, where is that? What stage is that? 13 This is Matt Smith with 14 MR. SMITH: I found this deep in my archives. 15 ORAU team. A brief explanation I can give right now is 16 17 there is an OTIB called OTIB-12, and that kind of gives a general outline of how Monte Carlo 18 tools were being used at the time this claim 19 went through the production process. 20 basically, what it was doing is to find the DCF 21 22 values of the distribution, the triangular

1	distribution, against the co-worker values
2	that we have for Y-12. At that time, a Y-12
3	co-worker model for external was, as everybody
4	is probably recalling, quite a complex set of
5	data that was based on some real statistical
6	analysis that had some variability associated
7	with it, depending on how much dose a worker got
8	after the, roughly after 1960.
9	So I'm at the point right now where
10	I'm going to put together a description of how
11	this whole thing was working back then. And
12	based on the data of the file I found, we're
13	talking almost ten years ago, so I want to put
14	together a good description that gets everybody
15	up to speed.
16	MEMBER MUNN: It sounds like this
17	remains open until Matt finishes his report to
18	us.
19	CHAIRMAN KOTELCHUCK: That seems
20	correct. Okay. That's open. So
21	MR. FARVER: Okay. 247.3,
22	inappropriate photon energy range used for the

NIOSH applied the 30 to 250 keV Y-12 site. photon energy range to all three sites. person worked at all three of the Oak Ridge So they're appropriate for the K-25 sites, but for the Y-12 site the OTIB lists 50 percent 30 to 250 keV and 50 percent greater than 250 keV. It may not have been appropriate for the Y-12 site, but it was applied across the board to all sites. That's what prompted the findings. You get a little higher REFs with the greater than 250 keV photons. So the dose is a little high, so that was kind of our concern.

And NIOSH gives a good explanation.

And we were concerned about it. A secondary concern was when I looked at the workbook. The maximizing assumption was to use the 30 to 250 keV, and we felt that that may not be maximizing and suggested that they look at it.

NIOSH provided a response and, basically, even though the DCF is a little higher, when it cuts into the IREP part, the

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1	REF, Radiation Effectiveness Factors, of the 30
2	to 250 photons are higher than for the greater
3	than 250 photons. So it seems they did do the
4	claimant-favorable thing. So I suggest
5	closing this because there's really no further
6	action.
7	MEMBER MUNN: Well, if that was the
8	claimant-favorable decision, then it appears
9	that we can close this item.
10	MR. FARVER: Yes.
11	CHAIRMAN KOTELCHUCK: Agreed.
12	Okay. Let's close, unless I hear objection.
13	Let's move on.
13 14	Let's move on.  MR. FARVER: Okay.
14	MR. FARVER: Okay.
14 15	MR. FARVER: Okay.  CHAIRMAN KOTELCHUCK: Which then
14 15 16	MR. FARVER: Okay.  CHAIRMAN KOTELCHUCK: Which then takes us to 247.6.
14 15 16 17	MR. FARVER: Okay.  CHAIRMAN KOTELCHUCK: Which then takes us to 247.6.  MR. FARVER: Yes, we'll get there
14 15 16 17 18	MR. FARVER: Okay.  CHAIRMAN KOTELCHUCK: Which then takes us to 247.6.  MR. FARVER: Yes, we'll get there eventually. 247.4 is the same prorating
14 15 16 17 18	MR. FARVER: Okay.  CHAIRMAN KOTELCHUCK: Which then takes us to 247.6.  MR. FARVER: Yes, we'll get there eventually. 247.4 is the same prorating issue, and this is for internal dose, where he

1	believe.
2	CHAIRMAN KOTELCHUCK: If I may ask,
3	is this two errors or one? I mean, they
4	calculated something incorrectly. They put it
5	in years, rather than months. But once they
6	did it, did they have to enter it again? I
7	mean, at some level, I would think the person
8	just said, oh, yes, I just calculated that here,
9	I'll put it in.
10	MR. FARVER: It was probably one
11	calculation to determine the fraction.
12	CHAIRMAN KOTELCHUCK: Right.
13	MR. FARVER: And then the incorrect
14	fraction was applied
15	CHAIRMAN KOTELCHUCK: Twice.
16	MR. FARVER: twice.
17	MEMBER MUNN: Yes, that would be my
18	assumption. I would see it as two errors
19	because it was a calculation had to be done
20	twice but only a single error in terms of
21	identifying the correction that was going to be

used. So it depends on how you look at it,

1	whether it was two errors.
2	CHAIRMAN KOTELCHUCK: Okay. I'm
3	thinking about our report, but you've answered
4	my questions and we had a discussion. Let's
5	continue on. Thank you.
6	MR. FARVER: Are we going to close
7	that item?
8	CHAIRMAN KOTELCHUCK: Yes.
9	MEMBER MUNN: I think so.
10	CHAIRMAN KOTELCHUCK: It's the
11	same issue.
12	MR. FARVER: Very similar for the
13	next one. It's prorated incorrectly.
14	CHAIRMAN KOTELCHUCK: Okay.
15	MEMBER MUNN: Another closure.
16	CHAIRMAN KOTELCHUCK: Yes.
17	MR. FARVER: Okay. Now, we
18	finally get to 247.6. The 1969 alpha dose was
19	not entered into the 2007 red bone marrow IREP
20	table. The employee had several cancers. He
21	had a 2006 red bone marrow, 2007 red bone
22	

made it into two of the IREP tables but not the third one, which we thought was a little strange. The calculation was done. It just was not, it wasn't there.

And the NIOSH response was, basically, it was a cut and paste error.

MR. SIEBERT: This is Scott. Yes, I mean, that's the generic issue. But let me just explain a little bit more. Since there were 2006 and 2007 cancers, you need to prorate the internal differently for those last years because, obviously, exposure that happened after the date of diagnosis did not impact the probability of that cancer occurring.

So what the dose reconstructor had to do was prorate the 1996, I'm sorry, 2006 doses and put those into the two that were correct. And then you had to also prorate the 2007 separately because for a 2007 diagnosis you would actually have the full exposure during all of 2006, and then they prorate the 2007 value.

When they did that proration, that intake, which is a co-worker from 1969, were the last, I want to say, 12 or so rows. And that's where they had to cut and paste so they didn't, they just didn't cut the last couple rows and paste it into the matrix when they did the prorating. So that's why there would be a difference between the 2006 and 2007 cancers.

The other thing I want to point out is we've updated, as we say all the time, we've updated our external tools. They now have the capability of importing some of the internal tools, such as CADW, IMBA, all those internal outputs that come from other tools. Instead of having to cut and paste those into the IREP sheet anymore, now we can pull that into the external tool, which is really more of an overall assessment tool, as opposed to just external. And it will automatically put it into the IREP sheet, as well, so we won't have those type of intake errors.

CHAIRMAN KOTELCHUCK: Yes, that's

good. Can I ask, I mean, that 247 was done by a person, right? It was checked by the supervisors, but it was done by one person. So we're dealing with the fourth error on one case by one dose reconstructor. Is that correct?

MR. SIEBERT: It would be the same person who did this, yes.

CHAIRMAN KOTELCHUCK: Yes. I mean, we discussed this a little bit this morning. I mean, at some point, as we begin to write up reports, we will need some documentation about that sort of thing.

And, again, I don't care about the name of the person and, you know, by no means are we [trying] to micro manage the operations from afar. But on the other hand, I can't help but note that we're going to be dinged for several errors, all of which were made in one case by one person, as opposed to three or four people, you know, each making a little error. That's a comment. That's not a discussion of how we dispose of this case because what you

1	said makes sense, and it is complicated.
2	So I think that your explanation
3	satisfies me that that was properly done, that
4	the dose reconstruction with respect to this
5	was properly done. Other folks?
6	MR. FARVER: It wasn't properly
7	done, it was a cut and paste error.
8	CHAIRMAN KOTELCHUCK: Right. I
9	should say not properly done, it was corrected.
10	I correct myself. It was found, and it was
11	corrected.
12	MR. FARVER: Well, it hasn't been
13	corrected yet.
14	MEMBER MUNN: We are assuming that
15	it's going to be corrected because we're
16	assuming that the Agency is going to redo this
17	entire claim.
18	CHAIRMAN KOTELCHUCK: Right. And
19	we don't have to, we don't have to keep it open
20	because the problems have been found, they will
21	be dealt with. There's not any question about

1	technical matter, quotes just a technical
2	matter. Expunge the word "just." It is a
3	technical matter, and I don't see that we need
4	to see it again.
5	MR. FARVER: Unless you want to
6	keep it open like 247.1, which we
7	MEMBER MUNN: Well, 247.1 is going
8	to keep the issue open
9	CHAIRMAN KOTELCHUCK: That's
10	right.
11	MEMBER MUNN: to ensure us that
12	the case is, in fact, being reworked and will
13	be handled appropriately.
14	CHAIRMAN KOTELCHUCK: Agreed.
15	MR. FARVER: So we'll close this
16	one.
17	CHAIRMAN KOTELCHUCK: Yes.
18	MEMBER MUNN: And you might even
19	refer back to .1 that it will be reworked.
20	MR. FARVER: Yes. Okay. 247.7,
21	NIOSH did not
22	CHAIRMAN KOTELCHUCK: Other Board

1	Members, by the way, just as we start, as we're
2	moving along and trying to move along, if there
3	are objections, please stop us and raise them.
4	That said, go on. 247.7.
5	MR. FARVER: 247.7, NIOSH did not
6	discuss all the incidents described in the CATI
7	report. We've seen this before where the
8	employee mentioned an incident, and we feel it
9	should be mentioned somewhere in the dose
10	reconstruction. NIOSH agrees that a comment
11	could have been added to address the event, so
12	we've talked about this before. There is not
13	much action we can take.
14	I will say they're getting better at
15	that. I've seen improvement. So I suggest
16	closing this.
17	CHAIRMAN KOTELCHUCK: Fine. This
18	might have been dealing with yes, in fact,
19	I think this could have been viewed as an
20	observation.
21	MR. KATZ: Doug, this is Ted. Can
22	I just ask the question: Are you recording the

nuances in terms of solutions where, for
example, NIOSH has instituted a systematic
correction for a kind of problem, like we've
talked about several of them today, like the cut
and paste and the cut and paste won't have to
be done anymore? Are you recording that along
with the finding, so that in the report that the
Board does it can sort of address the different
solutions that have been implemented, some of
which I guess are implemented independent of
the Board's finding because NIOSH made the
change before the Board made its finding.
But in any event, are you capturing
But in any event, are you capturing that information in your sort of matrix of
that information in your sort of matrix of
that information in your sort of matrix of information that you will then summarize and
that information in your sort of matrix of information that you will then summarize and produce for the Subcommittee, Doug?
that information in your sort of matrix of information that you will then summarize and produce for the Subcommittee, Doug?  MR. FARVER: For example, the last
that information in your sort of matrix of information that you will then summarize and produce for the Subcommittee, Doug?  MR. FARVER: For example, the last one, 247.6, I didn't include a lot of that
that information in your sort of matrix of information that you will then summarize and produce for the Subcommittee, Doug?  MR. FARVER: For example, the last one, 247.6, I didn't include a lot of that because it's included in the response.

guess, but will get statistics that will help the Subcommittee write its report to the Secretary? And some of those statistics will talk about sorts of different kinds of problems and how they were remedied and so on. They'll sort of need those data, and so somehow, I guess, you have to key this stuff so that you can easily summarize it.

