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UNITED STATES OF AMERICA
NUCLEAR REGULATORY COMMISSION

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ADVISORY COMMITTEE ON NUCLEAR WASTE

133RD MEETING

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WEDNESDAY

MARCH 20, 2002

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The meeting commenced at 1:00 p.m. in Conference
Room 2B3, Two White Flint North, Rockville, Maryland,
George M. Hornberger, ACNW Chairman, presiding.

PRESENT:

GEORGE M. HORNBERGER	ACNW Chairman
B. JOHN GARRICK	ACNW Member
MILTON N. LEVENSON	ACNW Member
RAYMOND G. WYMER	ACNW Member

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1 STAFF PRESENT:

2	JOHN T. LARKINS	Exec. Dir.-ACRS/ACNW
3	SHER BADAHUR	Assoc. Dir.-ACRS/ACNW
4	HOWARD J. LARSON	Spec. Asst.-ACRS/ACNW
5	LYNN DEERING	ACNW Staff
6	LATIF HAMDAN	ACNW Staff
7	MICHAEL LEE	ACNW Staff
8	RICHARD K. MAJOR	ACNW Staff

9

10 ALSO PRESENT:

11	SITAKANTA MOHANTY
12	RICHARD CODELL

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I-N-D-E-X

AGENDAPAGE

High-Level Waste Performance Assessment

Sensitivity Studies

Sitakanta Mohanty 120

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

(1:07 p.m.)

CHAIRMAN HORNBERGER: The meeting will come to order, the afternoon session here of the 133rd meeting of the ACNW. I have a note for the Committee. At three o'clock, we are to go over to the neighboring building to get new badges. And so we have an appointment at three o'clock. That shouldn't be a problem because we have a 2:45 to three o'clock break schedule and I don't think that I will steal so much of the time that we've given over to Dick and to Sitakanta to do a presentation.

Dick was originally scheduled to talk to us about the sensitivity studies for the waste package, and Howard tells me he's going to talk about anticipatory research instead.

(Laughter.)

Although I may be corrected and it may in fact revert back to the sensitivity studies. Sitakanta, are you going to go first or is Dick?

MR. MOHANTY: I'm going first. Good afternoon, ladies and gentlemen. My name is Sitakanta Mohanti. I will be -- myself and Dr. Richard Codell will make this presentation. I will go over the first part of the presentation, and Dr. Codell will make the

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1 presentation on the second half of this talk.

2 The title of this presentation is,
3 "Sensitivity and Uncertainty in the NRC Total System
4 Performance Assessment of TPA 4.1 Code," I should add
5 results. Okay. Here is an outline of this
6 presentation. First, we will briefly address the
7 purpose of this analysis of uncertainty and
8 sensitivity. Then we will present an overview of the
9 Total System Performance Assessment preliminary
10 results. And then we will talk about the sensitivity
11 analysis results that have been obtained so far. Then
12 some effects of treatment of data, especially variance
13 and uncertainty on the expected dose estimation. Then
14 finally we will talk about the preliminary risk
15 insights from the sensitivity and uncertainty
16 analysis.

17 Under the sensitivity analysis results,
18 that is the third bullet, we have three specific
19 presentations: One is characterized as the parametric
20 sensitivity analysis, then we will talk about
21 distributional sensitivity analysis, then the third
22 one will be with a subsystem of barrier component
23 sensitivity analysis. I will be talking about the
24 first two bullets, and a portion of sensitivity
25 analysis results, especially distributional

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1 sensitivity analysis and subsystem of barrier
2 component sensitivity analysis.

3 First, here are the purposes of the
4 analysis. As you all know, NRC staff, in conjunction
5 with the staff from the Center for Nuclear Waste
6 Regulatory Analysis, have been involved over several
7 years in developing the Total System Performance
8 Assessment Code. The TPA Code represents an
9 independent approach to assist NRC's review of DOE's
10 performance assessment.

11 NRC's performance assessment tools are
12 intended to be used for gaining risk insights and to
13 risk inform the pre-licensing and the potential
14 licensing activities proactively and reactively. For
15 example, the development of the Yucca Mountain Review
16 Plan that you will hear about tomorrow, the
17 development of analysis tools by various key technical
18 user groups, or KTIs, and the confirmatory testing,
19 all these have been and will continue to be influenced
20 by the analysis that is performed using the Total
21 System Performance Assessment or the TPA groups of
22 tools.

23 As far as the reactor work is concerned,
24 staff is particularly looking at improving capability
25 to review license applications, such as DOE's

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1 performance assessment results and probe DOE's
2 assertion regarding the repository performance,
3 identify probabilities, such as risk dilution. Some
4 of these examples we will cover during the course of
5 this presentation.

6 Staff will also look at DOE's sensitivity
7 and uncertainty analysis approaches and also will try
8 to identify by doing its independent analysis which
9 model assumptions analysis and what is the degree of
10 importance of all these to the overall performance.
11 And it will also verify DOE's assertion regarding the
12 barrier importance.

13 These activities will require staff's,
14 one, understanding of the system as a whole, therefore
15 getting into various components of the total system is
16 very important. Therefore, staff will use these tools
17 and knowledge in understanding the system as a whole
18 and understand the factors that are important to
19 safety performance.

20 Here I will give you just a very brief
21 background before we move on to the results. These
22 are also some of the caveats in the sense that you
23 have heard about the results -- you have heard the
24 results from TPA 3.2 sensitivity analysis in the past,
25 and this represents the latest -- DOE's latest design,

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1 which it designed. And we have not done any analysis
2 using DOE's low temperature concept, because their
3 high temperature concept is considered as the normal
4 case.

5 The Total System Performance Assessment
6 Code, or the TPA Code, currently has about 950
7 parameters, out of which 330 parameters are samples.
8 So this is a pretty large problem for any kind of
9 Monte Carlo analysis on conducting sensitivity and
10 uncertainty analysis. So that means 620 parameters
11 are not sample, they're fixed at constant values at
12 what we believe as the best available value. And if
13 necessary, those values can be varied if we want to
14 support the current sensitivity analysis.

15 The results that will be presented
16 alternative conceptual models -- the results on
17 conceptual model analysis will not be shown for the
18 second time. However, in the context of the Total
19 System Performance Assessment, conceptual model
20 studies are done on a case-by-case basis, alternative
21 conceptual model studies.

22 And we would like to add the note that
23 analysis are performed mainly for developing staff
24 understanding, and the analysis that would be
25 presented are not necessarily mandated by the

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1 regulatory requirements. And the results are
2 preliminary in the sense that this sensitivity
3 analysis is currently under development. The report
4 is not ready. So what you are seeing today is a
5 snapshot of the results that we have come up with so
6 far. The results will be perhaps finalized in several
7 months.

8 Here I will start with the performance
9 assessment results. The performance measure is the
10 peak expected dose to the reasonably maximally exposed
11 individual. And the results will be shown essentially
12 for two scenarios. The first one is the nominal case
13 scenario, which is characterized by the slow
14 degradation over time leading to ground water release.
15 And the disruptive events scenario only one we will
16 present here is the igneous activity. Other two
17 disruptive event scenarios are seismic activities as
18 well as faulting activity. However, seismicity is
19 included in the nominal case, whereas we are not
20 presenting results on faulting because there is no
21 sensitivity. We don't see more sensitivity to
22 faulting event results.

23 And as far as the nominal case scenarios
24 are concerned, essentially the risk is computed by
25 averaging the results from Monte Carlo realizations,

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1 which is in terms of dose as a function of time. And
2 then the peak is determined from the expected dose
3 curve. Whereas the disruptive event scenario requires
4 some specialized calculations because of the low event
5 probability. Therefore, special convolution has been
6 used to take into consideration all possible events
7 prior to the event time. Prior to the evaluation
8 time, if there are events, those should be
9 appropriately factored in so that we get a smooth risk
10 curve.

11 First, this is the result of the nominal
12 case scenario. In the figure, we are presenting dose
13 versus time, but before we go through the figure, let
14 me just highlight that by the time the regulation was
15 out, was finalized, this work was already underway.
16 Therefore, some of the things that you are seeing here
17 are still different from what is mandated by the rule.
18 For example, the well pumping rate is varied in these
19 calculations. The receptor group is located at 20
20 kilometers. Other than that, I think these are the
21 main ones that are different compared to the rule.

22 And just to highlight, what we have seen
23 so far is that there are no corrosion failures in
24 10,000 years, no seismic failures in the nominal case.
25 The nominal case is the one which is defined by

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1 probability pretty close to one, and we are presenting
2 here results from 350 realizations. So primarily the
3 doses are resulting from the initially defective
4 failure which is varied between one to 88 waste
5 packages. To compare that, we have a total of 8,877
6 waste packages, with each waste package having about
7 7.89 MTU of spent fuel.

8 CHAIRMAN HORNBERGER: Sitakanta, what does
9 that probability approximately one, what does that
10 mean?

11 MR. MOHANTY: Because this is the -- okay.
12 If we subtract the probability for the disruptive
13 events, such as volcanism, then it's a very large
14 number. This is number is pretty close to one.

15 Also, there are several important things
16 to observe in this figure. The rate curve represents
17 the expected dose curve, which is an arithmetic
18 average of the individual realizations which are
19 represented in this blue color. The dark blue color
20 is the 95th percentile curve, and the green color
21 represents the 75th percentile curve. What this
22 entails is that until about 6,000 years the expected
23 dose curve exceeds the 95th percentile. And
24 throughout the 10,000 years, the expected dose curve
25 exceeds the 75th percentile.

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1 So this gives some sort of indication that
2 the expected dose curve appears to be quite robust in
3 determining the expected dose. And the peak expected
4 dose is from the expected dose curve, and clearly this
5 indicates the expected dose curve is -- expected dose
6 arc is pretty close to 10,000 years. And to be exact,
7 in our calculation it is showing up at 9,769 years.

8 MEMBER GARRICK: Now, this is just from
9 defective failures.

10 MR. MOHANTY: These are all from defective
11 failures.

12 MEMBER GARRICK: Because this is not the
13 peak dose for much later times.

14 MR. MOHANTY: Right.

15 MEMBER GARRICK: Yes.

16 MR. MOHANTY: Okay. Corresponding to that
17 figure, here are some additional results. The figures
18 on the left represent the cumulative release from the
19 saturated zone because that is the end point after the
20 transport through the geosphere, and after that it is
21 the biosphere. So I'll talk first between the
22 biosphere and the geosphere. This is the release
23 rate, and the release rate -- cumulative release rate
24 are presented in the Y axis of this curve for 10,000
25 years here and 100,000 years here. And these values

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1 are presented in large scale so that you can see these
2 numbers which are very small, smaller than one. And
3 because these are smaller than -- some of these are
4 smaller than one, therefore the log of that is a
5 negative number here.

