have been used or have been designed for use in doing dose reconstruction; another group -- Joe's group doing site profile review while I'm doing my -- or our team is doing a basic review or an advanced review. If they're moving together in lock-step with continuous communication, the efficiencies will be incredible. If they're not, we're going to lose a lot of efficiency and it's going to have cost and schedule implications. So that's an observation.

DR. ZIEMER: Let me interject that I think it

DR. ZIEMER: Let me interject that I think it certainly was the Board's view and the working groups view that these four tasks are in a sense integrated in the fashion that you talk about. At the same time, recognize that in the sampling process I don't think a priori one could guarantee that a given dose reconstruction would -- that's being reviewed would be from a site that has been selected for site profile review, so --

DR. MAURO: I understand that, but -- it's a complex problem --

DR. ZIEMER: Right.

DR. MAURO: -- but we'll manage it, but these are some thoughts.

DR. ZIEMER: Yeah, but let's -- there's

another comment. Jim?

DR. MELIUS: Yeah, just to follow up on that,
I think that, given the way that NIOSH is doing
the individual dose reconstructions, they do -- as
I understand it, they do a site profile, then they
do a number of individual dose reconstructions.
So just on a random basis, it's likely they'll
overlap.

I think as we charge you with doing dose reconstructions and develop a way of making that selection, it is possible in the future we may want to focus some of the individual dose reconstructions away from facilities that had site profile or things. But I think in that case we should inform you ahead of time as you're, you know, responding as to how those cases will be drawn, or at least some more specific information on that. Again, that's one of the reasons that --some of the changes in the approaches we've made on these tasks.

DR. MAURO: Therein mind our budget, our work hour allocation per case, presumed that they would be working as a couple. If they're decoupled, we do run the risk of some inefficiencies. We actually costed (sic) out the work hour allocation

assuming optimum efficiency, okay? So bear -- I'm already being a project manager, recognize that we -- there are some, you know, loop -- places where we could run at these kinds of problems.

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Another observation -- I have two more observations, then I'm going to turn it over to Joe. Okay?

When I reviewed the Savannah River site profile, I presumed that the -- all of the site profiles will have the same fundamental organization. Let me just reiterate it to you. One is that you first look at the medical exporing this case, after the introduction there's the medical exposure records, review that carefully. That's, in my opinion, fairly straightforward. Once you understand the time and the type of equipment that was used, the protocols are pretty clear, in my mind, as a health physicist. And we have the staff -- medical health physicist -- we're okay. We're okay.

Environ-- now here -- the second one is the occu-- environmental occupational exposures. That is releases that occur from a facility that may expose some of the construction workers that we heard from yesterday. I noticed that what -- what

was done -- well -- with regard to that issue is
to draw upon the work that was done by RAC, Risk
Assessment Corporation. That is, they did the
reconstruction of the source terms, airborne
emissions from the facility for the purpose of
doing off-site dose calculations, dose
reconstruction. And certainly that very same
source term information is of value for evaluating
on-site by using appropriate meteorological
models.

What I guess I was expecting was that these documents would go down -- go to source -- original source documents. That is -- in -- in effect, by using -- and this is by no means a criticism, but in effect you're using a tertiary level document. That is, when you look at records -- I've been involved in a lot of off-site dose reconstruction work, and when you go into the literature you find a hierarchy of documents. There are very high level documents that represent summary level information. And then there are intermediate level doc-- then you get right down to the -- the strip charts. Okay? You get down to the nuts and bolts. My sense is, and here's where I'd like to see what your reaction is, we're

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going to use our judgment of when do we go down into the bowels of the problem; where -- when do we think that -- I'm not just going to trust some tertiary document as being a correct and complete representation. I'm going to go down -- because I've done this before, and I've found lots of surprises. So our plan is when we think it's important -- and here's where things get interesting. When we think it's important -- for example, let's say we're talking about the dose to a construction worker from an airborne emission from a particular facility at a particular time, inhalation exposure to airborne plutonium or cesium or noble gas. When we feel as if that particular scenario might be an important contributor to dose, we're going to dip in from working up here to working down there, and keep you informed. How much of that we're going to have to do, we don't know. So here we have another cost and schedule issue. It's a living process.

Now -- so we're -- we're going to -- we're going to keep you apprised of that, so we're not simply going to go back and take a look and say oh, okay, yeah, they -- they -- they used the RAC

work correctly. Here's the RAC numbers, here's the source terms, the times, yep. So we're going to check that. That's -- that's standard quality control. But then there is the more probing analysis, do we believe that source term. So that's our plan. I'm hoping that you agree with that 'cause that's the only way to do this.