MEMBER MUNN: I don't think we have had any programmatic way to do that. I don't think we set up our documentation in such a way that we can, certainly by machine, we can't just call out that information and have it come up for us.

MR. KATZ: I guess what I'm just saying -- I'm sorry. I don't mean to derail our progress, but this does relate to what we'll be talking about later with the report. I mean, it seems like it would be very easy to key the matrix with another column that makes some distinctions like this and helps you with summarizing.

1	CHAIRMAN KOTELCHUCK: You know, I
2	don't think I'd add another column to matrices
3	that we know and have been working with. I
4	don't see the value. It seems like maybe as
5	we start our report and maybe for the future.
6	But for the moment, it seems to me, if we're
7	going to go over the matrix, we're going to see
8	from what SC&A and NIOSH say that what were some
9	of the issues and what were some of the options.
10	I think that's about what we can do.
11	MR. FARVER: We should talk about
12	it later on and determine what information
13	would be most helpful to you so that we can start
14	applying it in our next, say, 14th through 18th
15	set findings.
16	CHAIRMAN KOTELCHUCK: That makes
17	sense.
18	MR. FARVER: What can we do? We
19	know
20	(telephone connection interrupted)
21	MR. KATZ: in the matrices, you
22	have a lot, you have a lot of material.

1	CHAIRMAN KOTELCHUCK: Well, okay.
2	I mean, I can think of columns it seems to
3	me a good suggestion to say let's start talking
4	about that for 14 through 18, and let's think
5	about that as we move along from 10 to 13. We
6	still got a long way to go. 247, observation
7	one.
8	MR. FARVER: Observation One is,
9	it's the rotational cycle DCFs from IG-001.
10	We've talked about this before. I believe we
11	transferred this to Wanda for her group to look
12	at.
13	CHAIRMAN KOTELCHUCK: Right.
14	MR. FARVER: And according to the
15	NIOSH response, it looks like it's going to be
16	handled under a PER.
17	MEMBER MUNN: Yes, I think that's
18	correct.
19	CHAIRMAN KOTELCHUCK: Okay, good.
20	Let's go number two.
21	MR. FARVER: Okay. Number Two.
22	And I'm not real sure about that one. It was

1	only an observation. It has to do with the
2	ratio of the neutron to photon dose, I believe.
3	Not applicable to the other years because the
4	co-worker dose was used.
5	Okay. I guess our concern was why
6	it was only used one year and not used the other
7	years.
8	And observation three talks about
9	what Scott was talking about, prorating the
10	doses for 2006 - 2007. And what they did
11	technically was kind of unusual. We don't
12	usually see that. They're allowed to do that,
13	to prorate the partial years.
14	CHAIRMAN KOTELCHUCK: Well,
15	actually, you'll excuse me, but I think that was
16	the proper way to handle it. I appreciated
17	what was said because I hadn't thought about
18	that issue, and it seems to me perfectly once
19	the cancer is diagnosed, then the exposure
20	afterward does not count toward the dose that
21	caused that.

MR. FARVER:

22

I understand, except I

1	don't recall seeing it before where they
2	prorated the internal dose. Like in this case,
3	diagnosed in March, so they prorated it for only
4	three months of dose in that final year. I
5	mean, it's correct. I just don't ever recall
6	seeing it before.
7	CHAIRMAN KOTELCHUCK: Yes, yes.
8	MR. SIEBERT: Doug, this is Scott.
9	Yes, we generally don't do that for the problem
10	that we had in this case. But every time you
11	prorate something, you have the option of doing
12	it incorrectly. So if we can just leave it as
13	a full year, generally we will do so. However,
14	it's not unusual, and especially if you've used
15	IMBA to calculate the doses, it only goes
16	through the date of diagnosis.
17	MR. FARVER: Yes, I understand. I
18	just hadn't seen it before for internal.
19	CHAIRMAN KOTELCHUCK: Let's move
20	on.
21	MEMBER RICHARDSON: Before we move
22	on, when you, you've got a recorded dose for a

1	calendar year, right? I mean, going back to
2	the external first, I guess. And you're saying
3	that you take the recorded dose for that year
4	and you take the fraction of the year prior to
5	their diagnosis and apply that? Is that the
6	prorating here for the external case first?
7	MR. FARVER: For the external?
8	MEMBER RICHARDSON: Yes.
9	MR. FARVER: That would be
10	prorating on the, it would be a total dose. And
11	is it a co-worker dose? I'm trying to find it.
12	MR. SIEBERT: Yes, I believe that
13	was co-worker dose. That's why it's prorated.
14	MR. FARVER: Yes, we're taking a
15	co-worker dose for the year and just prorating
16	it for the partial year the employee worked.
17	MEMBER RICHARDSON: Okay, okay.
18	Thank you.
19	MR. FARVER: I got to skip out for
20	a moment and call you right back because my
21	phone is dying. So I'll phone right back in.
22	CHAIRMAN KOTELCHUCK: Okay.

1	John, while he's doing that, why don't you move
2	up to four.
3	MEMBER MUNN: It's awfully quiet.
4	Are we still on?
5	CHAIRMAN KOTELCHUCK: Yes, we're
6	on. We're looking at the screen. I'm looking
7	at the dates, the employment dates here.
8	MR. FARVER: Okay. I'm back.
9	CHAIRMAN KOTELCHUCK: Okay.
10	December 17th, '46 to April '47. And then
11	January '69 to February. Whoa. The dose
12	reconstruction performed using DOL verified
13	employment. So the person who gave the CATI
14	report, the dates would suggest that the person
15	worked all the way through, whereas the
16	employment record indicates that that person
17	didn't work between '47 and '69, right?
18	MEMBER MUNN: We don't have any
19	choice about those things. What the
20	Department of Labor gives us as the employment
21	dates is what we use.
22	CHAIRMAN KOTELCHUCK: Oh, I'm not

1	even, I'm not asking for choice. I would
2	believe, in this case I would believe the
3	records. I have no question that the records
4	are if somebody paid to salary to the person,
5	we'd have records on that.
6	So what is there this was an
7	observation?
8	MEMBER MUNN: Yes.
9	CHAIRMAN KOTELCHUCK: Yes. Well,
10	then there's nothing more to talk about then.
11	MEMBER MUNN: No, there really
12	isn't.
13	CHAIRMAN KOTELCHUCK: No. Let's
14	move on, 248.1.
15	MR. FARVER: Yes, 248.
16	MS. K. BEHLING: Excuse me, Doug.
17	This is Kathy Behling. Before you start 248,
18	can I ask a question?
19	CHAIRMAN KOTELCHUCK: Sure.
20	MS. K. BEHLING: Okay. I'm sorry
21	to interrupt here. I was trying to talk
22	earlier. I guess my mute wasn't working. Can

1	we just quickly tell me, when we went to the
2	Huntington case and we closed that last
3	finding, does that mean that we also closed,
4	that we are not going to have Steve Marschke go
5	back and verify that all of his findings were
6	corrected in the updated TBD, or are those
7	findings still open?
8	I'm a little confused. I just
9	thought that perhaps Steve Marschke could go
10	back and just verify that everything, all of the
11	findings he had were actually updated in the new
12	TBD. Are those findings still open?
13	MR. FARVER: Well, Kathy, my plan was
14	that either Steve or I will go back in and verify
15	that those changes were made.
16	MS. K. BEHLING: Okay.
17	CHAIRMAN KOTELCHUCK: For the
18	Subcommittee, it's closed.
19	MR. FARVER: This is closed, but if
20	something comes up I guess we'll just have to
21	reopen it.
22	MS. K. BEHLING: Okay. I just

1	didn't want that to fall through the cracks
2	because we did say, although NIOSH has assured
3	us that the change has been made, I did think
4	that perhaps either you or Steve could maybe
5	write a brief summary as to what you find when
6	you go into the TBD to ensure that everything
7	has been corrected based on the initial
8	findings.
9	MR. FARVER: Yes. I will or Steve
10	will go back in and look at that and make sure
11	the changes were made.
12	MS. K. BEHLING: Okay, okay.
13	Since we had closed that one finding, I didn't
14	know if we'd go back to the 9th set at all. And
15	I just didn't want that to slip through the
16	cracks.
17	MEMBER CLAWSON: Hey, Doug, this is
18	Brad. I was just wondering, you know, as a
19	Subcommittee Member, can you just write up a
20	little report to us or a little paper letting
21	us know that, you know, we have closed it and

I understand why, but just so that everything

1	was found correct or not?
2	MR. FARVER: Yes, I'll just write a
3	little memo and just say I looked at it or Steve
4	looked at it and it's as expected.
5	CHAIRMAN KOTELCHUCK: Appreciate
6	that. Thank you. Okay. Back to 248.1.
7	MR. FARVER: Okay, 248.1. Okay.
8	Incomplete accounting of recorded dose. Our
9	reviewer saw that there was 115 millirem in the
10	52nd week of 1956 on a dosimetry card that was
11	not included in the dose assignment. But there
12	was 55 millirem from week 52 but not the
13	additional 115. Okay. So there was a little
14	confusion there.
15	Now, this is going to take us down
16	to the bottom there on Exhibit A. I've got the
17	dosimeter card.
18	CHAIRMAN KOTELCHUCK: Okay.
19	Let's go, let's go there.
20	MR. FARVER: For 1956, we kind of
21	see what it's talking about. Let me know when
22	you're on that page, and I'll kind of explain

1	what I can. This is kind of a piece it together
2	so you jump from week 39 and then you get down
3	to week 50 through 52. Are we there?
4	CHAIRMAN KOTELCHUCK: Yes.
5	MR. FARVER: Okay. If you see the
6	39, that's week 39, and that's going to be your
7	end of the third quarter. So that will be your
8	third-quarter doses. Then you can see 50, 51,
9	and 52. If you look under the penetrating
10	column for week 52, you'll see two numbers: 115
11	and the 55.
12	CHAIRMAN KOTELCHUCK: No, I don't
12 13	CHAIRMAN KOTELCHUCK: No, I don't see, I can't quite see it on my screen.
13	see, I can't quite see it on my screen.
13	see, I can't quite see it on my screen.  MR. FARVER: Okay.
13 14 15	see, I can't quite see it on my screen.  MR. FARVER: Okay.  CHAIRMAN KOTELCHUCK: Could you
13 14 15 16	see, I can't quite see it on my screen.  MR. FARVER: Okay.  CHAIRMAN KOTELCHUCK: Could you  lift it up? I think it's just, just
13 14 15 16 17	see, I can't quite see it on my screen.  MR. FARVER: Okay.  CHAIRMAN KOTELCHUCK: Could you  lift it up? I think it's just, just  below there we go.
13 14 15 16 17	see, I can't quite see it on my screen.  MR. FARVER: Okay.  CHAIRMAN KOTELCHUCK: Could you  lift it up? I think it's just, just  below there we go.  MR. FARVER: It's a little
13 14 15 16 17 18	see, I can't quite see it on my screen.  MR. FARVER: Okay.  CHAIRMAN KOTELCHUCK: Could you  lift it up? I think it's just, just  below there we go.  MR. FARVER: It's a little  confusing because you've got these three bottom

1	MR. FARVER: Well, the one, the top
2	part of week 52 where it starts with 180, those
3	are the same doses from the third quarter. So
4	those are the third-quarter doses that are down
5	there.
6	CHAIRMAN KOTELCHUCK: Yes, right.
7	MR. FARVER: Now, underneath that
8	would be, I'm assuming, week 52.
9	CHAIRMAN KOTELCHUCK: No, the
10	bottom is the sum of, the bottom line is the sum
11	of the two on line 52.
12	MR. FARVER: At the very bottom,
13	I'm not sure what the very bottom is.
14	CHAIRMAN KOTELCHUCK: Well, the
15	very bottom is, the very bottom is the sum of
16	the last of the two lines on 52.
17	MR. FARVER: Okay.
18	CHAIRMAN KOTELCHUCK: And the
19	first line is a repeat of week 39 and,
20	therefore, not correct. And, therefore, I
21	would interpret line 52 as the 105, 80, and 55.