6 Clearly, it shows that technetium is
7 dominating, also iodine-129 and chlorine-36. And here
8 is the corresponding curve. And this indicates that
9 most of the dose, which is about 52 percent of the
10 dose, contributes and is coming from technetium-99 and
11 about 25 percent of the dose coming from iodine and 20
12 percent coming from neptunium-237, and others are sort
13 of insignificant in terms of dose contribution.

14 I have just put a figure from 100,000
15 years just for comparison purposes. This shows that
16 if you go beyond 10,000 years, the dominant -- the
17 same nuclides are dominating, but you can see some
18 finite values. You don't have to see -- there are no
19 negative numbers here in the log space.

20 Next, here is the result from the
21 disruptive event scenario, as I mentioned earlier.
22 The faulting event -- we are not showing the results
23 for faulting event, and the seismicity was included as
24 part of the base case, and we did not see any failures
25 in 10,000 years. So, essentially, this is a

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1 comparison between the nominal case scenario and the
2 igneous activity scenario.

3 So, clearly, this shows that the peak in
4 the igneous activity scenario, which has a recurrence
5 rate of ten to the one minus seven per year, the dose
6 -- the peak expected dose occurs much earlier compared
7 to the nominal case. As I mentioned earlier, in the
8 nominal case scenario, the peak expected dose occurs
9 close to 10,000 years.

10 And to obtain this smooth curve, we needed
11 about 4,200 realizations, coupled with the convolution
12 integral approach that one was used to obtain this
13 curve. And this drop here is perhaps because we have
14 not taken one step beyond 10,000 years. Because if we
15 take a step beyond 10,000 years, this line is going to
16 flatten out or will perhaps slightly go up.

17 For the early release, this peak from the
18 igneous activity event, which is 0.35 milirems per
19 year, that occurs at 245 years. And the dominant
20 radionuclide is americium-241, and this dose is
21 primarily because of high activity nuclides, which
22 americium-241 is one of them.

23 Okay. Next, I will briefly go over the
24 stability of the peak expected dose. As such, because
25 it is an expected dose, we should expect a lot of

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1 stability in that number. We are using 350
2 realizations, and we considered that to be quite
3 stable. But I also wanted to show you some variation,
4 what happens if we go much beyond 350 realizations.
5 In this table, that shows that we have gone beyond
6 500. We have gone all the way to -- in fact, we have
7 gone to 4,000 realizations. Here, this one shows only
8 up to 3,000 realizations. And it varies between 2.48
9 ten to the minus two milirems per year and 3.24 ten to
10 the minus two.

11 So, essentially, we don't see nice and
12 smooth conversions. And we did some investigation to
13 find out what might be the reason for that. It turned
14 out that when we plugged the peak dose as a function
15 of the number of sampling realizations, there are some
16 extreme values. That is what is causing this kind of
17 change in the peak expected dose value. And we have
18 noticed that this kind of realization shows up in
19 about one to 2,000 realizations. And this is
20 something we are continuing to investigate further.

21 Next, we will start with sensitivity
22 analysis. I will be talking about the distributional
23 sensitivity analysis and subsystem value components in
24 sensitivity analysis. And after me, Dr. Codell with
25 start with the parametrics sensitivity analysis.

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1 This distributional sensitivity analysis
2 is done primarily to understand how the peak expected
3 dose is going to be influenced if the distributional
4 function assumption that we have received from various
5 KTIs are not correct or at least to identify if there
6 are some areas where staff need to focus more to
7 determine if anything can be improved.

8 Two approaches we have followed. One is
9 using a fixed range from here to here for changing the
10 mean of the distribution by ten percent. So we have
11 shifted the mean by ten percent. And in the second
12 approach, we have completely changed the distribution
13 function type. That means if in the nominal case we
14 had a normal distribution, we changed that to a
15 uniform distribution to see if that has a major
16 impact. Similarly, if the distribution had a log
17 uniform distribution, we changed that to log normal,
18 because in log space we thought that would capture the
19 difference.

20 Instead of working with all 330
21 parameters, we thought maybe changing for the top ten
22 influential parameters that we have identified by
23 using other methods would be more appropriate, because
24 those parameters are already showing a lot of
25 sensitivity. That's why in this talk we will

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1 primarily focus on the top ten influential parameters.

2 And we have used two different sensitivity
3 measures. One is the change to the peak expected
4 dose, before and after changing the distribution
5 function type and an effective distance between the
6 CDFs. CDFs are constructed by using the peak dose
7 from individual realizations.

8 CHAIRMAN HORNBERGER: Sitakanta, when you
9 say you looked at the top ten percent in importance,
10 how did you determine that, from a different
11 sensitivity analysis?

12 MR. MOHANTY: Yes. Those were determined
13 from parametric sensitivity analysis that Dick is
14 going to talk about.

15 For the distributional sensitivity
16 analysis, the two kinds that I saw, these two figures
17 are showing their results. The ten percent shift to
18 the mean with a fixed range, the results are shown
19 here. And for the complete change to the distribution
20 function, the results are shown by these blue curves
21 -- bars.

22 Let me describe the results from the shift
23 to the mean by ten percent. Clearly, it shows that
24 when the distribution function type is increased, the
25 mean is increased by ten percent, there is a 150

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1 percent increase in the waste package flow
2 multiplication -- a 150 percent increase in the peak
3 expected dose because of a ten percent change for the
4 waste package flow multiplication factor.

5 The second one that came out to be very
6 important is the spent fuel dissolution, which is a
7 pre-exponential term that defines that dissolution --
8 spent fuel dissolution rate. Fifty-seven percent
9 change to the peak expected dose occurred because of
10 a ten percent shift to the distribution function.

11 Similarly, when we changed the
12 distribution function type we did not see that kind of
13 effects for the two that showed up as important when
14 the mean was shifted. Rather, the two that turned out
15 to be important are the drip shield failure time and
16 the neptunium retardation in alluvium. So, therefore,
17 this clearly indicates that staff should revisit and
18 determine if the input parameter distributions were
19 not carefully looked at, at least the ones that are
20 showing up as important in the sensitivity analysis
21 should be looked at further, because these effects can
22 be cumulative. So when we add these things up for
23 many parameters, many sample parameters, that could
24 influence the peak expected dose that we compute from
25 the nominal case.

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1 Next we'll talk about the subsystem of
2 barrier component sensitivity analysis. This analysis
3 is just an extension of the sensitivity analysis we
4 are doing, first, to the parametric sensitivity
5 analysis, distributional sensitivity analysis, then
6 the system can be broken down in many different ways.
7 One can break the system along the line of
8 subprocesses, but here we have broken it down along
9 the line of physical components, and we are primarily
10 interested in seeing how much sensitivity we are
11 getting from individual components. But then breaking
12 down these components are very subjective, because one
13 can have more components than what we have shown here.
14 But it appears to be adequate for our purpose.

15 But it is very important to highlight here
16 that this analysis should not be mixed with multiple
17 barrier analysis. This is not a proposal to do
18 analysis this way. Therefore, it should be clearly
19 noted that this analysis is not required by -- there
20 is no regulatory requirement for this kind of
21 analysis.

22 And I would like to draw your attention to
23 the representation of the repository in this column.
24 The repository can be viewed at the top as an
25 unsaturated zone. Then next to that we have -- below

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1 that we have the drip shield. Below that we have the
2 waste package, then waste form, invert, unsaturated
3 zone and the saturated zone. So here you are seeing
4 barrier components, but in the results that I will
5 present in the next two slides, we show only six
6 because unsaturated zone above the repository and
7 below the repository will be treated as one entity.

8 And we will also show results from the
9 one-on analysis, one-off analysis and cumulative
10 addition analysis, because they all provide different
11 insights into the system.

12 CHAIRMAN HORNBERGER: Sitakanta, I don't
13 think I full understood something. You said that this
14 was not intended to be an analysis of barriers, but
15 then I've lost track of why you're doing this.

16 MR. MOHANTY: We are doing this purely to
17 supplement the sensitivity analysis. We are trying to
18 group them together. It's one way of looking at a
19 group of parameters, so we thought maybe grouping the
20 parameters along the line of a physical entity makes
21 it easier to understand.

22 My purpose in showing that column in the
23 previous slide was that these should be viewed as
24 individual cases. Each column here represents one
25 case. The group on the left represents the one-off

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1 analysis; the group on the right represents one-on
2 analysis. And the first column under the one-off
3 analysis represents the nominal case. And the row at
4 the bottom these represent the percentage change.

5 I would like to draw your attention to
6 these numbers in the sense that the numbers on the
7 left-hand side these are changes with respect to the
8 nominal case results. The numbers on the right-hand
9 side, these are with respect to the case where all
10 barriers are suppressed, and the suppression of a
11 barrier is represented by the gray color. That means
12 if we go to the second column here, this shows the
13 drip shield as a barrier has been suppressed. Under
14 the third column, the waste package as a barrier has
15 been suppressed. So, therefore, this number -- when
16 the drip shield barrier is suppressed, the number at
17 the bottom shows that the peak expected dose changed
18 only by a factor of 34 percent. When the waste
19 package value was suppressed, the peak expected dose
20 changed by 62,200 percent. So likewise, these numbers
21 represent changes with respect to the nominal case
22 result. But these are in percentages.

23 CHAIRMAN HORNBERGER: And can you
24 enlighten me just a little bit by what you mean by
25 suppressed?

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1 MR. MOHANTY: By suppression, we mean that
2 the function of the -- okay, from a purely technical
3 point of view here, the drip shield fails at a certain
4 time; it has a distribution. By suppression here we
5 imply that drip shield failure has been shifted back
6 to time zero. That means drip shield may have failed
7 at time zero, but if there is no infiltration because
8 of the thermal hydrology until 10,000 years, no water
9 is going to contact the waste package.

10 CHAIRMAN HORNBERGER: So the drip shield
11 -- the suppression of the drip shield is equivalent to
12 assuming that there is no drip shield.

13 MR. MOHANTY: Right, right.

14 CHAIRMAN HORNBERGER: Okay. Now, if we go
15 to the waste package, that's a little more difficult
16 for me. Does that mean that there is no waste
17 package?

18 MR. MOHANTY: Right. Here, what it -- let
19 me give you the detail here, if I can find my cursor
20 here. When the waste package is gone as a barrier,
21 what it implies is that the waste package has two
22 functions: when the waste package fails, and the
23 second is it contains -- it does not allow water to
24 enter into the waste package through flow
25 multiplication factor. Only a fraction of the waste

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1 package surface area will contribute to the water
2 getting into the waste package. Therefore, it will
3 come into contact to the spent fuel. So when the
4 waste package is gone, when the waste package is
5 removed as a barrier, that implies that the waste
6 package is failing at time zero. And, also --

7 MEMBER GARRICK: Does that also affect the
8 composition of the water?

9 MR. MOHANTY: No. So that is an important
10 point we want to make, that when we are doing this
11 analysis we are not changing the physical processes
12 that are going on.