Finally --

DR. ZIEMER: Let's again allow a moment for comments. I think Jim has one and I have one here. Oh, you don't.

Well, what you've described for us is in fact an audit procedure.

DR. MAURO: Yeah.

DR. ZIEMER: And it's not something we necessarily have to approve today. I think your plan will include something along the lines of what you just described to us. And in an audit procedure, a certain amount of that probing -- and then you see what your results are and report those back.

DR. MAURO: Yes.

DR. ZIEMER: You know, we probed down, we pulled the string here, here and here, and in all cases things made -- were fine or in all cases it

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didn't make sense, or some distribution in between there. And based on that, then the Board can say well, there's some issue here. And certainly even that kind of audit procedure doesn't have to be 100 percent audit. You selectively, based on judgments and so on, start pulling those strings where -- where it's appropriate.

DR. MAURO: But you --

DR. ZIEMER: And I assume your plan will describe to us what you --

DR. MAURO: Yes, but you see how this is an open-ended process.

DR. ZIEMER: Yes, yes.

DR. MAURO: And we'll keep you apprised. And when we think we're going to run into cost and schedule issues because of this 'cause we take -- we go where the information takes us, and so we're -- we're very vulnerable in terms of well, you know -- and we'll give you our reasons why we're going where we're going and -- and I -- but I guess in a way we're not going to be seeking approval if -- at any point -- we'll keep you apprised, and if you feel that what we plan to do, you're -- for some reason there are problems with it, then I think certainly intervene, say no,

don't do that, we don't -- regroup and give you further direction. But right now my plan is to keep you apprised, but to keep the train on the tracks and keep it going.

Another observation having to do -- well, two more and I'll be done. When I read chapters in the Savannah River report dealing with occupational exposure, internal and external, I was expecting to see databases of records, of either -- bioassay data, records -- the -- database upon database upon databases of air -- radiation area monitors, continuous air monitor data. In other words, just enormous -- an ocean of data that represents location and time when the material was collected.

What is there is something a little different. It really is almost a -- a guide to the dose reconstructor to help him fill in gaps, understand what the minimum detectable levels are, understand what mixes to assume, what chemical forms to assume. In other words, it's almost as if it's a helper, as opposed to a database. Okay? I think that's good that it -- you know, I guess my reaction was that's good that-- but as an auditor that's trying to independently evaluate, I

sure would like that database. Is there anything going on to compile that kind of data? I mean we're talking about the tons of -- of -- the big spreadsheets of Excel databases which show, as a function of time and location, individual measurements -- whether it's bioassay or airborne radionuclide particulate or it's radiation area monitors that are taken by location as a function of time and put into a database. That, to me, is an important information. Now --

DR. ZIEMER: I don't know that we'll answer that specifically today, but that -- as you get underway now, you will have an opportunity in fact to see a lot of underlying data that's beyond what's in the immediate report.

DR. MAURO: Okay.

DR. ZIEMER: And that's one of the things we'll want you to become familiar with is what all the supporting databases are for these things.

DR. MAURO: Okay, so --

DR. ZIEMER: And what's there and what isn't there.

DR. MAURO: Okay.

DR. ZIEMER: And you know, if -- if you, as our auditors, have some judgments on adequacy or

lack thereof of some -- at some site, that could be part of a report.

DR. MAURO: One of the --

DR. MELIUS: Can I just --

DR. ZIEMER: One other comment here.

DR. MELIUS: Can I just follow up on that, because I thought Jim Neton answered this question partially or in his presentation yesterday and so forth, and I don't know where -- I thought he had referred to the fact that they do have this compilation of information, dose -- exposure information or whatever. It's not necessarily referenced in the document, and --

DR. ZIEMER: That's why I say I think once you're into beyond what's on the web site, once the contractor has access to all those records, then you can perhaps make a better judgment on what additional things you think are needed or maybe you'll feel it's adequate and so forth.

DR. MELIUS: But just to follow up on that, and maybe it's -- maybe you've thought of it already, but it -- for NIOSH, in producing these documents, it seems to be a common question, a common concern that people have is why isn't this information look-- referenced, and it may very

well have been looked at and in some sense utilized, it's just not printed there as a reference. And maybe that's something you ought to consider adding to those documents as a way of just, you know, showing what kind of a guidance, you know, this is and what other information's available. I'm not familiar with the details to know how practical that is, but it -- you know, it might be helpful. It might be -- for other people as they're looking at these documents, also.

DR. MAURO: Last point has to do with the other deliverable, the tracking system. I was speaking to Don Loomis, who is the database manager task leader on that, and re-- he knew -- told him I was coming to this meeting today, and there -- in his -- his view is that there are no boundaries on how many fields we can handle, any kind of queries you want. But what would be helpful is the -- is when we build the relational database that we put in all of the fields and all of the types of reports built into the system.

