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in, so it's good from that. And I think, in all fairness to NIOSH, they need to review the report and then I think there are some steps that can be taken, you know, relatively soon to at least think of ways that the smoking issue can be addressed. And Russ, if you want to elaborate, you're...

MR. HENSHAW: I just want to say -- is this
on? I can't tell from -- yeah. We have something
from NCI. We haven't really had a -- we just got
it -- well, Tuesday, I believe -- Monday or
Tuesday. We haven't had a chance to really look
at it very carefully, so there's a possibility,
maybe a probability, we'll need to go back and get
some additional data to understand the few pages
of information we have so far.

DR. MELIUS: Epidemiologists always have an odd view of time and so forth -- trouble predicting when something will get done or complete. And it's never complete, always got to have more analysis.

The final issue really is related to the first issue, which is the issue of how to address other occupational exposures that might take place, particularly within the DOE sites. And I think that's really part and parcel of the first

issue, the occupational cohorts that are being looked at. And so when we get an update from HERB, I think we'll be able to ask more questions about that.

The final thing I wanted to just mention is that the update to BEIR is underway and I don't think we're expecting anything very soon on that. But that will clearly have a -- could have a large impact on -- terms of possible changes that might need to be made to IREP or something from the analysis and reporting that's underway there, that's at least a year away, as I recall, maybe even longer before we see that. You remember the

MR. ELLIOTT: My understanding from one of the members of the BEIR committee was that the report was due to surface in public last November, and we haven't seen that yet. So I had a call in to Eula Bingham to find out where it's at and what the holdup is, and I haven't got a comment back. But I don't believe it's a year away. I think it's closer than -- than maybe that, that we think -- should be here soon, I hope.

DR. ZIEMER: Now I believe that report is dependent upon official issuance by RERF of the

new risk coefficients. Is that correct?

DR. MELIUS: I believe so, yeah. That's my understanding.

DR. ZIEMER: I have heard, unofficially, that those risk coefficients are not likely to change very much. I don't know if any others have heard rumors, and certainly the record shouldn't show that to be definitive in any way, but my understanding is that the changes in the dosimetry -- which goes back to the Japanese dosimetry -- have been, for the most part, rather small changes and hence the risk coefficients, though they will change, will not change by great amounts. But it still remains to be seen what the impact will be on -- eventually on IREP and we want to certainly be tracking that.

MR. ELLIOTT: I certainly agree. That's similar to what I've heard. We were also anxious to see what the report would say, though, about occupational studies and their effect or non-effect on risk --

DR. ZIEMER: Right.

MR. ELLIOTT: -- estimates, so I think that's our focus on this report. That's where we want to see it come in.

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DR. ZIEMER: That may be of greater importance, actually, than the coefficients, which may not change very much.

Could I also ask, on the smoking issue, once you've digested that information, is there a plan to report -- maybe at the next meeting -- what those findings were? Or what -- what do we expect to get from NCI on the smoking issue?

MR. ELLIOTT: What we -- what we're talking about in receipt from NCI is basically the Pierce analysis data that was done to support their modifications on smoking and lung cancer. And what Russ alluded to was that we've got four or five pages of really what looks to us like a SAS* printout with no data dictionary and no explanation and no interpretation, and so that's what we're after right now. It would be our intent that we analyze that bit of information and come back to the Board with a proposal on the impact on the NIOSH-IREP cancer risk models for lung cancer and what we should do in that regard, what changes or non-changes should be made. And so we would present that to the Board. Of course we would have that peer-reviewed and vetted and then brought to the -- those comments and the

resolution that we provide to those comments 1 2 brought to the Board, as well. 3 DR. MELIUS: And that's my report. 4 DR. ZIEMER: Okay. Thank you, Jim. Let's see if there are additional questions relating to 5 6 the report of the research group. 7 (No responses) It appears that there are not, and there's no 8 9 specific recommendation beyond these general things that we're looking forward to. 10 11 DR. MELIUS: Correct. Yeah, it's -- I think 12 it's more of an information update at this point 13 in time. BOARD DISCUSSION/WORKING SESSION 14 15 DR. ZIEMER: Thank you very much. If you 16 would look at your agenda and make sure that you 17 have the correct version of the agenda -- which I didn't have. But the correct version of the 18 agenda now for our next item -- except for (off 19 20 microphone) the break, which (Inaudible) since we're a little ahead of schedule -- there's a 21 22 Board working session for dose reconstruction 23 review process --24 THE COURT REPORTER: His mike's gone. 25 DR. ZIEMER: -- is what you should have.