13 MEMBER GARRICK: So you're really not
14 accounting for the interactive effects.

15 MR. MOHANTY: Right, because your purpose
16 is primarily the sensitivity --

17 MEMBER GARRICK: So this is not so much an
18 attempt to see the physical event and the progression,
19 as it is to deal with this question of sensitivity and
20 uncertainty.

21 MR. MOHANTY: Right. Yes.

22 MEMBER GARRICK: Okay.

23 MR. MOHANTY: But I think it is also
24 important to point out here that this removal, this
25 one-off analysis, especially when it comes to waste

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1 package, only affects the waste package that are
2 already seeing water. So two things are happening:
3 It is seeing water early and the second, more water is
4 getting into the waste package. But there are lots of
5 waste packages that do not see that water, as long as
6 the unsaturated zone barrier is above -- unsaturated
7 zone is above the repository horizon.

8 So, similarly, the numbers on the right
9 maybe they should be viewed as a decrease, so there
10 should be negative numbers. So here it means that
11 when the drip shield -- when all barriers are
12 suppressed and drip shield is added as a barrier, just
13 the only barrier, and the spent fuel would be in the
14 waste package somewhere here, but now we have no waste
15 package, in that case the drip shield has a
16 performance of about 63 percent. So this allows us to
17 see how much performance individual are coming from
18 these individual barrier components.

19 So here is shows that when the waste
20 package barrier is added to a case where all barriers
21 are suppressed, we have a 99.9 percent reduction in
22 peak expected dose. Whereas, when the unsaturated
23 zone is added, we have a reduction of 96 percent, and
24 when the saturated zone is added, when others are
25 suppressed, it's about 94 percent.

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1 So we have carried this analysis a little
2 further, and we have added to the one-on analysis.
3 Here we are adding those cumulatively. The first
4 column here represents when all barriers are
5 suppressed. In the second column, this shows that
6 this is similar to the saturated column you saw on the
7 previous page. But when we add unsaturated zone to
8 the saturated zone, it says that the peak expected
9 dose has been reduced by 99.2 percent. Of course
10 there are several decimal places that I'm not showing.

11 When we add invert to the unsaturated zone
12 and saturated zone, then that reduces to 99.6 percent.
13 And by the time we reach the waste package, this is
14 99.99, but maybe the number will change in about
15 seventh or eighth decimal place. So then when we add
16 all the barriers, all the barrier components, then we
17 regain the nominal case.

18 Then we grouped all these barrier
19 components together to reflect the engineered barrier
20 and the natural barrier. Clearly, as expected, as we
21 observed from individual component sensitivity,
22 clearly, when we group them together, it shows when we
23 compare that with respect to the nominal case, there
24 is a -- and when the engineered barrier is suppressed,
25 then we see a substantial increase in the peak

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1 expected dose, which is about 808,233 percent. And
2 when the engineered barrier system is there, but the
3 natural barrier is suppressed, it's about 58,233
4 percent.

5 I think that ends my presentation. Now
6 Dr. Richard Codell will take over.

7 DR. CODELL: Please don't adjust your
8 sets. We're experiencing technical difficulties here.

9 CHAIRMAN HORNBERGER: While we're waiting,
10 I'll interject and try to ask Sitakanta a tough
11 question. Having looked at all of that one-on and
12 one-off analysis, I'm not sure what message I'm
13 supposed to take from that.

14 MR. MOHANTY: We are continuing to conduct
15 this analysis. These results are quite fresh. We are
16 also trying to figure out how they are going to
17 contribute to the risk significance. The main reason
18 we did this kind of analysis is to see if any barrier
19 component in suppressing is shadowing the effect of
20 other barriers. So to determine how these individual
21 barrier components are performing, we had to separate
22 those out to individual ones.

23 For example, if we go to -- I think it
24 should be Slide Number 13, the one-on analysis for the
25 drip shield, we see that when all other barriers are

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1 suppressed, the drip shield reduces dose by 63
2 percent. That number -- we have not devised any
3 better method at this time to determine whether the
4 effect of drip shield is 63 percent or something else.
5 Simply by looking at one-off analysis we could not
6 figure that out.

7 Also, another reason for doing this
8 analysis is that if something is modeled, then we can
9 capture that effect in the traditional sensitivity
10 analysis. But if the model doesn't represent that,
11 then sensitivity analysis cannot capture it, because
12 it is not in the model. We do our best to capture
13 everything possible in the model, but there are also
14 uncertainty about the models themselves.

15 So there are two important aspects here.
16 One is uncertainty in the model themselves, number
17 one; and number two, is the shadowing effect of one
18 barrier component over others. So, therefore, by
19 adding it cumulatively, starting from the saturated
20 zone and coming up all the way to the level of drip
21 shield or the unsaturated zone above that, that gives
22 us some insights. Simply from those numbers we can
23 derive if there are any shadowing effects.

24 MEMBER GARRICK: I guess the thing that --

25 (Pause.)

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1 DR. CODELL: Maybe I should, to save time,
2 just work from the viewgraphs. Hopefully we'll be --
3 in a minute or two we'll have the presentation, and
4 I'll be able to put it up on the screen.

5 I cover parametric sensitivity analysis.
6 We're talking about the nominal -- we're on Slide 15.
7 We're covering nominal scenario only. And the purpose
8 is to determine sensitivity of parameters singly and
9 also in groups. The grouping is something new this
10 year. We're using -- there are two methods we're
11 using for parametric sensitivity. The first is
12 statistical methods that evaluate sensitivity to a
13 previously calculated pool of vectors that were
14 generated by the TPA 4.1 Code. In this case, we're
15 using generally 4,000 vectors cover the range of the
16 parameters. And then there are non-statistical
17 techniques that get to a sensitivity a second way,
18 which is to redirect the calculations to get the
19 maximum -- extract the maximum sensitivities from the
20 models.

21 We generally look at the peak of each
22 realization and look at the sensitivity of that, even
23 though the standard is based on something else; that
24 is, the peak of the mean dose. Starting on the next
25 slide, the statistical methods, these include

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1 primarily regression on raw and transformed variables,
2 non-parametric tests, like Kolmogorov-Smirnoff tests
3 and assigned tests, the parameter tree approach, and
4 there's some new work on cumulative distribution
5 functions sensitivity method and some other work
6 recently developed by Sitakanta Mohanti and Justin Wu
7 at the Center.

8 Another method along these lines is a
9 method which is based on the mean -- calculating the
10 sensitivity of the mean dose directly with respect to
11 the means of the independent parameters and also the
12 variance of the independent parameters. These are too
13 new to really go into any detail but they're
14 developmental.

15 The non-statistical methods include
16 differential analysis, Morris method and FAST method.
17 These are things that we covered before. There's one
18 new method that -- factorial design of experiments
19 which has the unfortunate acronym DOE, it's usually
20 called DOE.

21 (Laughter.)

22 This is something that John Telford, of
23 the Office of Research, and I have been working on for
24 several months with some pretty impressive results, I
25 feel. I'll go into that in a little more detail.

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1 Let me just take one second here to open
2 up the correct file.

3 (Pause.)

4 Okay. So we're at the bottom of Slide 16
5 now, starting at 17. The next slide shows -- this is
6 a tried and true method that we've been using for
7 several years now. We call it a composite statistical
8 method. This is to look at, in this case, six
9 statistical tests of various kinds, looking at
10 transformed and untransformed variables. And it's
11 really a seat-of-the-pants kind of method but works
12 quite well.

13 We used six statistical tests with 4,000
14 realizations. And then looking at each test and
15 factor the number of times the variable in question is
16 statistically significant in each test and its rank
17 and then develop a single list of parameters, top
18 parameters from the number of times they appear and
19 the ranks of the six tests. And when you do that you
20 come up with a list arbitrarily cut off at ten
21 variables for 10,000 and 100,000 years, showing that
22 a lot of the parameters that show up have to do with
23 how much water gets in contact with the waste.

24 I'll show some comparisons of the methods
25 later, but to get into the new work we've done, John

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1 Telford and I, the factorial design of experiments.
2 Basically, factorial design, in the simplest form, is
3 to look at two values of each of the variables, a high
4 and a low value. We took fifth and 95th percentile of
5 each distribution, and since there were 330 variables,
6 if you looked at all possible combinations, you'd have
7 two to the 303 runs required, which at the present
8 rate would take ten to the 94 years. And, of course,
9 in maybe 1,000 years this will be --

10 MEMBER GARRICK: Are you looking for
11 permanent employment?

12 (Laughter.)

13 DR. CODELL: In 1,000 years things might
14 be much better, but right now it's out of the
15 question. So the fractional factorial design is what
16 you have to use, but it gives you reasonable time
17 estimates, but it's somewhat ambiguous.

18 So the way we did it is looking at the
19 sampling iteratively, that is running a fractional
20 factorial and then using other information from the
21 runs to refine the list and then repeating on the
22 refined list several times until we're quite sure that
23 we've gotten most of the important variables. And
24 this took a lot of trial and error, but we think we
25 hit on a good procedure for doing this.

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1 The advantages of this technique is it's
2 systematic and potentially precise. It's easily
3 interpreted with a powerful, statistical techniques
4 like analysis of variance and trees. And it does
5 reveal interaction among variables instead of looking
6 only at sensitivity of single variables. I think this
7 is an important point. Disadvantages are it's still
8 costly and difficult to implement. And looking at
9 only the high and the low value, you're not looking at
10 the range of entire variable.

11 So this is how we went through the
12 procedure. For the 10,000 year case, first looked at
13 a design set using some statistical software of 2,048
14 variables, and we identified 100 potentially sensitive
15 variables. We reduced that list to 37 on the basis of
16 other information. For example, even though some of
17 these variables appeared to be sensitive like seismic
18 parameters, you could see from other results that
19 there weren't any failures, so you knew those
20 variables were confounded and could be eliminated from
21 consideration.

22 And the second screening -- that was the
23 first screening -- the second identified ten variables
24 and then we went into a full factorial with only ten
25 variables, which is a reasonable number to deal with,

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1 and identified six to eight sensitive variables. When
2 you do this and you go through the analysis of
3 variance, one of the byproducts is a tree diagram.
4 And this shows very clearly that if you follow the
5 path of the cursor here, a low value of drip shield
6 failure time gives you -- and a high value of the flow
7 multiplying factor, the diversion factor and the fuel
8 dissolution factor and the waste package effective
9 fraction leads to the highest dose.

10 So this kind of information is much more
11 revealing than looking at sensitivity of single
12 variables at a time. This same sort of information,
13 incidentally, comes out of what we call the parameter
14 tree method, which is a statistical technique, but
15 this is much more precise, whereas there's a lot of
16 uncertainty in the parameter tree approach.

17 The next slide shows the same sort of
18 result for the 100,000-year run, and it also shows the
19 high and the low value of each variable contributing
20 to the highest dose.