1	CHAIRMAN KOTELCHUCK: What do
2	others think?
3	MR. SIEBERT: This is Scott. I can
4	explain what it actually is.
5	CHAIRMAN KOTELCHUCK: Oh, well,
6	I'm sorry. Okay. Pardon me. Alright.
7	MR. SIEBERT: Dr. Kotelchuck,
8	you're exactly right. That top line that's
9	listed in 52 is a repeat of week 39. And what
10	they were doing is they were bringing in the
11	third and fourth quarters for 1956. This is a
12	time frame when they were looking at this doing
13	different dosimetry structures. If we look at
14	1957, which we don't have here, the form is
15	different, and they were in the midst of looking
16	at changing over.
17	So the first one is a repeat of week
18	39, which is quarter three's values. The
19	second row in '52 are the actual week 52, which
20	is really all the fourth quarter results, the
21	150, the 80, and 55. And then the values below

it, as you said, are the totals. So those are

the totals for the third and fourth quarter for 1956.

The question arises with those numbers that are above those that appear to be in week 51, but they're actually really just above week 52. They're not week 51. What we're calling the D data, because this is in pretty much all the 1956 cards for X-10, and there's a little D there and then those values.

It appears what they were doing is they were running some ideas as to what their different values would be with different assumptions in their dosimeters or things of the sort. However, these are not actual dose values for the 51st week. They're just something that the site was using to figure out, to look at how they wanted to change things.

We also went back, this claim and the next claim we're going to talk about, 249, we looked at this issue in both of these claims, and that is what we're seeing with that data. We also went back -- let me back up a second.

What the data entry people enter are what they see on the card, so they actually enter that information in the 51st week with that D designation as it is right now. And it's up to the dose reconstructor to interpret that information, which is what I'm discussing right now.

CHAIRMAN KOTELCHUCK: Right.

MR. SIEBERT: We also went back and we looked at what is the cumulative through the beginning of 1956, adding on 1956 first half of the year and then these values the second half of the year, not the D values but the portions that are at the very bottom, and we compared that to the next year, the 1957 results, which brings forward the cumulative from 1956 back. And in every case that we've looked at, those numbers line up with the totals that are at the bottom of the sheet, not with the D values.

So what we determined is those D values are something the site was doing with their dosimetry, but they're not the doses of

1	record that they were writing in the record.
2	CHAIRMAN KOTELCHUCK: Sounds good.
3	MR. FARVER: Okay.
4	CHAIRMAN KOTELCHUCK: That sounds
5	correct, sounds like it's
6	MR. FARVER: But in this case,
7	those D values were used to calculate dose.
8	MR. SIEBERT: Correct. And at
9	that time, when the dose reconstructor did
10	this, they did not necessarily know that
11	information. And rather than remove anything,
12	they, from a claimant point of view, left that
13	information in.
14	CHAIRMAN KOTELCHUCK: Right.
15	Understandable. That's not an error. Right.
16	That was not an error, but it was a different
17	way of entering the data at the plant level.
18	MR. FARVER: Okay. Now that we
19	understand that, I mean, we're going to see the
20	next case is going to be a little different.
21	But this is what we're looking at where the 115
22	comes from and the 55. Really, our review is

1	looking at the 115 and the 55 for the 52 row and
2	thought that the 115 should be added, did not
3	realize that that was the third quarter. So it
4	was our mistake.
5	CHAIRMAN KOTELCHUCK: Okay.
6	MEMBER MUNN: Any one would be
7	confused. But, yes, I can certainly, given the
8	explanation, it makes sense.
9	MR. FARVER: In this case, there is
10	no mistake.
11	MEMBER MUNN: Right.
12	MR. FARVER: We still have a little
13	data entry concern, but we're going to talk
14	about that on the next case.
15	CHAIRMAN KOTELCHUCK: So this gets
16	closed because it was properly calculated, now
17	that we understand the interpretation, right?
18	MEMBER MUNN: Yes.
19	MEMBER RICHARDSON: I have a
20	question. This is David Richardson. So this
21	is a procedure that NIOSH has developed for the
22	handling of recorded information from the

1	dosimetry cards, and the basis for how you're
2	handling it, is that when you add up the data
3	through 1957, you find that it adds up if you
4	don't include that information?
5	MR. SIEBERT: That is correct.
6	MEMBER RICHARDSON: That's the
7	extent of the basis for deciding how you're
8	going to do this?
9	MR. SIEBERT: That's correct.
10	MEMBER RICHARDSON: And what's the
11	assumption that the 1957 value reflects the
12	cumulative value? I mean, ORNL doesn't have,
13	I mean ORNL actually, I'll start by saying
14	it the other way. ORNL has a lot of information
15	on historical dosimetry practices and
16	recording of doses. There's not documentation
17	saying that the dose of record, as recorded on
18	these dosimetry cards should be handled with
19	this algorithm.
20	MR. SIEBERT: I'm not sure what
21	you're trying to say.
22	MEMBER RICHARDSON: I'm trying to

say this is like a procedure that you're implementing based on an empirical observation from the cards and how you want to sum them, not based on a procedure for how to use the information that was documented some place by the health physics staff who collected and recorded that information? I'm just asking is there anything else, except for the max --

MR. SIEBERT: We have gone through the SRDB references for this time frame, and we have not been able to find any mention of this D data whatsoever.

MEMBER RICHARDSON: So the other thing I was wondering, and this may be helpful for you and maybe you know it and have done it, is Mancuso wrote very detailed guidance to the people when he wrote out protocol for using this information to calculate up the doses. I mean, he re-keyed all this information, as well, and had guidance based on discussions with the health physics staff on site at the time about how to interpret and handle all these dosimetry

cards.

And, I mean, at some place, we have and presumably you have because I think people came and scanned all our documents at one point, that guidance, as well. Is it not there? And under this protocol, does it add up to the value that Mancuso obtained when he keyed this information and summed it up?

MR. SIEBERT: I can't speak for that. Grady, I don't know if you want to speak to this. What I'm guessing is the fact that what we have from the site is the dose of record that we need to use, as opposed to a different study. That's all I can say on that. I mean, that's just what I'm thinking off the top of my head.

MR. CALHOUN: I have no information. I have no personal knowledge of the Mancuso study, other than hearing of it. And we certainly would not use or at least are not likely to use anybody else's interpretation of the data. We try to go back to the original

data.

MEMBER RICHARDSON: Oh, that's
what I'm saying, though. I'm saying you appear
to be interpreting the data of record, as I
took it, was the dosimetry cards. And you're
saying you want to discount a row of the
information not because there's a procedure
written that says discount this row of
information. You've found that you want to
institute a procedure for how to key it and then
interpret it to obtain a dose. And so far what
I've heard is that it's your, this is completely
something you've made up. And that's not bad
or wrong or right, but I'm just saying is there
no other guidance for how to use these dosimetry
cards? And there were people on site who used
them who worked with health physics staff who
wrote down procedures on how to interpret these
cards, not in 2014 but in 1965, which was much
closer to the time.

I just, it sounds like something that you've

And, again, we can let this go, but

1	decided to do relatively recently. And I'm
2	surprised that there's not another basis for
3	this, other than the fact that this seems to
4	make it work.
5	CHAIRMAN KOTELCHUCK: David, are
6	Dr. Mancuso's files available at ORNL? I don't
7	know whether he did this as a staff employee
8	there or
9	MEMBER RICHARDSON: He was
10	CHAIRMAN
11	KOTELCHUCK: researcher outside.
12	MEMBER RICHARDSON: He was
13	employed by the Department of Energy to
14	computerize their Oak Ridge dosimetry data, and
15	he had keypunch people who were doing exactly
16	what the Oak Ridge staff are doing now of keying
17	it. And the final tabulated results, of
18	course, are available to NIOSH has them
19	in-house. But the other, there were
20	procedures on how to use these cards.
21	CHAIRMAN KOTELCHUCK: Right.
22	Okay. So they're available. In terms of

answering my question, those data are available to check? They're available for NIOSH to check?

MEMBER RICHARDSON: Yes. But, you know, I'm just, it seems like it's reinventing the wheel from quite a distance to try and -- I mean, when I was listening to it, it was as though reverse engineering a protocol for interpretation of all these recorded values. And, again, you know, if I was doing this from a research basis, I wouldn't want my method to be we recorded these and then we excluded some, and there's not a historical citation as the basis for why we were or were not including. would feel much more comfortable if I had said, you know, based on documentation from the health physics department, these lines are not empirical readings, lines these were And I would think, more for an notational. epidemiologist's purpose, for a compensation purpose, you would like to be able to say here's the basis for saying that these are not doses

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that we want to include.

MEMBER MUNN: Well, one must also counsel caution in using -- I understand what you're saying with respect to methods that were developed at an earlier date when more familiar people were available for review. By the same token, it's always cautionary to use procedures and methods that were developed by other people for other purposes. I guess it's hard to recommend that we verify or at least calculate any different process because it was used by other people at other times. Well, that's a difficult thing to try to identify.

MEMBER RICHARDSON: Yes. Well, I mean, again, I would say .... I think George Kerr who works with the Oak Ridge staff, maybe still, and they all wrote reports and reports. And Donna Cragle was involved in this and Betsy Dupree, you know, before and after Mancuso, on the difficulties and the procedures in place for interpretation of the historical external dosimetry data and the internal dosimetry data

for the Oak Ridge workers on how to work with those records and interpret them. And it wasn't -- the guidance was based on conversations with the people who had run the dosimetry programs.

So this is another way of doing it, but it's kind of trying to figure out what subtraction leads to a logical summation. But I would say that's the least ideal way of figuring out how to interpret these data, and the best way is to understand the process that led to what values were recorded in what fields.

MEMBER MUNN: I guess, from my perspective, it's worthwhile to know that those studies exist and it's worthwhile to even use them as a part of background information. But to actually go so far as to make point-by-point comparisons of another study wouldn't seem to be appropriate in serving an entirely different use. But perhaps I'm being too specific about it.