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Does everyone have that version of the agenda?

And the reason I call that to your attention is because the earlier version showed the item as being Sanford Cohen & Associates as the next item, where in fact that has been --

THE COURT REPORTER: It's in and out.

DR. ZIEMER: -- that has been scheduled for tomorrow at 9:00, Board discussion/working session on Stanford Cohen & Associates with respect to the Board support for dose reconstructions. So our focus at this moment will be on the dose reconstruction review process. And we had set aside time on this I think from our last meeting to do any follow-up on that item, and I'm trying to recall, Mark -- and I'll ask if you can help me out on this -- where did we stand as far as the working group's recommendations were concerned after the end of the last session? I'll put you on the spot here a little bit.

MR. GRIFFON: Yeah, I know. I thought this was on the schedule for tomorrow, actually. You know, I'm not sure where we left off. We had a draft procedure for our review process, but beyond that, I don't know where the working group left off or if you...

MR. ELLIOTT: I, too, am at a little bit of a loss here. I think -- maybe we could recap to -- to the point of -- as to where we're at right now. We -- you -- we haven't announced yet, but we have -- you have awarded two of your tasks, and that's what you will be able to talk to Sanford Cohen & Associates tomorrow about. Tasks two and four have been awarded and they can start work under -- under those tasks. So you might want to think about those two tasks and whatever questions of clarification you have for your contractor or anticipating what questions they might have of you.

The other two tasks, one and three, are -are not awarded. Those are still in the
negotiation process. Those are what you're going
to discuss in closed session tomorrow, so you're you're limited in what you can discuss in open
session about those. You could discuss -- you
know, we've still I think been wrestling with how
you're going to come up with your selection of
cases in a stratified -- representative or
stratified random sample. What are the variables
-- we would ask you what are the variables you
want to target for your selection of those cases.

We have bantered around this idea of a subcommittee or not subcommittee. I think you've come to grips with that. You want the whole Board to be involved, but you might still think about -- you know, as you proceed here, do you really -- is that the way you want to go. You know, there's some work here to be done as far as identifying cases for review when that task three is awarded, and assigning who's to review those cases and what that process really looks like.

So I mean I'm just trying to throw out ideas for topics for discussion here for this afternoon and perhaps tomorrow. And I'm certainly not -- want to lead you in one way or the other here, but these are things that kind of we have questions in our mind about how -- how do -- how do we go about doing these reviews. We're still -- we're still wrestling with what your approach and your process is going to be and how we will attend to making sure that we protect the privacy of individual claimants, how -- what your report is going to look like at the end of your review, you know. We're still awaiting to hear your thoughts on that, so those are just my thoughts off the top of my head.

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DR. ZIEMER: Thank you, Larry. And tomorrow during the official session with SC&A -- that is, during the morning session -- we will have a chance for them to ask questions and for us to ask questions pertaining specifically to task two and four, which have been awarded. That is -- and John Monroe (sic) and Joe Fitzgerald I understand will both be here from SC&A and there will be an opportunity for them to seek clarification on those tasks and for us to ask them questions and discuss those in more detail.

Okay. Now Jim and then Wanda.

DR. MELIUS: Well, one question they might ask us tomorrow is what site profiles do they want us to review, so I think, you know, sort of meaty issue is going to be how do we select those to get them underway -- get those reviews underway, but -- and I was thinking that in a more general sense the way of approaching this is to think -- much as some of the examples Larry just used is to think about what are the different activities that are involved here. How do we as a Board want to handle them. How do we want to select the site profiles, then the individual cases. We've still got work to do on that. How are we going to

interact with the contractor. Is that going to be done -- you know, the contractor has questions, who do they call, how do we get clarification on that. There's some issues that I think we have to be -- be careful both from the contracting point of view, but also in terms of the credibility of the process and making sure that's taken care of. And I think we just need to work through those and decide what's the best way to do that and are we going to need a subcommittee to do that, how much guidance do we give the subcommittee, do we do it as a committee -- the whole committee for -- for each of those. And then try to categorize them and come up with a timetable for dealing with them.

DR. ZIEMER: Okay. Wanda?

MS. MUNN: I hate to admit this, but I no longer remember what tasks two and four were. I remember what one and three were because -- for obvious reasons, but not having brought previous notes with me, I'm at a loss. Will someone please refresh my memory?