21 Just to show that we think we've captured,
22 with a very small number of variables, most of the
23 uncertainty, the next few slides reconstruct some of
24 the results of the original run with the 330 variables
25 and the reduced set, both either from regression or

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1 from fractional factorials. This is a cumulative
2 distribution function of the peak doses showing that
3 especially for the high end of the dose curve that
4 we've captured most of the uncertainty just with ten
5 variables from the regression analysis or eight
6 variables from the factorial design method. The lower
7 curve shows the mean dose for the same calculations,
8 330 versus ten for regression and eight for the
9 factorial design. So even though it's not perfect,
10 we've captured most of the uncertainty with a very few
11 number of variables. And the same is true for the
12 100,000-year result.

13 Now, the next set of slides, moving away
14 from the factorial design now, there were two options
15 for looking at sensitivities. The first one is that
16 we have -- we can look at the peak of each individual
17 run and look at the sensitivity based on the number or
18 we could look at the time of the occurrence of the
19 peak of the mean and look at that. The upper graph
20 shows the sensitivity result looking at the mean dose;
21 the bottom is looking at the peak of the individual
22 doses. Except for the first two columns here, the
23 results are quite different, and we're tempted to say
24 that the sensitivity measure and statistical parlance
25 has more power using the peak dose from each

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1 individual run rather than the mean.

2 But there's one interesting factor here.
3 If you look at this particular variable, drip shield
4 failure time came out about number 20 on this measure
5 using the mean of the peak dose, and yet it came out
6 quite high looking at the individual peak doses. This
7 is not an error. This is -- there's a real reason for
8 this that isn't obvious, and the next couple of slides
9 really get to this -- why does drip shield failure
10 time differ?

11 MEMBER GARRICK: Dick, could you say
12 something about the sensitivity measure, what it
13 really is?

14 DR. CODELL: Well, I'll let Sitakanta
15 address that. He prepared these slides.

16 MR. MOHANTY: For the sensitivity
17 analysis, we need a point value. That means we have
18 dose as a function of time, but when we do the
19 analysis it has to be a point value that represents
20 for the 10,000 years. So it's a matter of whether we
21 should choose the peak from that realization or should
22 we choose the value, dose value corresponding to the
23 peak expected dose? That will be the point value.

24 So in other words, the red bars, those are
25 showing the sensitivity analysis using the dose values

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1 corresponding to the time when the peak expected dose
2 occurred. Whereas the figure at the bottom that is
3 indicating that peak can occur any time in 10,000
4 years so that it is independent of the time of
5 occurrence. So, therefore, it reflects sort of the
6 whole time domain, whereas the one at the top
7 represents a particular time.

8 CHAIRMAN HORNBERGER: But what are the
9 units on that sensitivity measure?

10 MR. MOHANTY: Oh. We have different
11 sensitivity measures from different methods. This
12 particular one is representing one method that we have
13 used for the two graphs. So that measure is
14 essentially, but it's kind of hard to explain. This
15 is extracted from the Morris method in which we take
16 the sensitivity from individual points and average
17 that, and we determined the mean of the ΔY over ΔX
18 where X is the variable that is being changed. ΔY
19 is the dose value that is being changed. So this is
20 an ensemble statistics so that sensitivity measure is
21 based on the ensemble statistics, both mean and
22 variance.

23 MEMBER GARRICK: But it is in a change in
24 dose per unit change in parameter.

25 MR. MOHANTY: In the parameter, right.

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1 DR. CODELL: Okay. Thanks, Sitakanta.

2 CHAIRMAN HORNBERGER: But you must
3 normalize somehow.

4 MR. MOHANTY: Yes. The parameters are
5 normalized.

6 CHAIRMAN HORNBERGER: Okay.

7 DR. CODELL: On the next slide, I wanted
8 to talk about the treatment of data variability and
9 performance assessment modeling. This particular
10 piece of work came up during the SAS review of the
11 TSPASR and the SSPA, and it was called the galsean
12 variance partitioning. It was basically how you treat
13 data in the model.

14 It isn't exactly how DOE is doing it, but
15 it leads us to some interesting conclusion on how we
16 should deal with experimental data uncertainty either
17 because of lack of knowledge or variability. And the
18 difference between these two kinds of uncertainty,
19 epistemic and aleatory, is often blurred, for example,
20 treatment of corrosion rate data for the waste
21 packages and its effects on dose.

22 To get at this phenomenon, we put together
23 a model based very loosely on NRC's model and DOE's
24 model, but it's a separate model. NRC's TPA model we
25 represent variability and waste package corrosion by

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1 a few representative waste packages -- only one per
2 subarea, ten in all. Whereas DOE has in its Total
3 System Performance Assessment uses the patch failure
4 model that allows significant spatial variability of
5 failures.

6 We could look at the data on corrosion,
7 and we could say it's either -- this is real data from
8 the corrosion experiments on the coupons and say it's
9 either a fixed but uncertain rate or a spatially
10 varying rate due to the material and environmental
11 variability.

12 On the next slide, I showed this very
13 simple model that deals only with a few parts of the
14 model, particularly the waste package corrosion and
15 the dissolution of waste in a fixed number of years
16 once the waste package has failed. Now, there are
17 three possible models. Model 1 is the whole
18 repository. That's where you take this corrosion rate
19 data, shown here as a density function, and you apply
20 it to each and every waste package identically; that
21 is, they'll all fail at exactly the same rate, pretty
22 much, there is some slight variation, but at the same
23 time. Whereas the other extreme is Model 3 where each
24 patch of each waste package is sampled from the
25 distribution so that each and every waste package and

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1 each and every patch has a different failure time.
2 And Model 2 is in between those two extremes.

3 Now, if you take this model and you just
4 look at, for the present time, five realizations, as
5 shown in this figure, you'd see that Model 1, where
6 every waste package fails at about the same time,
7 gives you five individual peaks, and they're all
8 rather high, because when they fail at the same time
9 you get a big release and therefore a big dose.
10 Whereas Model 3, where you have this patch failure,
11 they're all pretty much the same and smaller.

12 The interesting thing is that I wanted to
13 point out here is that the dose and the way the NRC
14 has defined in the rule as the peak of the mean is
15 very sensitive to the timing of the peaks, so that
16 even though these individual peaks are high, when you
17 look at the way that Model 3 always fails the same,
18 each new realization looks pretty much like the last
19 one, these all line up. So when you take the peak of
20 the mean dose, Model 3 actually gives you a higher
21 dose, which I will show.

22 And how does that relate to a few slides
23 before where I showed the drip shield failure time
24 being an important parameter? Well, drip shield
25 failure time determines the timing of the dose. If it

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1 fails early, then the release is early. If it fails
2 late, the release is late. That's the same effect as
3 changing Model 1 to Model 3. And that's why it showed
4 up in one way when you look at the peak of the mean
5 dose and another way when you looked at the peak to
6 the individual doses. But that was an interesting
7 conclusion.

8 MEMBER GARRICK: Again, that's dependent
9 upon the corrosion model that you --

10 DR. CODELL: Well, yes. And the way we've
11 treated drip shield failure time in the TPA model is
12 just a sample failure time. It just relates -- it's
13 just a -- it could have been another example of the
14 same phenomenon.

15 MEMBER GARRICK: Right. And the other
16 thing here is that there's going to much greater
17 variability in the setting than in the waste packages.
18 So whatever you take advantage of with respect to the
19 similarity of the waste packages could be very offset
20 by the variability because of spatial considerations.

21 DR. CODELL: It could be but we don't have
22 enough information from the corrosion rate data to
23 know which is which, and that's a dilemma. A very
24 important factor in our analysis is how quickly the
25 waste packages will corrode. And even though it seems

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1 to be a very long time, if it were not a long time,
2 then we'd want to know whether the variability in the
3 data was due to real spatial differences or
4 experimental -- or other unquantifiable errors.

5 MEMBER GARRICK: And, of course, in the
6 DOE model, they'd decouple the drip shield
7 contribution from the diffusive transport out of the
8 waste package.

9 DR. CODELL: Yes.

10 MEMBER GARRICK: So it really depends upon
11 how you structure the thing. I'm curious about how
12 you screened your parameters.

13 DR. CODELL: I'm sorry, which slide were
14 we?

15 MEMBER GARRICK: Well, when you go from
16 900 to 330, to 100, to 37.

17 DR. CODELL: Well, actually, the 990 were
18 screened by experience. We've, at various times in
19 the past, looked at all those variables varying and
20 decided that most of them didn't contribute anything
21 to the results. So those were held at fixed values.
22 The screening that took place in the factorial design
23 was more systematic, because we started with 330 and
24 worked our way down. And it was either based on the
25 sensitivities we observed in the analysis or variance

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1 or --

2 One of the problems with fractional
3 factorial design is a problem called confounding, and
4 that's where a variable can be mistaken -- sensitivity
5 in a variable can look like it's sensitive but it's
6 actually a combination of several other variables.
7 And it's just a numerical combinatorial problem, not
8 a real physical problem. But by looking at the
9 physical outputs of the code, for example, seeing that
10 the factors that looked sensitive, that had to do with
11 seismicity, couldn't have been because there weren't
12 any failures due to seismicity. So we could eliminate
13 those. So it took a little bit of a combination of
14 the silicon and the carbon computers to reach this
15 conclusion.

16 MEMBER GARRICK: Did you call this the
17 confounding phenomenon?

18 DR. CODELL: Yes.

19 MEMBER GARRICK: That's a good name.

20 DR. CODELL: It's not my -- that's what
21 it's called in the factorial design method.

22 CHAIRMAN HORNBERGER: How are you sure at
23 the end of the day that you don't have some aliasing
24 left in your final ten or whatever --

25 DR. CODELL: Well, there can't be any when

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1 you do the full factorial.

2 CHAIRMAN HORNBERGER: Oh, right.

3 DR. CODELL: That's the --

4 CHAIRMAN HORNBERGER: So once you choose
5 the ten it's okay.

6 DR. CODELL: Right. Yes. And the final
7 test was seeing how well you did by comparing it to
8 the original.

9 Well, getting back to this little
10 experiment on the two kinds of uncertainty, the
11 epistemic and aleatory, the first result shows for a
12 full set of realizations that the peak of the mean
13 dose the Model 3, where you have the patch failure,
14 gives you the highest result. This may seem
15 counterintuitive, but as it turns out, if you're
16 sampling each and every patch, you end up getting the
17 similar kinds of failures each new realization. And
18 that's why they look identical and fall on top of each
19 other, leading to a high peak of the mean dose. And
20 the other models give you much lower doses.

21 However, this is sensitive to other
22 parameters in the model, and what we determined when
23 all was said and done was that if you look at a much
24 slower release, say 100 times slower than what we used
25 in this example, all three models pretty much fall on

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1 top of each other. And when looking at the ranges of
2 parameters in the Department of Energy model, probably
3 it is more like the case on the right more than the
4 left. But it's still an interesting phenomenon and
5 explains some other interesting features like the drip
6 shield failure time result we got.