CHAIRMAN KOTELCHUCK: I don't

1	think you need to go point by point. It would
2	seem to me that, if the data is available, one
3	would take a few samples out of the data that
4	was keyed in back in '65 and just see that it
5	agrees with what is being said now. What's
6	being said now makes real sense, and the fact
7	that the person put a D there suggests that
8	something is special about this. And
9	MEMBER MUNN: I guess the point I'm
10	trying to make is that we in order to use
11	another study as a point of verification, we
12	have to work on the assumption that that study
13	underwent the same kind of rigorous review that
14	we're giving this particular study and that
15	there were no errors like the ones that we're
16	finding in our own reviews. You understand
17	what I'm saying
18	CHAIRMAN KOTELCHUCK: Yes, yes.
19	MEMBER MUNN: and that it's a
20	good point of reference, but I don't understand
21	that it's necessarily a point of proof.
22	CHAIRMAN KOTELCHUCK: Well, I'd

like to hear a suggestion for resolution.
MEMBER MUNN: Well, let's go back
to what our original finding was.
CHAIRMAN KOTELCHUCK: Right.
MEMBER MUNN: Are we simply trying
to verify that no dose was excluded and that,
since no dose was excluded, we do, in fact, have
a valid basis for making the calculation? Is
that what we're trying to determine? I thought
it was.
CHAIRMAN KOTELCHUCK: But one dose
was excluded. One set of doses was excluded,
and the issue is can we really ascertain that
it was properly excluded, [that this] makes
sense? But it's not verified elsewhere, and it
could be.
MR. FARVER: For this case, the
third-quarter doses, which is on the top level
of row 52, were excluded correctly. They were
not double they were considered once and not
double input. They were input once for the

third quarter, but they were not input again on

1	week 52.
2	MEMBER MUNN: But that's correct.
3	MR. FARVER: That's correct. And
4	the D doses up in row 51, they were included.
5	MEMBER MUNN: Yes, it makes sense
6	to me.
7	MR. FARVER: Okay.
8	CHAIRMAN KOTELCHUCK: So I
9	thought
10	MR. FARVER: Well, where we're
11	going next is, when we get to the next case we're
12	going to see that the D doses were not included.
13	MEMBER RICHARDSON: So the D doses
14	should not have been included, according to the
15	procedure which is being described as
16	implemented today but
17	MEMBER MUNN: Correct.
18	MEMBER RICHARDSON: previously
19	had been included. So, one, it doesn't make
20	sense if we say that's what the D doses are
21	not true doses.
	MEMBER MINING Vo

MEMBER MUNN: Yes.

MEMBER RICHARDSON: So either they were erroneously included there, but they shouldn't have been. But as of now, the decision about their status is kind of, is what appears to make sense to NIOSH as a way of handling the dosimetry information --

CHAIRMAN KOTELCHUCK: You know Right. You are arguing that that's a what? very good sensible explanation to what the D but it really isn't verified data are, Maybe we can resolve it by going elsewhere. back to the principle of being claimant friendly. We could put that in for week 51, and that would be claimant friendly because it would increase the dose. Even though I believe that NIOSH is doing a sensible thing and it's not verified, we could just say put it in because it's claimant friendly.

MR. KATZ: This is Ted. I mean, that just seems arbitrary, Dave. But I don't see why someone can't go look at the Mancuso study and look at what the associated protocols

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1	for handling the data, and see at least whether
2	it diverges from this or if it's consistent with
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4	CHAIRMAN KOTELCHUCK: Well, Wanda
5	is saying that who knows how well the Mancuso
6	study was
7	MR. KATZ: But that matters not. I
8	mean, you can look at it and see what they did
9	and at least know if that's consistent or not,
10	and then at least you know something more than
11	we know now. And perhaps it's perfectly
12	consistent, in which case there is no more
13	discussion, because that's then, in effect,
14	supporting what
15	CHAIRMAN KOTELCHUCK: That's
16	supporting evidence, yes.
17	MR. KATZ: I mean, why not someone
18	look instead of I mean, it seems like we're
19	beating our heads on something that could use
20	some more information.
21	MR. CALHOUN: This is Grady. I
22	don't know. To me, it seems like the better

approach may be to be better define why we're doing what we're doing. Unless we buy in completely to this Mancuso study, comparing us against what they did is worthless.

So, you know, maybe we just need to better define and talk to our site experts who wrote our documents and find out why we're doing what we're doing, and at least be able to describe it a little bit more in detail. I think that's a better first step than trying to compare something to a document that we're not sure that we buy into 100 percent either. And maybe we do. I just am not knowledgeable --

CHAIRMAN KOTELCHUCK: Well, on the other hand, one could say this: What NIOSH has described makes sense for interpretation of the D values. If another study confirms it, that's evidence. If the other study does not agree with that, then there is not supporting evidence, and you folks can continue to think about, you know, whether this is right or not and come back and say, well, thinking about it

1	more. We still believe it's correct and Mancuso
2	was wrong.
3	But it is possible I mean, what
4	I'm saying is an affirmation from Mancuso=s
5	data is evidence. A lack of agreement is not
6	necessarily evidence. It doesn't confirm what
7	you've done.
8	MR. CALHOUN: I am certainly not
9	ready to commit to doing that. I'll talk to our
10	management here and see what they think about
11	that. I'd be much more apt to try to better
12	describe what we do than compare our work to
13	another study. So that's really where I'd like
14	to start it.
15	CHAIRMAN KOTELCHUCK: Well, okay.
16	Let me ask you this: do you think it would be
17	a big job to find Mancuso's data and check it
18	out?
19	MR. CALHOUN: I have no idea.
20	CHAIRMAN KOTELCHUCK: I have no
21	idea either. I mean, I respect that yours may
22	be better than his. I don't know. He's a

1	respected academic researcher, but on the other
2	hand, you folks are good researchers and well
3	trained, you know, and experienced. So I'm not
4	saying one is better than the other.
5	(Simultaneous speaking.)
6	MR. CALHOUN: the topic right
7	now, to say one thing one way or another, you
8	know, I always - I'm quite confident in the work
9	we do here. But I'm certainly willing to go
10	back and look at it to make sure that it really
11	does make sense.
12	CHAIRMAN KOTELCHUCK: Well, that's
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14	MEMBER MUNN: Unless there's
15	another study that specifically mentions the D
16	data here, then it's probably futile. But how
17	to know whether another study does mention that
18	is almost impossible, unless everything is
19	computerized and you can word search for such
20	a thing as D data.
21	It seems very obvious. You know,
22	some things are obvious. You look at these

obvious things: 180, 135, 115. Clearly, that is simply drawing down the figures from the third quarter. That's not a mystery. D data, if you look at the D data, then you'd have to assume, if there's any validity at all to that, that every single aspect of dose increased radically over the course quarter, which seems unlikely somehow. And the addition figures below are obvious. know, you can probably eliminate the obvious things, which leaves only the question of: and exactly what was that D data stuff? But it clearly was not the readings for the fourth It would not have been identified in quarter. that way.

CHAIRMAN KOTELCHUCK: Well, I will say we've got a lot of cases to cover. We have cases that are open that we're going to have to come back to. I suppose we could just simply leave this open and let the folks at NIOSH and SC&A reconsider the evidence that exists and discuss with others there about whether there's

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1 access to the Mancuso data and whether they want to look at it. 2 3 Clearly, what I'm thinking is we're not going to resolve this right now, and we are 4 So I'd 5 going to have to come back to this data. 6 certainly be willing to give the NIOSH folks a chance to re-look, rethink, based on this 7 discussion and then come back to us. 8 9 MR. FARVER: This is Doug. I quess 10 I'm sorry I brought this up because my concern 11 was far more basic. For Tab 248, they included 12 the D data in their calculations. We go down to Tab 249, they did not include it. 13 It was not in the dosimetry input file. 14 So even though it's on the card, it's not in the dosimetry 15 16 input file. It wasn't keyed in. So I have a Two people are looking at 17 quality concern. this and they're interpreting it differently. 18 19 CHAIRMAN KOTELCHUCK: That's 20 concern. MR. FARVER: I don't know what it is 21 22 or if it should or shouldn't be used, but at

1	least be consistent.
2	CHAIRMAN KOTELCHUCK: Right.
3	MEMBER MUNN: Agree.
4	MR. FARVER: So that's my concern.
5	It's in the dosimetry input file
6	MR. SIEBERT: Wait, wait, wait.
7	This is Scott. I'd like to clarify that. That
8	was not necessarily correct. That D data is
9	listed in the data entry file for both of those
10	cases. The dose reconstructor in the second
11	case decided to remove it based on this process
12	of saying that the cumulative doses did not
13	match the record. However, the data entry
14	people entered it the first time and the dose
15	reconstructor made the decision to remove it.
16	So it is not a data entry issue.
17	They were both done consistently.
18	MR. FARVER: Okay. So now we have
19	inconsistencies among the dose reconstructors.
20	MR. SIEBERT: Which we agree
21	wholeheartedly that we need to look at that, and
22	we're documenting that process already. But,

1	you know, I know Grady is going to talk to his
2	management.
3	MEMBER MUNN: Did I understand
4	incorrectly that this occurs only during this
5	brief period of time, which, if I think I heard
6	what you said earlier, was a transition time
7	from one reporting method to another?
8	MR. SIEBERT: That is correct, the
9	end of 1956.
10	MEMBER MUNN: This is the only time
11	we see this D data, right?
12	MR. SIEBERT: Correct.
12 13	MR. SIEBERT: Correct.  MEMBER MUNN: So it's not that big
13	MEMBER MUNN: So it's not that big
13 14	MEMBER MUNN: So it's not that big an issue. We have a few figures here at the
13 14 15	MEMBER MUNN: So it's not that big an issue. We have a few figures here at the tail-end of one type of reporting, as they're
13 14 15 16	MEMBER MUNN: So it's not that big an issue. We have a few figures here at the tail-end of one type of reporting, as they're moving into a different mode of reporting data.
13 14 15 16 17	MEMBER MUNN: So it's not that big an issue. We have a few figures here at the tail-end of one type of reporting, as they're moving into a different mode of reporting data.  CHAIRMAN KOTELCHUCK: Right.
13 14 15 16 17 18	MEMBER MUNN: So it's not that big an issue. We have a few figures here at the tail-end of one type of reporting, as they're moving into a different mode of reporting data.  CHAIRMAN KOTELCHUCK: Right.  MEMBER MUNN: On this case, for
13 14 15 16 17 18	MEMBER MUNN: So it's not that big an issue. We have a few figures here at the tail-end of one type of reporting, as they're moving into a different mode of reporting data.  CHAIRMAN KOTELCHUCK: Right.  MEMBER MUNN: On this case, for example, we have three numbers. Gee, the

is a logical view and we're talking about very 1 small numbers here. It seems irrational for us 2 3 to contribute enormous amounts of time to worrying about these three numbers. 4 Well, the 5 CHAIRMAN KOTELCHUCK: three numbers, right, the three numbers are 6 I mean, the three numbers here 7 three people. are one person, but I wouldn't want to -- for 8 that one person, it's very important. 9 10 MEMBER MUNN: But even if those 11 three numbers appeared on every single solitary 12 report that we have, which I think is unlikely, even if it did appear, my point is, given the 13 records that exist for each individual employee 14 and the fact that this occurs only during this 15 last set of data recording sheets prior to the 16 institution of new reporting system, I don't 17 see -- it seems to me we're making a "I love a 18 19 mystery" out of a molehill. 20 CHAIRMAN KOTELCHUCK: Okay. Suppose you believe that. What conclusions 21

do you draw from that last statement?

conclusion, 1 MEMBER MUNN: Му personally, is that, for this individual, the 2 3 numbers that were drawn from the dose for this last fourth quarter of this year are 105, 30, 4 and 55. That's my conclusion, regardless of 5 6 whether the numbers are written on there. It's in line with -seems obvious to me. 7 MEMBER CLAWSON: This is Brad. Ι 8 can't believe -- I do not agree with that at all. 9 That D data is there for a reason. 10 If you can't tell me why it isn't there, then it's either 11 going to be put in there or we're going to figure 12 This arbitrarily deciding, 13 out what it is. 14 yeah, we don't need to worry about this data, it doesn't really mean -- I don't think that 15 really says that we're doing a very good job 16 And quess what? It may only be for 55. 17 But this is for one person out there. 18 19 I was that one person or a family member or something else, I guess it would matter to me. 20 The thing is, we're trying to make 21

this as clear as possible. And we want to be

1 able to be able to explain why we do what we do. To be able to hear that, well, we're just 2 ignoring it because it really doesn't matter, 3 I don't buy that. I don't think that's right. 4 5 CHAIRMAN KOTELCHUCK: I think we don't have much -- I think we need to let NIOSH 6 people rethink, based on this discussion, not 7 mandate anything other than ask you folks to 8 reconsider. If you want to talk to SC&A, fine. 9 10 That's up to you. And then come back to us with a resolution of this and the next case, your 11 12 suggested resolution of it and your whatever, and the body of evidence you're using to decide 13 it, although I think you're giving it to us. 14 But you may want to think about it a little bit 15 more to make us feel confident that what you 16 said was true. 17 Right now, we're not assured that 18 19 your D data - we're not assured of what you said about the D data, although it certainly, to me, 20 seems to make sense. 21