MR. ELLIOTT: Well, I'll try to do that, and
I'm certain that Mark will correct me in any way
that I might err here. Task two is to review site

profiles, and task four is to develop a database, a data management system for you all. Remember, task four was to design that, develop that, put that into place. And I think that involves, you know, tracking the cases that are assigned, when they were assigned, who's working on them, what the findings were, perhaps even -- you know, database management aspect of -- of how many site profiles have been examined within, you know, task two, as well as under task three where we -- you're looking at individual completed dose reconstructions. So you know, I think there's a lot to be talked about under task four. It may seem apparently obvious what has to be done, but I think you need to probably talk through that.

DR. ZIEMER: Task -- task two more specifically was --

UNIDENTIFIED: (Off microphone) Paul, .
(Inaudible) the mike.

DR. ZIEMER: Sorry. Task two was to prepare a site profile review procedure, not to do site profile reviews.

MR. GRIFFON: The task was to develop the methodology and also to do the reviews of I think ten to 12 DOE sites and two to four AWEs, so it

involved both.

DR. ZIEMER: Oh, yeah, you're right. You're right. The first step was the procedures, and then ten to 12 DOE sites and two to four AWES. So it may -- it may be that the actual determination of selecting the sites, we can start to be talking about that, but we have to have a -- we also need to know what the procedure is that the contractor will use, and we've asked them to do that as a first step in the process.

wondered if we have a copy of the procedure for processing individual dose reconstruction reviews, the one that we voted on and approved. I have it on the computer here, but I don't have a hard copy. The reason I say that is a lot of the bullets right at the front end of this procedure -- maybe we didn't flesh out everything, but we at least identified several of these issues that Larry and Jim have brought up that maybe we just need to run through again and clarify how it's really going to work now that we know a little more of what the contractor's proposed, et cetera.

MR. ELLIOTT: I don't know if Cori brought that particular document along for reference, but

2 get it off your laptop. 3 We could put it up on the screen. Let me find Cori and we'll see if ... 4 (Pause) 5 6 DR. MELIUS: While we're asking for what 7 information's available, that -- I don't know if 8 Martha or somebody has with them the award for 9 tasks two and four that would lay out the 10 timetable we -- 'cause -- gave the contractor 11 because I think -- we're going to have to know 12 that timetable on those tasks in order to sort of 13 figure out meeting schedules and how -- when 14 they're going to get feedback and so forth, so ... 15 MS. DIMUZIO: I don't -- I have them upstairs 16 in the room, so I'll go upstairs and get a copy of 17 that and I can bring it down. 18 DR. MELIUS: You actually make copies for the 19 Board? 20 MS. DIMUZIO: Yeah. Yeah. 21 DR. MELIUS: Would it be best to take a short 22 break or something, get some of this stuff copied? DR. ZIEMER: Yeah, let's -- let's take ten. 23 24 Uh-huh, that's fine. 25 (Whereupon, a recess was taken.)

we can certainly I think get it printed if we can

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DR. ZIEMER: I have a technical instruction for the Board and for myself. We've been instructed that when you're holding down the push button on your mike, be sure to hold it in the center or push it in the center and hold that steadily. Don't rock to the right or to the left 'cause it cuts the mike in and out.

Now Cori is distributing the document that came from the working group on procedure for processing individual dose reconstruction reviews. Task two, which we had been talking about, on site profiles -- task two has as a first item, prepare a site profile review procedure, and that's a deliverable one month after the authorization to proceed. So we're -- we're actually two weeks into that, aren't we, John?

DR. MAURO: One day.

DR. ZIEMER: Oh, you didn't get your authorization as fast as I thought you --

DR. MAURO: Just got the authorization yesterday.

DR. ZIEMER: Okay. I was thinking you'd be ready to report on the -- just kidding.

Okay, he's -- but the clock is ticking on that one.

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The issue of selection -- well, there will be an issue we want to talk about with regard to that. That procedure will be ready in one month. Then we have the issue of who then looks and reviews and approves that procedure and how the Board wishes to do that. Then the selection of the sites to be reviewed, and it may be that the Board would like to identify some criteria. I mean we have a number of sites -- we saw the matrix earlier today -- that are close to being ready for review. Some are already completed. But given that list, even after it's all completed, how do we decide which ones to review. And you might want to identify some criteria. For example, one criteria might be a site that has generated a large number of dose reconstruction cases. Or we might say let's look at the top five sites as a kickoff, or something like that, in terms of cases. So think about criteria of that sort that we could use so that selection of the site is not just based on gut feeling -- I like one site better than another -- but some sort of objective criteria on which to make those decisions.

Now let's open the floor -- Jim, your flag is