7 Something related to this previous
8 exploitation is risk dilution. This is something we
9 worry about. It's not good enough just to increase
10 the range of distribution if you don't know it.
11 There's some cases if you do that, you'll end with
12 actually a lower dose, which isn't what you wanted at
13 all. And here's an example. Once again, drip shield
14 failure time. If you have a narrow range, this green
15 curve, or a wide range, the blue curve, you'll get
16 different results. And the narrow range gives you a
17 higher dose than the wide range. Once again, this is
18 one of the parameters that has to do with the timing
19 of the doses, and when you increase the range of that,
20 you're going to end up with a lower result.

21 So summing this all up --

22 CHAIRMAN HORNBERGER: Dick, let me -- I
23 don't know, I think I'd like to challenge you on that
24 one, because you said you put in a wide range and you
25 may get a lower dose, and that isn't what you want.

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1 If in fact you have a broad uncertainty range, why
2 isn't it what I would want?

3 DR. CODELL: Well, I think the --

4 CHAIRMAN HORNBERGER: I mean if you really
5 -- if your uncertainty and failure times for drip
6 shield really is -- I mean I suppose it ties back into
7 your aleatory versus epistemic, because if you really
8 believed that every single drip shield was going to
9 fail on day 372, then it really matters. Is that what
10 you're saying?

11 DR. CODELL: That's right. That's right.
12 Yes. And this is interesting because I think prior to
13 NRC's regulations for high-level waste, I think most
14 people considered looking at the peak of the
15 individual doses as a factor. And automatically if
16 you put a wider distribution in, you're going to get
17 a higher -- one of those is going to be a higher peak.
18 But it doesn't work that way when you look at the peak
19 of the mean curve; it's just the opposite.

20 CHAIRMAN HORNBERGER: Which I assume is
21 why even though the regulation calls for the peak of
22 the mean dose, you will require the potential licensee
23 to display all sorts of things, including all of the
24 uncertainty?

25 DR. CODELL: Well, I wouldn't go too far

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1 there. I think I'd be stepping out of bounds. I
2 don't think we would require anything like that. If
3 Tim is in the audience, he could probably rescue me
4 right now.

5 CHAIRMAN HORNBERGER: Maybe I could frame
6 it another way. The ACNW will want to see that.

7 DR. CODELL: Yes.

8 MR. McCARTEN: Well, as Dick's slide
9 indicates, I mean the key there is the inappropriate
10 use of a wider distribution. We are clearly
11 interested in the distribution. And if the
12 uncertainty is there, we're not saying don't include
13 the uncertainty you have.

14 If there are some arbitrary decisions that
15 are done and sometimes done in the sense of
16 conservatism, let me make this bigger, I'm uncertain
17 about it, we want to look at that to make sure, well,
18 you may think it's conservative to make it bigger but
19 you've actually, in essence, produced a lower dose.
20 And so you want to have an appropriate range. As Dick
21 indicated, I think we are going to look at all the
22 information the Department gives us.

23 MEMBER GARRICK: Let me understand
24 something. Is this distribution a random variable?

25 DR. CODELL: Yes.

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1 MEMBER GARRICK: Because what we're really
2 interested in is our uncertainty about a fixed
3 variable.

4 DR. CODELL: Well, it actually is an
5 uncertainty about a fixed variable in almost every
6 case. I think in every case in the TPA Code it's
7 uncertainty about a fixed variable. I would consider
8 that a definition.

9 MEMBER GARRICK: And if that's the case,
10 then of course you know want to be very careful about
11 manipulating a broad distribution into -- or a peak
12 distribution into a broad distribution as to what
13 information you might be losing in that process.

14 DR. CODELL: Right. Well, these density
15 functions that we use are based on either data or
16 people's idea of what the data should look like. And
17 it isn't always -- they aren't always precise.

18 MEMBER GARRICK: Well, as George says,
19 we're going to be very interested in following this.

20 DR. CODELL: Preliminary insights from the
21 sensitivity analyses for 10,000 years, factors that
22 control water/fuel contact seem to be the most
23 important and most doses from low retardation, long
24 half-life radionuclides, like technetium. For 100,000
25 years, it's interesting that waste packages usually

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1 fail by 100,000 years, so the parameters aren't always
2 showing up as being conservative, because you'll
3 usually have failure anyway. Changing them isn't
4 going to make any difference.

5 The fuel/water contact is still important,
6 and the dominant radionuclide, neptunium-237, seems to
7 be very important, so parameters associated with that
8 are important. For barrier sensitivity, the
9 preliminary results that both natural barrier and
10 engineering barrier make substantial contributions.

11 This is some additional work in progress
12 that was too callow to talk about today, but we've
13 acquired some neural network software, which seems to
14 be very powerful, and this is basically doing non-
15 linear regression. We've had some limited success
16 with it so far.

17 Dave Esch and I took some training in it,
18 and I think you'll probably see this next we make a
19 presentation. We're looking at new sensitivity
20 measures consistent with the peak expected dose, as we
21 showed in the previous slide, and looking for
22 efficient distributional sensitivity methods like the
23 cumulative distribution function sensitivity that
24 allows us to look at the sensitivity at different
25 parts of the cumulative distribution; that is, high

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1 dose and medium dose or low dose and also the means --
2 the sensitivity of the mean dose directly.

3 Some other work, we're trying to get a
4 handle on barrier performance in a couple ways. We're
5 trying to define what a degraded state of a barrier
6 means. This is a very difficult problem trying to
7 figure out how to define a barrier as failed, like
8 what does a failed waste package mean.

9 Just looking at the kinds of barrier
10 sensitivity analysis that Sitakanta presented earlier,
11 there are six barriers, so two to the six is 64
12 possible combinations of failures, from everything
13 failed to everything working. There has been 29 of
14 the 64 analyses completed, and we've made some
15 preliminary shot at making a tree structure, but it's
16 possible to draw a tree with this result but looking
17 more powerfully at -- looking at it with more powerful
18 methods like analysis of variance there's not enough
19 runs yet to do that, but we're hoping to do that in a
20 future presentation.

21 In conclusion, parametric sensitivities
22 provide useful risk insights. The method we've been
23 using, the sensitivity method where we're combining
24 ranks of the various statistical methods still works
25 very well.

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1 Factorial design shows great promise and
2 clearly defining the sensitivities and the
3 interactions of the variables. The distributional
4 sensitivity technique that Sitakanta presented is an
5 effective approach identifying the impact of the
6 choice of parameter distribution and the shape and the
7 shift in the mean. We've shown that inappropriate
8 parameter ranges can lead to risk dilution in some
9 cases, and the treatment of uncertainty as lack of
10 knowledge, epistemic or variability, can affect the
11 peak risk calculation. That's the end of our
12 presentation. We'd be happy to take additional
13 questions.

14 CHAIRMAN HORNBERGER: Actually, before --
15 now that John is back. I started before you got back
16 from lunch, John, but this is really your bailiwick,
17 so why don't you run it.

18 MEMBER GARRICK: All right, well, let's
19 see if there's some questions. Of course, I have a
20 few.

21 Milt?

22 MEMBER LEVENSON: One comment.

23 MEMBER GARRICK: Microphone.

24 MEMBER LEVENSON: One comment and then one
25 question based entirely on ignorance. One of the

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1 things is that I guess I sort of disagree with your
2 use of terminology because no matter what you do in
3 the way of assumptions, you are not going to change
4 the risk. So you can't dilute the risk or increase
5 the risk. You may change your calculated number, but
6 it's not really the risk.

7 But on Slide 13, I'm having trouble
8 relating this to the physical world in that on the
9 one-OFF, if you remove the waste package, you have a
10 62,200 percent change.

11 MEMBER GARRICK: Can we see that on the
12 screen? Can you put the projector back on, please?

13 On the one-OFF analysis, when you remove
14 the waste package, you have a 62,200 percent change,
15 but with the one-ON analysis, none of the other
16 barriers are there. You just add the waste package.
17 You only have 100 percent change. Factor 2. It
18 doesn't seem consistent with the physical world, as I
19 visualize it.

20 MR. MOHANTY: Let me explain the
21 difference. Under the one-OFF analysis, the first
22 column represents the nominal case. For the nominal
23 case, the peak expected dose is .021 millirems per
24 year, whereas under the one-ON analysis, we are
25 determining the percent in change based on the first

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1 column under one-ON analysis. And that number, I
2 don't remember what that value is, but we are using
3 that number to determine the change. So that means at
4 most it can be that. So when we put the waste
5 package, so 99.9 percent represents a reduction in
6 what we observe under column 1.

7 CHAIRMAN HORNBERGER: You can't take away
8 any more than 100 percent. But if you have something
9 to start with, you can change it by 62,000 percent.

10 MEMBER LEVENSON: I guess without the
11 numbers, it's very difficult to determine what the
12 significance of what this chart is.

13 CHAIRMAN HORNBERGER: Even with the
14 numbers, I would maintain it's very difficult to
15 figure out what the significance of what this charge
16 is.

17 (Laughter.)

18 I don't mean to be too severely because I
19 know we are interested, like you are in barrier
20 performance. But it strikes me, Dick just casually
21 said it's a really difficult problem to figure out
22 what it means to have a barrier suppressed. And I
23 agree with that. It just doesn't make sense to me to
24 even consider changes as if all the drip shields
25 failed at Time Zero. I don't understand what you're

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1 doing.

2 MEMBER LEVENSON: It's not so much to
3 understand what you're doing. It isn't clear to me
4 what the significance is.

5 CHAIRMAN HORNBERGER: Right.

6 MR. HAMDAN: One of the main objectives of
7 this is one that sensitivity analysis is one that is
8 not risk, but in response to your question and that is
9 to test one other. And one could argue that these
10 barriers individually or in combination is to see if
11 one is working and this has not been in all these
12 slides clearly that you can elicit in slide 3, clearly
13 the emphasis also I think he did answer and answer
14 very well. But the question as to what added value
15 the sensitivity adds to the model and whether the
16 model has been improved has not been addressed.

17 MEMBER GARRICK: Yes, but the black box
18 here is the degree to which the model represents
19 reality and I think that's part of what Milt is
20 struggling with.

21 You know, it's this question of if you
22 tried to look at this as a system and you apply the
23 basic equations of continuity and conservation of mass
24 and momentum, etcetera, etcetera and you flow through
25 the system, this model isn't doing that because the

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1 800-pound gorilla here is the water and the chemistry
2 of the water. And the chemistry of the water is
3 extremely sensitive to each of the stages it passes
4 through.

5 So we're not talking about something
6 that's so much represents reality as we are talking
7 about some very interesting concepts that you can
8 apply in a Monte Carlo-type analysis, but at least
9 that's my perspective.