Can we do that? Leave this open and

1	let you come back?
2	MR. SIEBERT: That's okay with me.
3	CHAIRMAN KOTELCHUCK: And think
4	about it. I mean, it's something that sort of,
5	if you will, blew up, I mean, in terms of a
6	bigger issue than maybe we thought it was going
7	to be. And I gather that will also include the
8	next what is it 249, where the same issue
9	comes up.
10	Then if that were the case, it is now
11	if we agree on that, then it is 3:20. We've
12	been going about two hours. I will propose a
13	10-minute break from 3:19 to 3:30, 11 minutes.
14	Would people like to do that?
15	MEMBER MUNN: Yes.
16	MR. KATZ: Good idea.
17	CHAIRMAN KOTELCHUCK: Okay. And
18	then we'll start back on is there anything
19	else on 246? Are there other I can't
20	MR. FARVER: There's 248. We
21	still have a couple of findings.
22	CHAIRMAN KOTELCHUCK: Okay.

1	MEMBER MUNN: We're on 248, right?
2	CHAIRMAN KOTELCHUCK: We are? I
3	can't read it? We're on 248, yes.
4	MR. FARVER: And there's two more
5	findings and a couple observations.
6	CHAIRMAN KOTELCHUCK: Okay. Then
7	we'll come back to the rest of 248.
8	MEMBER MUNN: Great. Thank you.
9	CHAIRMAN KOTELCHUCK: Okay.
10	Thank you. See you folks at 3:30.
11	(Whereupon, the foregoing matter
12	went off the record at 3:20 p.m.
13	and went back on the record at
14	3:30 p.m.)
15	CHAIRMAN KOTELCHUCK: 248.1 is
16	open. Let's go to 248.2. Doug?
17	MR. FARVER: 248.2. The reviewer
18	concludes that the employee should have been
19	assigned missed neutron doses for the years '53
20	through '55. And this was at X-10, I believe.
21	Or, no, it could have been Y-12 for these years.
22	He worked at different places.

1	And they base that on that the
2	employee=s files contain some NTA neutron
3	monitor film results for the last half of '53,
4	all of '54, and three weeks in '55. All the
5	results were zero, with the exception of three
6	badges which had one track edge each. So based
7	on this, we feel they should have assigned
8	missed neutron doses. And NIOSH
9	agrees,unmonitored neutron doses should have
10	been applied.
11	CHAIRMAN KOTELCHUCK: Close.
12	MR. FARVER: Yes.
13	CHAIRMAN KOTELCHUCK: And it will
14	be done?
15	MR. FARVER: That I don't know.
16	MR. SIEBERT: This is another one
17	that, since it's X-10 and there is an X-10 PER
18	coming down the pike, it will be addressed when
19	we reassess under the PER.
20	CHAIRMAN KOTELCHUCK: Okay. But
21	it will
22	MEMBER MUNN: Since we have an

1	obvious that's been called to our attention and
2	NIOSH agrees that a dose is overlooked, doesn't
3	that automatically trigger a rework?
4	MR. CALHOUN: It would only trigger
5	a rework if it was automatically comped, so I
6	don't know if they've actually looked at it to
7	that level yet.
8	CHAIRMAN KOTELCHUCK: Right.
9	MR. SIEBERT: We have not looked at
10	it based on present-day standards and reworking
11	it that way.
12	CHAIRMAN KOTELCHUCK: But I think
12	CHAIRMAN KOTELCHUCK: But I think from the Subcommittee's point of view, I think
13	from the Subcommittee's point of view, I think
13 14	from the Subcommittee's point of view, I think this should be closed. It'll be done.
13 14 15	from the Subcommittee's point of view, I think this should be closed. It'll be done.  MEMBER CLAWSON: Yeah. This is
13 14 15 16	from the Subcommittee's point of view, I think this should be closed. It'll be done.  MEMBER CLAWSON: Yeah. This is Brad. I guess my question is: How do we know?
13 14 15 16 17	from the Subcommittee's point of view, I think this should be closed. It'll be done.  MEMBER CLAWSON: Yeah. This is Brad. I guess my question is: How do we know?  Is it going to be corrected or what?
13 14 15 16 17 18	from the Subcommittee's point of view, I think this should be closed. It'll be done.  MEMBER CLAWSON: Yeah. This is Brad. I guess my question is: How do we know?  Is it going to be corrected or what?  MR. KATZ: This is Ted. This is
13 14 15 16 17 18	from the Subcommittee's point of view, I think this should be closed. It'll be done.  MEMBER CLAWSON: Yeah. This is Brad. I guess my question is: How do we know?  Is it going to be corrected or what?  MR. KATZ: This is Ted. This is just standard. This has been since the

1	address that. Otherwise, they don't
2	necessarily redo a DR because there's an error
3	in it if it's not going to change the outcome
4	of it.
5	So I think what Grady just said was
6	someone will look at it to see if this is, how
7	much impact this would have. But, otherwise,
8	then it would get addressed under the PER. And
9	that's just normal course. That's always been
10	the case.
11	MEMBER RICHARDSON: So was this
12	claimant close to a threshold already?
13	MR. CALHOUN: Well, pretty much all
14	of them that you guys look at now are.
15	MR. FARVER: Seven percent.
16	MR. CALHOUN: Three percent of all
17	the cases that we have in-house are between 45
18	and 52 percent.
19	MEMBER RICHARDSON: Yeah, because
20	they already had, like, 20 rem to the bladder,
21	right? And they're adding another rem or two?
22	CHAIRMAN KOTELCHUCK: Yeah.

1	MEMBER MUNN: Barely over one.
2	CHAIRMAN KOTELCHUCK: Look, this
3	is going to be when the PER comes out, all
4	of these will be looked at. And it's not even
5	a question that they will be looked at. So
6	MR. CALHOUN: But here's the deal.
7	And I hate to commit us to do any more work here,
8	but a lot of the times when we make our defense
9	of these things we say, well, it wouldn't affect
10	the PoC. So I think at least we need to look
11	at this to see if it affects the PoC. If it
12	affects the PoC and makes it look like it will
13	flip, we'll ask for a rework. But if it
14	doesn't, we won't.
15	MR. KATZ: Right. That's what I
16	was trying to say
17	CHAIRMAN KOTELCHUCK: That's
18	right.
19	MR. KATZ: was, I thought, the
20	normal course.
21	MR. SIEBERT: Let me just interrupt
22	because this is something I should have known

1	before you guys were talking about it, and I
2	apologize. This claim was reworked in 2009 and
3	compensated based on additional cancer
4	information. I apologize.
5	CHAIRMAN KOTELCHUCK: Okay.
6	You'll put that in the SRC action or somewhere.
7	You'll put it in. But it is closed. 48.3.
8	MR. FARVER: 48.3.
9	CHAIRMAN KOTELCHUCK: Wait a
10	minute. 248 is already compensated, so what
11	are we looking at three for?
12	MEMBER MUNN: Because we still have
13	to clear the item.
14	CHAIRMAN KOTELCHUCK: That's
15	right. Okay. And it may reveal a procedural
16	mistake that might be affecting other people.
17	MEMBER MUNN: But NIOSH and SC&A
18	agree and it has already been compensated, so
19	it's correctly closed.
20	CHAIRMAN KOTELCHUCK: That sounds
21	right.
22	MR. FARVER: This is another QA

where they misapplied the OTIB-49 correction factor. That's for Super S plutonium. They should have applied beginning in 1953, and it would have increased the employee=s doses. But they didn't apply it until 1963. Once again, that's something that you should catch in a peer review. You know, you're off by ten years.

MEMBER MUNN: But this is Yes. emblematic of what we were discussing earlier today when we were talking about viewing these things in their aftermath to identify what category they fall into. The wording here is It's a QA review issue, and it appropriate. should have been caught. It wasn't, but the closure says it's a QA issue. So any time we're reviewing it in the future, we'll understand And if we're tallying the kinds of that. errors that we find, the kinds of findings that we wish to pass on, then this clearly tells us that this is QA and that's what we wanted to do. We've got it, we understand it. The client has

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1	been compensated, and this item appears to be
2	closed and properly identified as QA.
3	CHAIRMAN KOTELCHUCK: Agreed.
4	MR. FARVER: Okay.
5	CHAIRMAN KOTELCHUCK: Observation
6	1.
7	MR. FARVER: Observation 1. The
8	short story is, when you look in the CATI report
9	and you look at what buildings the employee
10	worked in, he says he worked in one building in
11	Y-12, 9735. And when we went to verify the
12	interview distribution, we could not find that
13	building listed anywhere in the Y-12 or X-10
14	TBDs.
15	So that's pretty much what prompted
16	the observation was, well, we don't know what
17	it should be because we hadn't found this
18	building listed anywhere. The building that
19	was used
20	CHAIRMAN KOTELCHUCK: Are we doing
21	248, Observation 1?
22	MR FARVER: Vec 248 Observation

1	1.
2	CHAIRMAN KOTELCHUCK: I don't see
3	anything about the building.
4	MR. FARVER: Well, it was
5	CHAIRMAN KOTELCHUCK: Oh, okay.
6	MR. FARVER: It wasn't included in
7	theThe full observation is kind of lengthy,
8	and it wasn't all included there.
9	CHAIRMAN KOTELCHUCK: Oh, okay,
LO	alright. Okay.
L1	MR. FARVER: We couldn't find the
L2	building to verify the energy distribution, is
L3	what it comes down to. It wasn't a big deal
L4	because the one they used was okay.
L5	CHAIRMAN KOTELCHUCK: Alright.
L6	MR. FARVER: So we just wanted to
L7	point out that that building was not in any of
L8	their documents.
L9	MEMBER MUNN: So we have a phantom
20	building, but we have an overestimation in any
21	case.

MR. FARVER: Yes.

1	MEMBER MUNN: And so the
2	observation has been duly noted and been
3	properly evaluated. There's no additional
4	information we can add, so this observation is
5	closed.
6	CHAIRMAN KOTELCHUCK: Right.
7	Well, it's an observation, so, okay. Next?
8	MR. FARVER: The next one has to do
9	with some missed dose. When we were going
10	through the records, we came up with three
11	additional missed doses for '51 and three
12	additional missed doses for '53, based on what
13	was written in the margins of the dosimetry
14	records. It really wasn't going to affect
15	anything, so we didn't make it a finding.
16	Dose-wise, it's like 80 millirem, so it's not
17	very significant, which is probably why it was
18	just written up as an observation. And NIOSH
19	has replied there were visitor dosimeters same
20	week as the permanent, so they used the

CHAIRMAN KOTELCHUCK: Okay.

permanent.