Now -- that was part -- it was -- that's -- the other deliverable a month from now is that program. So we already have a list. We understand from your request, your torp, and from

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our proposal what we do plan to put in there. But I plan to put a lot more in there, and let me explain what I mean.

For example, all of our project management data where tak -- we took each task, one, two, three, four, and we're breaking them down into subtasks and sub-subtasks. For example, on task two, the site profile work, we expect to have a number of site profiles. Each site profile's going to have its own point number for tracking costs. Each case that comes in on task one is going to have its own point number for tracking costs so that as a project manager I understand where the money is going and why. If there are -in a similar way -- I guess what I'm -- I'm asking you is that anything that you want to do, I don't care what it is, related to queries and sorting of data and reports that you'd like to be able to elicit from this database, we can handle. But the sooner you give it to us, the better. We could revise it later, but it's a little more difficult, I'm told, to do it after the fact than before.

And I guess that concludes my I guess initial reaction to things. If there are any questions -- DR. ZIEMER: Yeah, further questions? Joe,

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do you have additional comments or items you want to add to ...

MR. FITZGERALD: Thank you, John. Well, it's good to finally be here after some years. I think John covered the highlights, but one thing I want to just mention -- I'm very comfortable with the task, very comfortable with the touchpoints in the task, but I want to emphasize that, you know, to me, this is really doing a vertical sampling, boring down and asking probably questions that if you were doing a horizontal -- and getting the necessary as opposed to maybe totally sufficient data for dose reconstruction, you might not get to or might judge that you might not need. And when you do the vertical and you push down and you actually get beyond what's on the shelf, what the paper says, then you get into situations where you will be asking for data, you'll be probably wanting to interview people that haven't been touched by the process to date. And from some limited experience over 20 years, that's going to enjoin probably some challenges that we will bring back to you in the way of access, the way of perhaps getting information. I know that's been some of the experience to date. But I think doing

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this kind of review is probably going to engender more of those kinds of challenges in terms of getting to the right kind of information and digging into areas that haven't been dug into. I've done it my entire career, so I know what's involved in doing that, and persistence will pay. But I just want to sort of lay that observation -it's not a question for the Board, but just an awareness of what -- what's involved when one truly does a vertical sampling to answer the hard question of adequacy and completeness. that's the -- sort of the tail end of necessary and sufficient. We're answering a sufficiency question. And so that's -- that's something that I think as we go into this it'll become clearer what -- where we might need your role perhaps in some cases with the Department of Energy, where we might need some clarification as to, you know, how deep does the vertical go in some cases. pretty comfortable definitely with the scope and the tenets and certainly we'll be able to articulate a plan that will reflect what we proposed in the beginning, and also what it's going to take to answer that question. And I certainly do understand the challenges that NIOSH

and ORAU have undergone in terms of doing this -the necessary part, but this is going to be a -certainly a somewhat different process. And you
know, the question of access to information,
access to people, workers, all that, I think will
be certainly decidedly answered by our first
forays into this. So that's -- that's really my
only observation.

I think John covered some of the more tactical questions, but sort of on the 30,000 foot level, that's -- that's going to be, I think, the biggest challenge and the question of how we can deliver that for you and intend to deliver that for you. So thank you very much.

DR. ZIEMER: Thank you, Joe.

MR. FITZGERALD: Any questions?

DR. ZIEMER: Let's see if there's any questions for Joe. Jim?

DR. MELIUS: I have a -- I'm not sure who it's for, but in terms -- in -- I'm not familiar with the details of what you've been awarded, or at least -- or I don't recall them, but the -- in terms of making the assignments in the site profiles, are there -- and we have to -- going to try to, I think in our later discussions, sort of

narrow down where to get started. In your planning and sort of to do that efficiently, I guess sort of how many does it make sense to be assigned initially or is it -- make sense to say here's the -- whatever it is, ten, 12, whatever; go get started and, you know, they'll be done under this task order under the -- a year, or is it, you know, let's wait three -- you know, do five now, five in three months, what -- I guess I'm trying to get some sense of what your expectations are at this point.

MR. FITZGERALD: Well, you know, I think -we haven't chatted about the specifics of this,
but certainly my expectations, we would certainly
want to know what the so-called menu would look
like for the year. And I think there's some merit
-- and again, this is the Board's purview and
decision, but some merits perhaps in ramping into
it with perhaps somewhat less complex sites
because, again, we're establishing on the ground
the procedures that we're establishing on paper,
and it certainly would perhaps facilitate things.

Nonetheless, the people that we intend to put into these reviews are not coming into it as neophytes. They have the operational experience