10 Ray?

11 VICE CHAIR WYMER: I'll be a little bit
12 facetious. I certainly admire the sophistication and
13 complexity of these analytical tools and the variety
14 that were used in these analyses and that's
15 impressive, but I couldn't lay a finger on it myself.
16 But I was pleased to see that in your preliminary
17 sensitivity analysis that you confirmed what I thought
18 for the last two or three years that --

19 CHAIRMAN HORNBERGER: Chemistry is
20 important.

21 VICE CHAIR WYMER: Natural version as a
22 substantial contribution, that looks pretty good.

23 (Laughter.)

24 Waste packages fail corrosion parameter is
25 not sensitive. Fuel water contact, that's important,

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1 pretty good.

2 And retardation of neptunium seems to me
3 like that's important. But I thought I knew that
4 stuff.

5 Factors controlling water fuel contact
6 dominate performance. That's right. And most dose
7 from low retardation and long half-life radionuclide,
8 sure, I know that trivializes the degree to which you
9 understand these things and the sophistications, but
10 nonetheless the answers are sort of self-evident for
11 whatever that's worth.

12 MEMBER GARRICK: George?

13 CHAIRMAN HORNBERGER: I just wanted to
14 make a final comment on the barrier component
15 sensitivity. I really do understand what Latif said
16 and that is that you do a lot of these things to try
17 to understand what's going on with your modeling. I
18 certainly have no problems with this.

19 The issue that I have, the difficulty I
20 have with slides like this is that there's too much
21 chance for mischief making with the numbers by people
22 who will want to use them for purposes that are not to
23 understand how your model is working. And I guess I'd
24 just ask you to give a little thought to that as you
25 present these things.

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1 MR. CODALL: We've given a great deal of
2 thought to that. In fact, at every level of review,
3 we've been asked to be sensitive to this and to put
4 disclaimers in that this not underlying not required
5 by the regulations.

6 I think people who want to make mischief
7 of this will do so regardless.

8 But this is the kind of analysis that's
9 often done for safety. You look at the failure of a
10 system. You look at what happens when an engine on an
11 airplane dies. It's nothing not wrong with it, in my
12 opinion. It's just my opinion.

13 MEMBER LEVENSON: I think there's a
14 significant difference in fact, that's one of the
15 points I think George made earlier that an engine is
16 either on or off and you can -- this is a legitimate
17 analysis for that sort of thing. The waste container
18 is not either in existence or not in existence, and
19 therefore, I think you have to be very careful about
20 using what is a legitimate analysis under other
21 conditions or this one that might be much more
22 legitimate to say what happens if 10 percent of the
23 waste packages fail early, etcetera, rather than --
24 they're either on or off.

25 MEMBER GARRICK: Yeah. I think that

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1 there's no question that the modeling test exercise
2 that you've done here is very interesting and very
3 powerful. As I was saying earlier thought, I think
4 that what we're really interested in is information
5 that would give us confidence that the models that are
6 being employed are doing a reasonable job of
7 representing reality in terms of what's happening.

8 Now maybe this can contribute to that, but
9 what is really something that concerns us is the
10 interactive effects of these different barriers and
11 how the one thing that would suppress some of the
12 mischief that we talk about would be to do this same
13 exercise for different models.

14 Take for example, the TPA model and do the
15 exercise, and then take the diffusive transport model
16 of DOE and do the exercise and you would certainly see
17 that things would line up differently. And it would
18 clearly indicate that how model sensitive it is.

19 But again, I guess the question I would
20 ask is what contribution comes from this work towards
21 creating a model that we have increased confidence in
22 in terms of representing the performance of the
23 repository?

24 MR. MOHANTY: Let me start with one-ON
25 analysis. What that figure tells us is that on

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1 saturated zone, unsaturated zone is making quite a bit
2 of contribution and these individual contributions
3 perhaps could not have been seen if we did not isolate
4 those from other either components or in a broad sense
5 subsystems.

6 So that tells us something. And also when
7 we compare that say with an invert, we are seeing only
8 2 percent change. Then we are going back to the total
9 system for performance assessment code to determine
10 why we are only getting .2 percent and we did go back
11 and find that the way in what is modeled in what is
12 supposed to reduce flow or delay transport, but it
13 just so happens that the flow through invert is
14 predominantly fracture flow.

15 So when the flow is a predominantly
16 fracture, in the fractures and we are not assigning
17 any retardation fractures to the fractures, therefore,
18 the invert is almost completely bypassed in the TPA
19 approach. So that is a kind of insight we gain when
20 we do this kind of one-OFF one-ON analysis or
21 cumulative analysis.

22 VICE CHAIR WYMER: I would have liked to
23 have seen that sort of thing in your table of
24 preliminary insights, the two cases you just cited are
25 much more informative.

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1 MR. MOHANTY: If I can make another point
2 that there are two ways we can determine all
3 components for how well the components of the code is
4 functioning. To give you an example, if the packages
5 were going to fail at 1 million years, then the only
6 way we can find out that what is what is affecting the
7 packages is we go with continual calculus for 1
8 million years, are we deliberately suppressed that to
9 find out what the impacts are if we are to fail early.

10 So by doing this kind of analysis that
11 prevents us from going to much further into the future
12 to million years because we can gain similar kind of
13 insights by deliberately doing the sensitivity
14 analysis by suppressing components.

15 CHAIRMAN HORNBERGER: I have a question
16 now that you have mentioned your one-ON analysis. If
17 we look at that lefthand column, okay, you have a dose
18 associated with that, is that right? What you said
19 was that we could read those as 99.9 percent reduction
20 in dose?

21 MR. MOHANTY: Right.

22 CHAIRMAN HORNBERGER: So my question is
23 what is the dose in that lefthand column and how did
24 you calculate it?

25 MR. MOHANTY: Under the one-OFF analysis

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1 or one-ON analysis?

2 CHAIRMAN HORNBERGER: One-ON analysis.

3 MR. MOHANTY: On the one-ON analysis, this
4 is not a real dose where barriers are suppressed. We
5 do honor the various processes in the TPA code. We do
6 not veer away from the processes too far because we
7 know that we are limited simply by this is just a
8 technique we are using.

9 CHAIRMAN HORNBERGER: All right, so then
10 if I go to the first column, the drip shield, the 63
11 percent reduction in what?

12 MR. MOHANTY: We do have a dose value.

13 CHAIRMAN HORNBERGER: How did you
14 calculate it?

15 MR. MOHANTY: This is by suppressing all
16 components.

17 CHAIRMAN HORNBERGER: Okay, which means
18 what did you do, dissolve all the fuel instantaneously
19 in the --

20 MR. MOHANTY: Yes.

21 CHAIRMAN HORNBERGER: And so it's a high
22 dose.

23 MR. MOHANTY: It is a high dose, yes.

24 CHAIRMAN HORNBERGER: I mean that's a high
25 calculated dose. I don't want to get in trouble with

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1 --

2 (Laughter.)

3 So I just point out that again, even on
4 this one we're saying oh yes, the natural, the insight
5 that you gain, it's quite artificial and I'm much more
6 comfortable with Latif's interpretation that what
7 we're doing is learning how the components of the
8 model are working.

9 MEMBER GARRICK: Also, those kind of
10 reductions in the kind of doses we're talking about
11 are not very relevant.

12 I don't know that that really tells us a
13 great deal about the protection provided by those
14 components. I'll have to think about that, a good
15 deal more. And I still worry about the fact that
16 there is interaction between the barriers and what the
17 waste package sees in terms of input material is
18 different than what the invert sees as different what
19 the unsaturated zone sees and so on.

20 And that could be a major factor in what
21 really happens. All the peer reviews of the TSPAs
22 have given great attention to the importance of water
23 composition because that's the mechanism by which
24 everything happens and that's just the process of
25 applying principles of continuity from the

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1 infiltration model, if you wish, namely the geology
2 above the waste package through to the waste package
3 and so on.

4 So again, there's no question this is an
5 intriguing process and it does, as Latif says, but
6 we're going to have to be a lot more diligent students
7 and studiers of this before we can really see what it
8 contributes to reality.

9 MR. CODALL: In terms of projected work,
10 we are getting together soon to talk about what
11 degradation of barriers means. I hope in the next few
12 weeks, Tim McCarten is convening a group to better
13 define in terms of what is expected in the regulations
14 what barrier degradation means.

15 This one-OFF, one-ON analysis probability
16 overkill.

17 CHAIRMAN HORNBERGER: yes.

18 MR. CODALL: But getting at the -- getting
19 a finer level is the next step. This maybe you
20 consider this a first step and gaining information
21 about importance of barriers.

22 MEMBER GARRICK: Go ahead, Latif.

23 MR. HAMDAN: But do we need a more refined
24 -- it seems to me that from this beautiful
25 presentation the staff has already has all the tools

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1 it needs what it needs to do from the subject. After
2 all, the barrier requirements which was about 60 has
3 been removed by 63. This morning Commissioner
4 McGaffigan complained about things could grow and grow
5 and grow.

6 Now with the two that you have here, it
7 seems that either you are doing a very nice job with
8 what you have, so why bother with to come up with new
9 tools that you want to try and new analysis,
10 specifically tools to do the same thing again and
11 again.

12 I would suggest that you rethink this
13 because this is really maybe you can do what you want
14 to do with what you already have and keep in mind
15 again that the barrier capability performance for
16 individual barriers which was about 60 is now omitted.

17 MR. McCARTEN: I guess -- we hear what the
18 Committee is saying and there's no question we've had
19 a lot of discussions internally and that's the reason.
20 It's clearly stated. These types of analyses are not
21 required to demonstrate compliance with the barrier
22 requirements in 63. But what we're trying to do and
23 for the Committee I think Dick and Sitakanta both
24 tried to give all the things we're looking at with
25 potential to increase our understanding, and I think

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1 the concept of all these things, we're just throwing
2 out where we're going.

3 There is a huge downside to doing barrier
4 neutralization because people jump to those numbers at
5 the bottom and the value is not in those numbers at
6 the bottom. And what we're -- I like to think that
7 when this is done, it's a way to probe and test your
8 own thinking of how the code is working, how you think
9 the system is working and this is just another way to
10 poke at you, your brain a little harder, to think a
11 little more.

12 Ultimately, what you're not seeing and
13 we're not there yet is what kind of information about
14 the system can you pull out and that's the key. And
15 I think this is a way to kind of jiggle the system a
16 little more.

17 Maybe it's not the right way to go, but I
18 think it's a way to push your understanding and I
19 think that's the key we need to get to as you guys are
20 indicating. And we're not there yet. We owe you
21 something more. Where's the understanding in this?
22 And right now, it clearly is not at those bottom
23 numbers. It's deeper than that.

24 MEMBER LEVENSON: Let me ask a question
25 sort of in that relationship. In the one-ON analysis,

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1 the unsaturated zone and the saturated zone have for
2 all practical purposes the same significance, whatever
3 that is. But in the one-OFF analysis the significance
4 is different by more than an order of magnitude. What
5 do we learn from that?