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1	MR. FARVER: Observation 3,
2	incorrect MDA values were used for strontium-90
3	and uranium. Okay.
4	CHAIRMAN KOTELCHUCK: 249.1.
5	MR. FARVER: Well, 248,
6	Observation 3, basically used the incorrect MDA
7	values. They used the highest MDA values in
8	the TBD. They may err on the high side. So
9	it's an overestimating method that would not be
10	used today. We didn't write it up in the
11	finding because it was an overestimate.
12	CHAIRMAN KOTELCHUCK: Right.
13	MR. FARVER: 249.1, incomplete
14	accounting of reported dose. This takes us
15	back to 248.1. So we go down to our exhibit.
16	We go down to Exhibit B. And if you look at
17	Exhibit B, it looks very similar to Exhibit A,
18	except in this case the D values were not used
19	in the calculation. I have there that they
20	were not included in the dosimetry input file.
21	Scott says that's incorrect. And I haven't

been able to get to that file to check that, but

Τ	I Will.
2	But, in any case, the doses, in this
3	case, were not used. So it could be the dose
4	reconstructor decided not to use them, where
5	the previous ones in 248 decided to use them.
6	So in either case, they have an inconsistency.
7	MR. CALHOUN: I'm actually going to
8	try to get an answer for you by tomorrow on that.
9	I just talked to Tim, and this is not some big
10	secret. So this is something that we can come
11	up with that has to do with the change in
12	dosimeters and how they were read, and our
13	preliminary answer is the D data should not be
14	used. But we're going to get you a more
15	CHAIRMAN KOTELCHUCK: I would love
16	that. We would love that.
17	MR. CALHOUN: I would love it more.
18	CHAIRMAN KOTELCHUCK: Okay, good.
19	MR. FARVER: The basis for this
20	finding was that in one case they used the data,
21	in this case they didn't use the data.
22	CHAIRMAN KOTELCHUCK: Right.

1	MR. FARVER: And they should be
2	consistent.
3	CHAIRMAN KOTELCHUCK: Absolutely.
4	We all agree.
5	MR. FARVER: It's a quality
6	concern. Whether it's a data entry or dose
7	reconstructor, it's a quality concern.
8	MEMBER RICHARDSON: Doug, the page
9	scrolled down. I was trying to check on the
10	phantom building. Is it 9203?
11	CHAIRMAN KOTELCHUCK: 9203, right.
12	MEMBER RICHARDSON: Yes. So it
13	does exist in the Department of Labor's Site
14	Exposure Matrix. It's a Y-12 building called
15	Laboratory Developments Facility.
16	MR. FARVER: Right. That wasn't
17	the phantom building. The phantom building
18	was 9735.
19	MEMBER RICHARDSON: Oh, 9735.
20	Okay. Which is the Research Services
21	Laboratory at Y-12.
22	MR. FARVER: Okay.

1	MEMBER RICHARDSON: I mean, as an
2	observation, it might be useful if the
3	Department of Labor and NIOSH agreed on what
4	buildings exist.
5	I mean, if the problem is arising
6	from it being I don't know how many I mean,
7	there's a lot of buildings at these sites. But
8	if there's a problem with them not agreeing, I
9	think that DOL has done quite a bit of work to
LO	make an index of the buildings at the facilities
L1	and what hazards are there.
L2	MR. FARVER: Right. It just
L3	wasn't in any of the Y-12 or X-10 documents that
L4	we saw, that building.
L5	Where are we? Oh
L6	MEMBER RICHARDSON: The other good
L7	part is that it shows I mean, this building
L8	was reported in CATI. Is that where it was
L9	coming from?
20	MR. FARVER: Yes, the employee
21	information?
22	MEMBER RICHARDSON: Yeah. So they

1	weren't making up buildings. It was there.
2	CHAIRMAN KOTELCHUCK: Well, that's
3	good. Yes. Okay.
4	MR. FARVER: How do you want to
5	handle 249.1, where at one time they used the
6	D data and in this case they didn't?
7	CHAIRMAN KOTELCHUCK: Yeah, that's
8	open, and we're going to hear tomorrow.
9	MR. FARVER: Okay.
10	CHAIRMAN KOTELCHUCK: Something, I
11	hope.
12	MR. CALHOUN: You'll hear
12 13	MR. CALHOUN: You'll hear tomorrow. That's my goal.
13	tomorrow. That's my goal.
13 14	tomorrow. That's my goal.  CHAIRMAN KOTELCHUCK: Okay.
13 14 15	tomorrow. That's my goal.  CHAIRMAN KOTELCHUCK: Okay.  Let's hope. So right now that will be open.
13 14 15 16	tomorrow. That's my goal.  CHAIRMAN KOTELCHUCK: Okay.  Let's hope. So right now that will be open.  How about 249.2?
13 14 15 16 17	tomorrow. That's my goal.  CHAIRMAN KOTELCHUCK: Okay.  Let's hope. So right now that will be open.  How about 249.2?  MR. FARVER: 249.2. I got to close
13 14 15 16 17 18	tomorrow. That's my goal.  CHAIRMAN KOTELCHUCK: Okay.  Let's hope. So right now that will be open.  How about 249.2?  MR. FARVER: 249.2. I got to close  one and get the other open. Hang on.
13 14 15 16 17 18 19	tomorrow. That's my goal.  CHAIRMAN KOTELCHUCK: Okay.  Let's hope. So right now that will be open.  How about 249.2?  MR. FARVER: 249.2. I got to close  one and get the other open. Hang on.  CHAIRMAN KOTELCHUCK: Sure.

1	CHAIRMAN KOTELCHUCK: Sure.
2	MS. K. BEHLING: Grady, you had
3	made mention that to get a PER out it may be a
4	year. I was just curious as to why the length
5	of time for some of the PERs. I'm thinking
6	along the lines of
7	MEMBER MUNN: I'm not hearing you
8	well, Kathy.
9	MS. K. BEHLING: Okay. I'm sorry.
10	Is that any better?
11	MEMBER MUNN: Yeah, lots better.
12	Thanks.
13	MS. K. BEHLING: Okay. I'm sorry.
14	I was just thinking along the lines of like the
15	IG-001 where that table 4.1(b) that we keep
16	talking about. I mean, it's four different
17	types of cancers. It's pretty specific. And
18	I hope I'm not asking a naive question, but I'm
19	just wondering what takes so long to get
20	something like that, a PER, issued.
21	MR. CALHOUN: Any time a Technical
22	Basis Document is changed, even a millirem, for

whatever reason, whether it's our discussions 1 here or anything else, we have to do a PER. 2 3 MS. K. BEHLING: Right. MR. CALHOUN: So that is literally 4 5 thousands and thousands and thousands 6 claims. So we've got dozens of PERs, at least 7 tens of PERs in the system right now that we have planned to do. And we do those, and they come 8 over, every week we get evaluations that come 9 10 But the sheer magnitude of them is very over. 11 And when we have documents that are limiting. 12 being reviewed by you guys or whoever and 13 they're not done, we're not going to do the PER until they're complete. 14 So the PER is initiated once the 15 document that drives it is signed, approved and 16 when we have back and forth 17 done. So discussions amongst ourselves and ORAU, back 18 and forth between you guys, the document is not 19 done, so we don't do the PER. 20 21 MS. K. BEHLING: Okay. Yeah, I was

just trying to get a better understanding of --

about that there may be cases that would be perhaps overturned, so I was just trying to get a better understanding of what --

MR. CALHOUN: Well, the ones that we really quickly are the ones that aren't PERs, but they're PADs, and that's when we actually receive a new piece of data, such as a new dosimetry file. Then we can just do that case according to whatever document exists. But we can't do a PER unless the document that drives it is approved because you've got to be doing the dose reconstruction through an approved document. So that's what pushes those back.

And we will receive, probably, I'm going to guess and say a hundred a month we receive of individual cases that were reviewed for one reason or another. Well, the reason for PERs is always because there's been a dose increase of some sort in one of the driving documents.

MS. K. BEHLING: Okay. Thank you.

1	I appreciate that explanation.
2	CHAIRMAN KOTELCHUCK: Doug, are
3	you
4	MR. FARVER: I'm here.
5	CHAIRMAN KOTELCHUCK: Okay.
6	MR. FARVER: I'm looking at the
7	table at the moment. Let's see.
8	CHAIRMAN KOTELCHUCK: Wait a
9	minute. Did we do 249.2? We just scrolled up
10	to Observation 1.
11	MR. FARVER: 249.2 is what we're
12	working on right now.
13	CHAIRMAN KOTELCHUCK: That's
14	right. Okay.
15	MR. FARVER: Incomplete accounting
16	of medical x-ray dose.
17	CHAIRMAN KOTELCHUCK: Okay.
18	Hopefully, John, if you might run the screen
19	down now.
20	MR. FARVER: The finding has to do
21	with we didn't see where they assigned a
22	pre-placement PFG exam. When we looked at the

1 documents and when we looked at the workbooks, it appeared that things were listed as PA when, 2 3 according to the TBD, it should have been a PFG, 4 okay? However, for this case, and for the 5 6 ovaries, at this time period, the dose is the 7 same. We're still CHAIRMAN KOTELCHUCK: 8 not at 249.2. Pardon me. Yes, thank you. 9 10 MR. FARVER: So, a chest PA is the 11 same as a stereo PFG for the prior-to-1947, which is 25 millirem. 12 Instead of assigning 13 both the PFG and a PA in the same year, they 14 assigned the one. It shows up in the workbook as a PA, but it doesn't really matter in this 15 case because the PFG and the PA exams have the 16 But the finding was because we 17 same dose. looked at the workbook and saw PA and didn't see 18 a PFG exam was considered. 19 That's why we wrote the finding. And the big picture, it doesn't 20 matter because the doses are the same. 21

KOTELCHUCK:

CHAIRMAN

22

Comments,

1	anybody?
2	MEMBER MUNN: It sounds like an
3	observation.
4	MR. FARVER: Well, it would have
5	been a finding if it was correct, but it's more
6	like a
7	CHAIRMAN KOTELCHUCK: Well, it
8	appears then that we should close it.
9	MR. FARVER: Okay.
10	CHAIRMAN KOTELCHUCK: Okay.
11	Let's go now to the observation.
12	MR. FARVER: Observation 1, NIOSH
13	did not reference where the less than 30 keV
14	photon DCF of approximately 0.2 originated
15	from, nor did they use the special plutonium DCF
16	provided in OCAS-IG-001. It appears that the
17	method that NIOSH used to determine was based
18	on OTIB-12, and it's not used anymore. This
19	really didn't have a big impact. It was more,
20	where did you get this number from?
21	CHAIRMAN KOTELCHUCK: Right. And
22	we know now.