6 MR. McCARTEN: Well, these numbers in
7 order of magnitude, I don't think is necessarily that
8 significant, but the next step is what -- the key is
9 understanding the capability of the barriers and
10 what's going on and why those numbers came out. I
11 think that's what -- you may use this analysis to push
12 you a little harder about the understanding of the
13 capabilities of the barriers, so you clear -- oh yeah,
14 that's why those numbers came out that way, but that's
15 sort of the next step with this and whether this is
16 the right way to go or there's other approaches re
17 better or there's as Dick indicated some intermediate
18 steps or this is the first step, that's where we're at
19 now and we're just trying to as a group, we're always
20 trying to do additional analyses to see if it's
21 helpful.

22 CHAIRMAN HORNBERGER: Tim, you remind me
23 of one of my favorite quotes, the purpose of computing
24 is insight, not numbers. The purpose of computing
25 numbers is not yet in sight.

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1 (Laughter.)

2 I do have a question for Dick, actually.
3 I was really intrigued because I hadn't thought of
4 that before, but DOE's patch model really does lead
5 them to get essentially the same failure rates on
6 every realization.

7 Do you see this as a problem in DOE's
8 code?

9 MR. CODALL: No. I think it's probably
10 somewhat more realistic than what we chose, so that's
11 a point for DOE's conservatism.

12 CHAIRMAN HORNBERGER: But it strikes me
13 that -- I mean, that's equivalent, I think to your
14 saying that all of the uncertainty is aleatory. I
15 hate those terms, but environmental variation and it
16 strikes me that they're probably -- potentially could
17 be another component if for no other reason that you
18 would have differences in fabrication of casks.

19 MR. CODALL: Right, but the problem is
20 that the data don't tell you which is which. And then
21 I think though that the answer that -- is that it
22 doesn't -- it seems that for the ranges of parameters
23 that we're dealing with, the results aren't too
24 different no matter what you assume and that's
25 somewhat reassuring.

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1 CHAIRMAN HORNBERGER: Actually, I had --
2 I've often somewhat facetiously suggested that DOE
3 could build a better safety case by purposefully
4 damaging canisters in a certain pre-determined rate so
5 that they all wouldn't fail at the same time.

6 (Laughter.)

7 MEMBER GARRICK: This reminds of the old
8 days of reliability analysis when they had no
9 failures, someone would assume a failure, a horrible,
10 horrible thing to do.

11 Well, this is very interesting and we'd
12 like to continue. I think that my perspective on this
13 that what you're doing needs to contribute to a couple
14 of things or we would have to challenge it.

15 It's wherewithal. One, a better
16 understanding of the contribution of the individual
17 barriers. And two, a greater confidence that we can
18 build a model that represents reality a little more
19 effectively after doing this work than we could
20 before. And if it doesn't do -- contribute to those
21 things, then I would have to wonder.

22 MR. CODALL: Well, I'd just like to point
23 out this is not part of this presentation, but we're
24 starting development on TPA 5.0 and we're putting back
25 in that code, the diffusion model. It was taken out

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1 earlier. I think I was probably responsible for
2 putting it in and taking it out because it didn't seem
3 to make any difference, but since DOE is depending on
4 that release pathway, we're putting that back in too,
5 so we'll have a handle on it. So there are changes to
6 the code that will improve it.

7 MEMBER GARRICK: I sure wish you'd put
8 something in there that would account for the chemical
9 effects inside the waste package.

10 MR. CODALL: There may be something like
11 that going in. Are you aware of something, Sam?

12 Nothing comes to mind. But where people
13 are chemists here and at the center who worry about
14 such things.

15 MEMBER GARRICK: That's where the action
16 is relative to the mobilization of the waste and the
17 creation of the source term and I think a lot of
18 attention on that would pay high dividends.

19 All right, well, we're running a little
20 behind. We thank you very much.

21 MR. CODALL: Thank you.

22 MEMBER GARRICK: And I think we'll adjourn
23 for a recess.

24 (Off the record.)
25

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CERTIFICATE

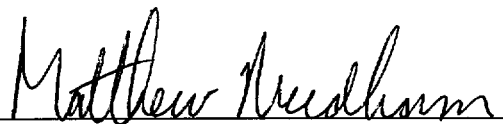
This is to certify that the attached proceedings
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Docket Number: (Not Applicable)

Location: Rockville, Maryland

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Matthew Needham
Official Reporter
Neal R. Gross & Co., Inc.

Sensitivity and Uncertainty in the NRC Total-system Performance Assessment Version 4.1 Code

Sitakanta Mohanty
Center for Nuclear Waste Regulatory Analyses
(210) 522-5185 (smohanty@swri.org)

Richard Codell
U.S. Nuclear Regulatory Commission
(301) 415-8167 (rbc@nrc.gov)

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Briefing to ACNW Members
Rockville, Maryland
March 20, 2002



Outline

- Purpose of Analyses
- Overview of Total-system Performance Assessment Preliminary Results
- Sensitivity Analysis Results
- Effects of Treatment of Data (Variance and Uncertainty)
- Preliminary Risk Insights

Purpose of Analyses

- Gain Risk Insights and Risk-Inform the Program
- Understand the System as a Whole
- Understand the Factors Important to Performance
- Improve Capabilities to Review Potential License Application
 - DOE performance assessment results (e.g., potential risk dilution)
 - Sensitivity and uncertainty analyses approach

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Background

- Total-system Performance Assessment 4.1 Code Uses DOE EDA II Design
- No Analysis of DOE Low Temperature Operating Mode Concept
- Total-system Performance Assessment Code Has More Than 950 Parameters; 330 Sampled
- Alternative Conceptual Models Treated on a Case-by-Case Basis (i.e., alternative models not sampled)
- Notes:
 - Analyses performed mainly for developing staff understanding; they are not mandated by regulation
 - Results are preliminary; input received from other key technical issues could change results

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Performance Assessment Results

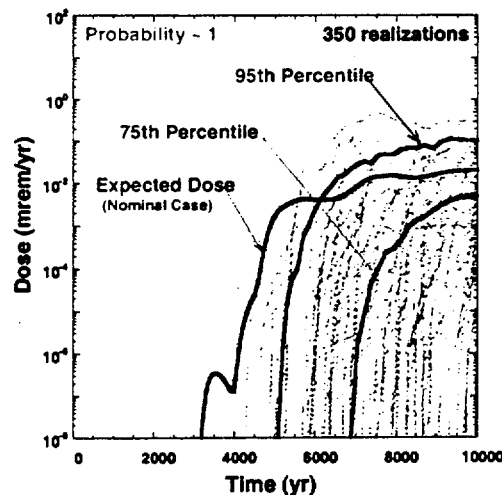
- Performance Measure: Peak Expected Dose to the Reasonably Maximally Exposed Individual
- Results Shown for Two Scenario Classes
 - Nominal scenario
 - Slow degradation over time leading to groundwater release
 - Seismic and climate included
 - Contributions over all realizations averaged at each time step
 - Disruptive event scenario (igneous activity)
 - Short duration, low probability, potential for high consequences
 - Requires special techniques to evaluate risk with reasonable number of runs
 - Convolution of expected doses for all possible events prior to evaluation time

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Nominal Scenario Results

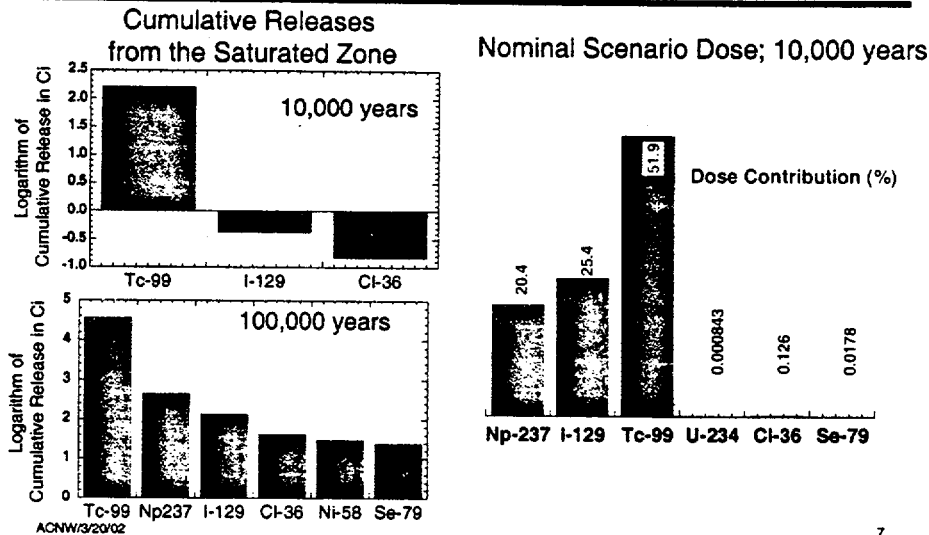
- Initially Defective Failure:
 - Average: 45
 - Range: 1 - 88 [uniform]
- Well Pumping Varied
- 20 km Receptor Location
- No Corrosion or Seismic Failure Within 10,000 years
- Peak Risk 0.021 mrem/years at 9,769 years



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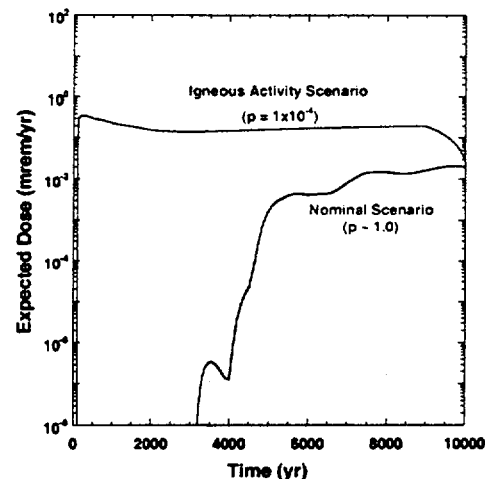
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Nominal Scenario Results (continued)



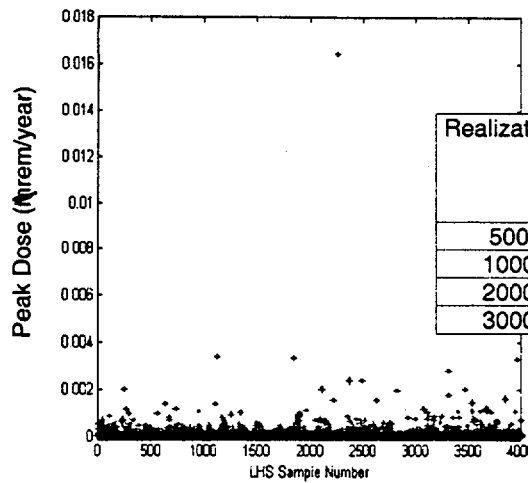
Disruptive (Igneous) Event Scenario Results

- Conditional Dose Computed at 12 Event Times (i.e., 4,200 Realizations)
- Peak Risk 0.35 mrem/years at 245 years
- Early Dose from Short-Lived but High-Activity Nuclides (e.g., ^{241}Am)