1	MR. FARVER: Now we know, but now
2	it's not going to matter because they're not
3	using [it] anymore.
4	CHAIRMAN KOTELCHUCK: Okay.
5	Let's go to 2.
6	MR. FARVER: Two. This was just to
7	point out, it seems to me, a little
8	inconsistency between PROC-61 and the
9	technical basis related where PROC-61 really
10	doesn't mention PFG exams for X-10 in their
11	Table 1, whereas the information is mentioned
12	in the technical basis, TBS-12-3.
13	So you get a little confused if the
14	dose reconstructor goes to PROC-61 instead of
15	the TBD and he may not include the exam. So
16	that was just to point out this little
17	inconsistency.
18	CHAIRMAN KOTELCHUCK: Okay,
19	alright.
20	MR. FARVER: And in NIOSH's
21	response, they say they do have guidance. The
22	guidance was in both of the documents about

1	PFGs. And you can see in our response, it was
2	just a different in revisions in PROC-61.
3	CHAIRMAN KOTELCHUCK: Okay.
4	MR. FARVER: Observation 3.
5	MR. SIEBERT: You know, I just want
6	to point out that we did do what was correct for
7	the documentation that was in place at the time.
8	MR. FARVER: Correct.
9	MR. SIEBERT: Okay.
10	MR. FARVER: Observation 3. SC&A
11	could not find any rationale for using a value
12	of a thousand times the environmental iodine
13	intake where the TE-132 intake [is] 10 percent
14	of the iodine intake for this incident okay.
15	A little more interesting.
16	MEMBER MUNN: That sounds like a
17	decimal point that really got moved.
18	MR. FARVER: I'm still not sure
19	that there was a rationale for the thousand, but
20	in the big scheme of things, there's no dose
21	anyway.
22	CHAIRMAN KOTELCHUCK: I wondered

1	why this wasn't a finding, rather than an
2	observation.
3	MR. FARVER: Probably because
4	there was no dose.
5	CHAIRMAN KOTELCHUCK: There was no
6	dose of iodine? There was.
7	MR. FARVER: There was not a dose,
8	the dose resulted in very small doses. There
9	was not a significant dose. Just what's the
10	basis for a thousand, I don't know. Scott, do
11	you have any input?
12	MR. SIEBERT: It's just
13	professional judgment that the dose
14	reconstructor was coming up with an
15	overestimate and was trying to address the
16	issue and showing it still had no dose [effect]
17	when he used the large number. So I can't tell
18	you the specific reason.
19	CHAIRMAN KOTELCHUCK: Okay.
20	MR. FARVER: Now, I can see if they
0.1	11
21	used a thousand times an intake and it comes up

1	as a finding.
2	CHAIRMAN KOTELCHUCK: Right.
3	MR. FARVER: Okay. So we don't
4	know where it came from. Just judgment.
5	Okay. Observation 4.
6	MEMBER RICHARDSON: This is dose to
7	the bladder, is that right?
8	MR. FARVER: Yes, yes. We noted
9	that the Type SS plutonium was found to deliver
10	the most dose using the bioassay results.
11	However, Type M provided the most dose using the
12	coworker data, so we just noted the difference.
13	And in the response, the solubility type
14	supplied in the dose reconstruction matched the
15	type discussed in the observation, Type Super
16	S for '49 to '61 and then Type M for '62 to '82,
17	based on most claimant-favorable. I think
18	we're just noting there that it would be a
19	solubility change, you know, that they were
20	using, but they were using the most

CHAIRMAN KOTELCHUCK:

21

22

claimant-favorable.

Alright.

1	MR. FARVER: Okay. And that wraps
2	up that case.
3	CHAIRMAN KOTELCHUCK: I think 268
4	is the next one.
5	MR. FARVER: 250.
6	CHAIRMAN KOTELCHUCK: 268? Oh,
7	I'm sorry, excuse me. We have to discuss it and
8	suggest that it's closed.
9	MR. FARVER: Oh, okay, 250.1. A
10	PFG examination was most likely used for
11	pre-employment. When we were reviewing the
12	TBDs, each of the TBDs for Y-12, X-10 and K-25,
13	they all state that PFG equipment was most
14	likely used for pre-employment examinations
15	during the time period in question. So we felt
16	that they should have assigned a PFG dose
17	instead of a PA dose for your pre-employment.
18	And they do give an explanation, and they were
19	following their guidance.
20	However, given the time period
21	between '44 and '45, we still felt that they
22	should have used a PFG exam, because in Oak

1	Ridge at that time they were all using PFG
2	equipment.
3	MR. SIEBERT: Hey, Doug, this is
4	Scott. I mean, I think people might be
5	digesting that answer. We agreed with that,
6	actually, which is why when OTIB-52 was updated
7	it reflected the fact that you should, for those
8	larger sites like that, you should use the
9	default x-rays at the site of interest, rather
10	than assuming they may have occurred offsite.
11	So that did get changed in OTIB-52 because just
12	for what you're saying. That makes more sense.
13	MR. FARVER: So that has been
14	changed since this?
15	MR. SIEBERT: Yes, Revision 1 of
16	OTIB-52, which was effective in 2011, made that
17	change.
18	MR. FARVER: Okay.
19	MR. SIEBERT: The version that was
20	in place when this one was assessed stated to
21	assume it was PA because it was likely not
22	screening at a large site because they were

1	MR. FARVER: Okay. So it's been
2	corrected or changed?
3	MR. SIEBERT: Right.
4	MR. FARVER: Okay.
5	CHAIRMAN KOTELCHUCK: Alright.
6	So it will be closed, should be closed.
7	MEMBER RICHARDSON: Can that be, or
8	will that be noted with the closing?
9	MR. FARVER: I'm writing something
LO	in there about OTIB-52 being revised to
L1	
L2	MR. SIEBERT: Doug, that's in our
L3	response.
L4	MR. FARVER: Oh, good, I'll paste
L5	it.
L6	Okay. 250.2. Incorrect time
L7	period was used for the internal dose
L8	calculations. The employee worked 7.9 months,
L9	not 7 months, which would change the time
20	period. It does result in an increase of about
21	12 rem. NIOSH agrees that the incorrect time
22	was used it's going to result in a DFR

1	CHAIRMAN KOTELCHUCK: Could I ask,
2	for 7.9 months, what about the 250-day minimum
3	that I thought was a requirement for
4	compensation?
5	MR. FARVER: I believe this was
6	just one time period.
7	CHAIRMAN KOTELCHUCK: Right.
8	MR. KATZ: Dave, you're thinking of
9	the SEC requirement, Special Exposure Cohort
10	requirement. Nothing to do with here.
11	CHAIRMAN KOTELCHUCK: Oh, right,
12	yes, correct. Thank you.
13	MR. FARVER: No, this looks like
14	the total time period for this calculation.
15	CHAIRMAN KOTELCHUCK: No, no, Ted
16	is right. I knew that there was a 250-day
17	minimum requirement for SEC qualification, not
18	for individuals.
19	MR. FARVER: Okay.
20	CHAIRMAN KOTELCHUCK: Okay.
21	MR. FARVER: That could impact the
22	PoC.

1	CHAIRMAN KOTELCHUCK: Yeah. So
2	given that it may have an impact on the outcome,
3	shouldn't we leave this open?
4	MR. FARVER: I was thinking the
5	same thing, because we're at 48 percent
6	MR. KATZ: Well, again, this is
7	Ted. You don't need to leave this open. As
8	long as the findings are agreed upon by the
9	Subcommittee, you can close the findings, and
LO	NIOSH does the follow-up as a matter of course.
L1	CHAIRMAN KOTELCHUCK: Well, mostly
L2	we don't think it will have a great impact.
L3	MR. KATZ: No. But whether it has
L4	an impact or not on the case doesn't change
L5	[what] the Subcommittee's done with this, and
L6	NIOSH will do the follow-up as to whether it
L7	needs to redo the dose reconstruction now or at
L8	a later point. But that's independent of what
L9	the Subcommittee does here.
20	CHAIRMAN KOTELCHUCK: I thought
21	we, earlier even today, said, well, let's keep
22	an eye on this and make sure that something is

1	done.
2	MR. KATZ: Right. But I think that
3	was a little different situation from this
4	case. I don't want to go back, but there was
5	some uncertainty about that case, [which] was
6	why you were leaving it open for resolution.
7	CHAIRMAN KOTELCHUCK: Okay.
8	Alright. What do others think on the
9	Subcommittee?
10	MEMBER RICHARDSON: This is David
11	Richardson. I don't disagree with Ted. I
12	agree with Ted. What I'd like is a little bit
13	more explanation of the issue again here and
14	NIOSH agreeing with it.
15	MR. FARVER: The issue?
16	MEMBER RICHARDSON: Yeah.
17	MR. FARVER: The employee
18	employment period was 7.9 months, not 7 months.
19	And so when they did their internal dose
20	calculations, they used the wrong time period.
21	MEMBER RICHARDSON: So when they do
22	the calculation, they're using the NIOSH IMBA

1 or something like that? I believe this has to MR. FARVER: 2 3 do with OTIB-49 adjustments, Super S plutonium. MEMBER RICHARDSON: This is a hand 4 5 calculation or something? Why are theemployment dates -- how is there an issue of 6 7 entering the employment dates in for calculation of an internal dose? That's not 8 I guess I'm asking is this an issue 9 imported? of having to re-enter information and, at this 10 11 point, the dose reconstructor not entering it in properly? Or how did this come about? 12 I believe this is the 13 MR. FARVER: 14 prorated issue where you're multiplying it by a fraction. You're getting a yearly dose, and 15 you only want to apply part of that year or part 16 of that dose. 17 MEMBER RICHARDSON: This is what 18 was described as a calculation that used to be 19 marginal and something added in, and now they 20 had to do fractions? Is this something that's 21

going to happen again, I guess?

22

Or has

	anything changed?
2	MR. FARVER: I'd say it's possible
3	it'd happen again because the prorating is
4	something that the dose reconstructor would
5	enter.
6	MEMBER RICHARDSON: So the
7	employment dates are validated, they come from
8	the Department of Labor as a part of the basis
9	for establishing that the worker has a valid
10	claim. And there's not a way for those dates
11	to be that information, to be incorporated,
12	it has to be taken, re-entered again by the dose
13	reconstructor?
14	MR. FARVER: Well, Scott's
15	probably digging in the files right now to look
16	this up. But I would say that it comes up with
17	a dose, and then the dose reconstructor applies
18	a fraction by calculating what the time period
19	is, and sometimes they calculate it
20	incorrectly.
21	MEMBER RICHARDSON: But they're
22	calculating a fraction off of dates that are

1 in electronic form, I guess is what I'm asking. That I don't know. MR. FARVER: 2 MR. STIVER: This is John Stiver. 3 It might be a good question for Grady or Scott. 4 5 MR. CALHOUN: I'm waiting for 6 I'm sure he's digging, just like --Yeah, I guess what I MR. SIEBERT: 7 can say is, yes, it's electronically available 8 through NOCTS. However, we do not have a tool 9 10 as such that does prorating based on employment No, we don't have that directly. 11 in NOCTS. MEMBER RICHARDSON: The person has 12 13 the ability to look at the dates, or do they exist on this same sheet that they're doing 14 with the internal 15 their work on dose 16 calculation? I mean, are you saying that the information is siloed and they need to flip 17 between an employment-history-siloed database 18 internal-dose-calculation 19 and an siloed database, and they do something which is to 20 create a fraction based on employment dates, 21

and they do that manually moving from one type

## of information to another?

MR. SIEBERT: Yeah, they'll use the information from NOCTS and do the prorating into the tool for, in this case, if I remember correctly, it's using the OTIB-18 tool for the early years of X-10. So, yes, they have to do that manually.

MEMBER RICHARDSON: Is there any information that that tool takes from other databases? Are there kind of identifiers that you have that are populating already as a unique subject ID or anything else?

MR. SIEBERT: I can't speak to that because we haven't looked into specifically having a tool to do that type of thing.

MEMBER RICHARDSON: I'm not talking again about the tool. I'm just trying to, you know, learn and understand the process. I mean, so that tool sort of stands alone and it doesn't have any information about a particular case, and the value that it outputs is not even uniquely identified for a claim?

1	MR. SIEBERT: Well, the
2	information that is entered is based on full
3	annual years of exposure because we don't have
4	smaller time frames than that. Then you take
5	the output and you prorate it to the employment
6	time frame. So, yes, the dose reconstructor
7	does that themselves and does the prorating.
8	CHAIRMAN KOTELCHUCK: Why, with
9	all the detail that we have in so many of the
10	calculations where it's done for people,
11	something so simple as the employment period is
12	not computerized?
13	MR. CALHOUN: It is computerized.
14	This is Grady. It is computerized in NOCTS.
15	However, comma, there's a lot of things you've
16	got to think about. The entire employment
17	history is what we receive. They could be from
18	multiple sites. The person could have
19	multiple cancers that were diagnosed on
20	multiple dates.
21	So you've got to actually look at
22	the employment period to determine how much of

that employment period is assigned to each 1 specific cancer and how much from each site, if 2 3 it's a multiple site case. Now, I guess that could be done, but 4 5 it's not quite as simple as you might think off 6 the top of your head. CHAIRMAN KOTELCHUCK: Part of it is 7 that most -- is it not true that most cancers 8 are singular, that is one type? Or we really 9 10 do have so many -- well, not skin cancer. 11 cancer, there are many primaries. But the 12 other cancers are --13 MR. CALHOUN: I can't give you a 14 percentage, but it's a lot. CHAIRMAN KOTELCHUCK: 15 Okay. MR. CALHOUN: You know, there's a 16 17 lot of cases with multiple cancers. frequently see bladder and prostate cancer 18 together. I don't have a number, but I would 19 say that it's approaching, I don't know, maybe 20 a third, even. I don't know that. I could 21

find that out.

CHAIRMAN KOTELCHUCK: I would just say this: I mean, even that way, suppose there were two types of cancers. Give me the dates worked up until the date of diagnosis, and that can't be calculated automatically? And wouldn't that avoid these simple but significant mistakes? Because this is not the first one we've dealt with. We were dealing with it earlier today. People just make a mistake, you know, in the time period worked, and it affects everything.

MR. SIEBERT: Well, this is Scott.

One thing I want to point out is, although, you're correct, we've run into it in this set a few times, in the overwhelming number of cases we do not do prorating, based on the fact that either we can overestimate, leave the whole year in, or we can underestimate, leave a whole year out, or use IMBA to do the calculations themselves. It's only when we're into this best-estimate territory that we'd be doing the prorating.

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1	CHAIRMAN KOTELCHUCK: Right.
2	MR. SIEBERT: So that's why, you
3	know, I don't think it's been a priority to
4	develop specific tools for that issue.
5	CHAIRMAN KOTELCHUCK: Right. So
6	overestimating and underestimating obviously
7	require different approaches, and people have
8	to use their wits and understanding. So, okay,
9	that satisfies me as to why that isn't just
10	automatically done.
11	MR. FARVER: So do we want to keep
12	this case open, or this finding open? Or close
13	it? Because we're not sure exactly how much
14	it's going to impact
15	CHAIRMAN KOTELCHUCK: I think the
16	argument was that we should close it. That's
17	what I heard. And that it will be taken care
18	of, that we don't need to come back to it.
19	MR. FARVER: Okay.
20	CHAIRMAN KOTELCHUCK: Others
21	agree?
22	MEMBER MUNN: That would be my

1	recommendation, yes.
2	CHAIRMAN KOTELCHUCK: Okay.
3	MEMBER CLAWSON: That's fine with
4	me. This is Brad.
5	CHAIRMAN KOTELCHUCK: Okay.
6	Dave?
7	(No audible response.)
8	CHAIRMAN KOTELCHUCK: 268.1.
9	MR. FARVER: 268.1.
10	CHAIRMAN KOTELCHUCK: By the way,
11	folks, I think we only have a few more cases to
12	deal with in Oak Ridge, and I think there's a
13	chance we can finish this up.
14	MR. FARVER: I don't know. You're
15	awful hopeful.
16	MEMBER MUNN: Yeah.
17	CHAIRMAN KOTELCHUCK: Well, I
18	can't scroll through to actually see. I just
19	took some notes yesterday before our meeting.
20	But, okay, let's go ahead. Forget the comment.
21	It's irrelevant.
22	MEMBER MUNN: Everybody loves an

1	optimist.
2	CHAIRMAN KOTELCHUCK: Well, we got
3	to yeah, we've got to be optimistic.
4	MEMBER MUNN: True.
5	CHAIRMAN KOTELCHUCK: Especially
6	late in the day in the middle of a two-day
7	session.
8	MEMBER MUNN: Yes.
9	CHAIRMAN KOTELCHUCK: 268.1. You
10	go ahead. Sorry, Doug.
11	MR. SIEBERT: This is Scott. I'm
12	sorry. This first one, 268.1, is a very
13	technical issue with the scaling factors at
14	Y-12. And just my opinion, it may be wise to
15	not start this one today.
16	MR. FARVER: I agree, Scott,
17	because that finding is misleading when you go
18	back and look at the original report. It's
19	very complicated.
20	MEMBER CLAWSON: Okay. There went
21	the positive attitude. Now let's get to the
22	pessimists.

1	(Laughter.)
2	MEMBER MUNN: Thanks for that
3	recommendation.
4	CHAIRMAN KOTELCHUCK: 268.1 open.
5	Alright. Optimism has to give way to
6	experience. Let's leave that open. 268.2
7	then.
8	MR. FARVER: 268.2. Incomplete
9	accounting of all recorded doses. The records
10	show that the employee was monitored for a short
11	period in '87 with a resulting dose of 49
12	millirem. They did not assign a dose, but they
13	assigned ambient here. They should have
14	assigned the 49 millirem.
15	CHAIRMAN KOTELCHUCK: Yes.
16	MR. FARVER: It looks like it was a
17	data entry error. In other words, it just
18	didn't get entered into the file.
19	CHAIRMAN KOTELCHUCK: And NIOSH
20	believes that there will be little impact.
21	MR. FARVER: Probably.
22	CHAIRMAN KOTELCHUCK: And we can be

1	confident that that will be checked so that we
2	can close it?
3	MR. FARVER: Well, you've got a PoC
4	of 39 percent. I don't think 50 millirem is
5	going to
6	CHAIRMAN KOTELCHUCK: Oh, you're
7	quite right. Okay. Well, I don't know what
8	the PoC is.
9	MR. FARVER: You're right.
10	CHAIRMAN KOTELCHUCK: Okay.
11	Well, if it's 39, then it will not have an impact
12	[to flip the PoC].
13	MR. FARVER: I don't believe so.
14	CHAIRMAN KOTELCHUCK: Okay. In
15	which case, we should close it.
16	MR. FARVER: It is just another QA
17	concern.
18	CHAIRMAN KOTELCHUCK: That's
19	right.
20	MEMBER MUNN: Yes, agreed.
21	CHAIRMAN KOTELCHUCK: Okay.
22	Let's go on to 268.3, which is on our screen.

1	MEMBER MUNN: Not on mine. I've
2	lost the whole thing.
3	CHAIRMAN KOTELCHUCK: Oh, okay.
4	MEMBER MUNN: I'm gone.
5	CHAIRMAN KOTELCHUCK: Oh my.
6	Well, it will come back. 268.3 is on the
7	screens of those of us who have screens. Yeah,
8	my computer keeps going out all the time. I
9	have to get my password to get back in, but so
10	far I've been able to.
11	268.3, Doug.
12	MR. FARVER: 268.3. SC&A
13	questions the solubility type used for the RU,
14	recycled uranium, components. And once again,
15	this is for K-25, Paducah, Y-12, X-10. Pretty
16	much it just wasn't clear where they got their
17	values from. With all the types, it was a
18	little confusing. So they cleared that up in
19	their response. It came from OTIB-60, Section
20	57, which covers the recycled uranium. It then
21	tells which types to be assigned.

1	this an observation?
2	MR. FARVER: I don't know.
3	CHAIRMAN KOTELCHUCK: This doesn't
4	seem to be a finding. You had a question,
5	correctly
6	MR. FARVER: It's a finding because
7	we just didn't know where they came from.
8	CHAIRMAN KOTELCHUCK: And now you
9	know, and it was right all along.
10	MR. FARVER: It's not that it was
11	incorrect. It was there was not enough
12	information in there to tell where these
13	assumptions came from.
14	MEMBER MUNN: Lack of information.
15	CHAIRMAN KOTELCHUCK: And that's
16	what B is?
17	MR. FARVER: I'm not sure.
18	CHAIRMAN KOTELCHUCK: Okay.
19	Well, clearly, it can be closed.
20	MR. FARVER: Okay. Observation 1,
21	there was no workbook to show the derivation of
22	the doses assigned by NIOSH in the files, they

1	could not reconcile the doses assigned in the
2	IREP input. But because they were
3	claimant-favorable, this was listed as an
4	observation. And this is for x-ray doses.
5	CHAIRMAN KOTELCHUCK: You might
6	scroll up just a little bit now to see the end
7	of the write-up. Thanks.
8	MR. FARVER: In other words, when
9	we went through our calculations, we calculated
10	what we thought it should be compared to what
11	the NIOSH calculations were. It seemed
12	reasonable, very close. But there was no
13	workbook in there actually showing the
14	calculations.
15	CHAIRMAN KOTELCHUCK: Right.
16	MR. FARVER: Okay. Observation 2,
17	the employee had whole body counts for
18	different years that were labeled as
19	insignificant for all but cesium in '64.
20	CHAIRMAN KOTELCHUCK: Okay.
21	Observation 2 is not on the screen yet. There
22	we go.

1	MR. FARVER: But during this time
2	period, a urinalysis was used to assign the
3	internal doses. Therefore, the whole body
4	count information was not needed to assign dose
5	in this case. We thought it would be best if
6	they would just include some kind of statement
7	in the report acknowledging that they had whole
8	body counts. NIOSH agreed that it would have
9	been nice to have some more information in
10	there. Okay.
11	CHAIRMAN KOTELCHUCK: Alright.
12	MR. FARVER: Do we want to move
13	forward?
14	CHAIRMAN KOTELCHUCK: Yes. 269.1
15	is open.
16	MR. FARVER: Okay.
17	CHAIRMAN KOTELCHUCK: No, 268.1,
18	I'm sorry, is the one that's open. 269 would
19	be the next one. Sure, let's keep going.
20	Folks, it's 4:30. What is the do we have
21	just a few more or
22	MR. FARVER: I really don't think

1	we're going to finish this today.
2	MR. KATZ: This is Ted. People are
3	sounding like they're out of gas. I don't know
4	if that's true but
5	CHAIRMAN KOTELCHUCK: Yeah. Then
6	what do we have left, I can't see it here, for
7	Oak Ridge? We have 269.
8	MR. SIEBERT: There are three more
9	cases.
10	CHAIRMAN KOTELCHUCK: Okay.
11	294.1, 324.1. Okay. Look, then, given that
12	we have two days in a row, maybe we should just
13	call it quits now at 4:30. I'm more than open
14	to doing that and just resume again in the
15	morning with 268.1.
16	MR. KATZ: That makes sense to me.
17	And Mark Griffon will be joining us tomorrow.
18	CHAIRMAN KOTELCHUCK: Good.
19	Okay, fine. And hopefully Dr. Poston also.
20	MR. KATZ: And hopefully Dr.
21	Poston, as well. Right.
22	CHAIRMAN KOTELCHUCK: Folks, then

1	I think we've agreed. Let me thank you all for
2	today. A long day and we have another long day
3	tomorrow. So have a good evening.
4	MR. KATZ: Yes. Thank you,
5	everybody.
6	(Whereupon, the foregoing matter
7	was concluded at 4:32 p.m.)
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