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Stability of Peak Expected Dose



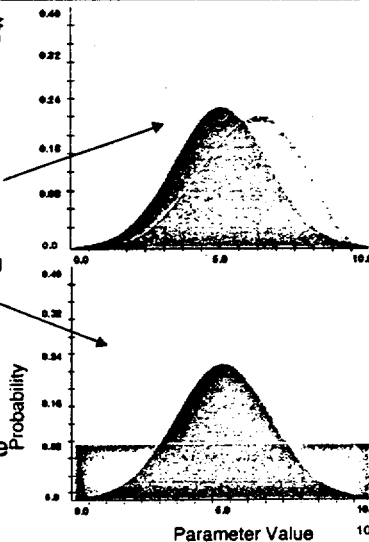
Realizations	Peak Expected Dose (mrem/year)	Peak Occurs at (year)
500	2.48×10^{-2}	10,000
1000	3.05×10^{-2}	8490
2000	3.24×10^{-2}	10,000
3000	2.46×10^{-2}	10,000

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Distributional Sensitivity

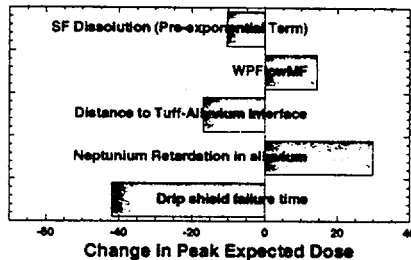
- Relative Impact of the Change of Input Distribution
- Test Cases:
 - Top 10 influential parameters
 - Fixed range, but shift mean by 10% of the range
 - Fixed range, but change distribution type
- Sensitivity Measures
 - Change in peak expected dose
 - Effective distance between nominal case and sensitivity case cumulative distribution functions



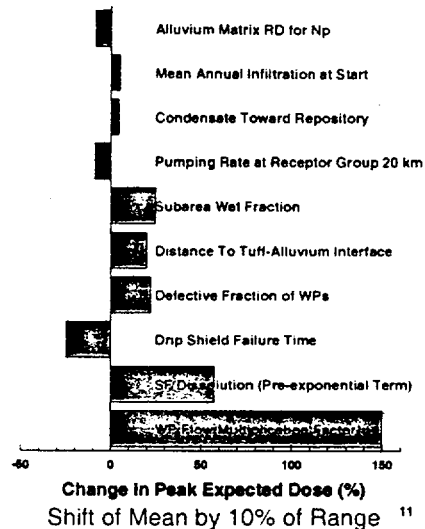
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Distributional Sensitivity (continued)

- Nominal Data Set
- One Parameter Varied at a Time
- Fixed Range

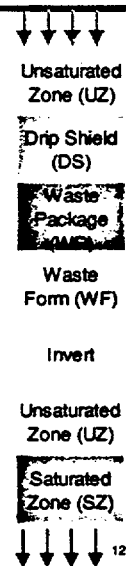


Change of Distribution Type
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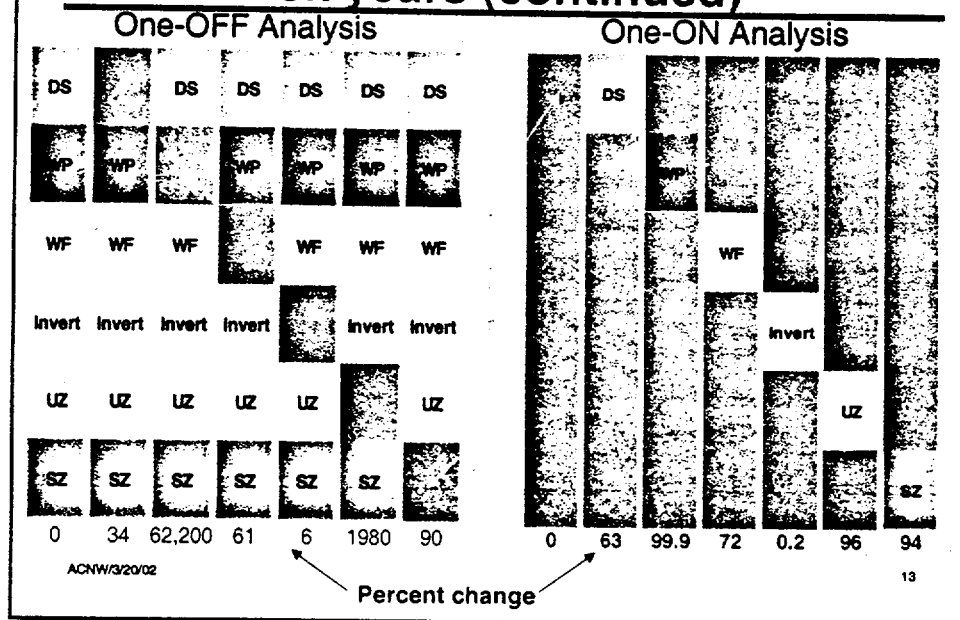
Barrier Component Sensitivity Analyses

- Barrier Divided into Barrier Components
- Sensitivity = Change in Performance Caused by Barrier(s) Suppression With Other Barrier(s) Active, or Barrier(s) Acting Alone With Other Barriers Suppressed
- Results of One-on, One-off, and Cumulative Addition Calculations Provide Different Insights
- Definition of System State for Barrier Suppression Problematic
- Not Required by Regulations

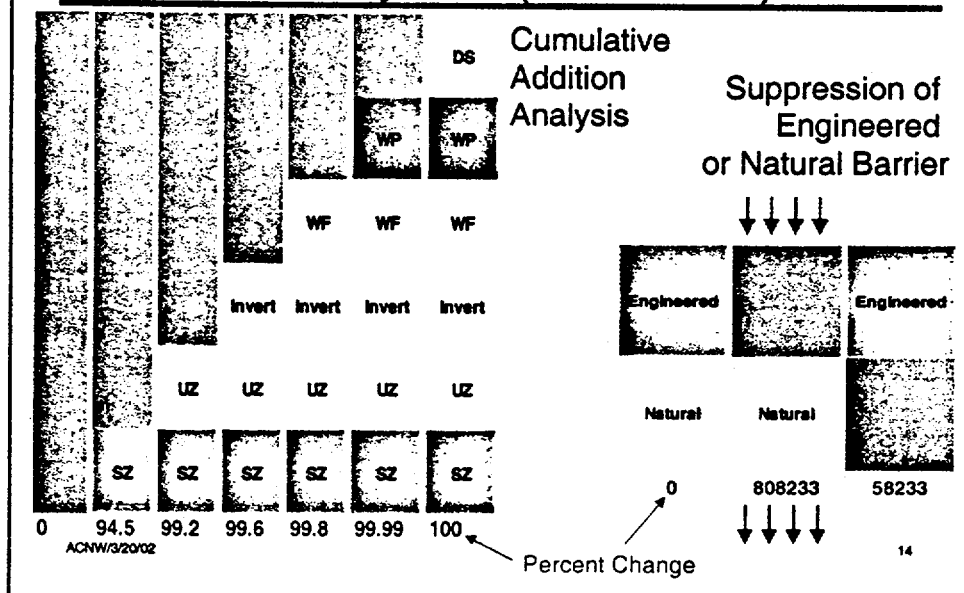


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Barrier Component Sensitivity- 10k years (continued)



Barrier Component Sensitivity- 10k years (continued)



Parameter Sensitivity Analysis

- Nominal Scenario Only
- Determine Sensitivity of Parameters Singly and in Groups
- Statistical Methods Evaluate Sensitivity for Previously Calculated Pool of Vectors
- Nonstatistical Methods Direct How Models Are Sampled to Get Sensitivity
- Results Generally Based on Peak for Each Realization

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Sensitivity Analysis Methods

- Statistical Methods
 - Start with 4000 latin hypercube sampling vectors; 10,000 and 100,000 years
 - Regression of raw and transformed input and output variables
 - Nonparametric tests
 - Parameter-tree method
 - Cumulative distribution functions sensitivity method (developed at the CNWRA)
- NonStatistical Methods
 - Differential analysis
 - Morris method
 - Fourier Amplitude Sensitivity Technique
 - Factorial Design of Experiments

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Composite Statistical Method

- Look at 6 Statistical Tests for 4000 Realizations
- Factor Number of Times Variable Is Statistically Significant for Each Test, and Its Rank
- Develop a Single List of Top Parameters and Their Ranking

Rank	10,000 years	100,000 years
1	Drip shield Failure Time	SF Dissolution Model 2 Pre-exponent
2	Subarea Wet Fraction	Subarea Wet Fraction
3	WP Flow Multiplication Factor	AA_1_1 (corrosion rate parameter)
4	SF Dissolution Model 2 Pre-exponent	Alluvium Matrix RD for Np
5	Defective Fraction Of WPs in a Subarea	Tuff Alluvium Interface Distance
6	Condensate Toward Repository	WP Flow Multiplication Factor
7	Mean Annual Infiltration At Start	Alluvium Matrix RD for Pu
8	Condensate Removed	Mean Annual Infiltration At Start
9	Tuff-Alluvium Interface Distance	Precip. Multiplier At Glacial Max.
10	Well Pumping Rate	Well Pumping Rate

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Top Parameters by Statistical Method

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Factorial Design of Experiments

- Sample Each Parameter at Two Values; 5th and 95th Percentile of Distribution
- Would Require 2^{330} Runs (10^{94} years) for All Possible Combinations
- Fractional Factorial Design More Reasonable, but Ambiguous
- Perform Sampling Iteratively, Using Other Results to Reject Insensitive Parameters
- Advantages
 - Systematic and potentially precise
 - Easily interpreted with Analysis of Variance and trees
 - Reveals interactions among variables
- Disadvantages
 - Can be costly
 - Doesn't cover entire ranges of variables

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Factorial Design Results

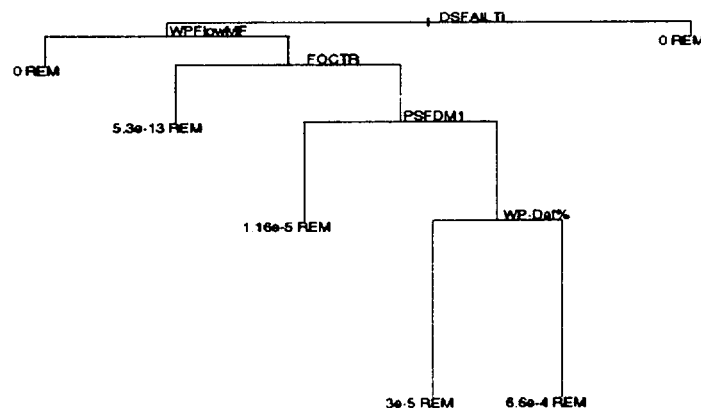
- First Set of 2048 Vectors Identified 100 Potentially Sensitive Variables
- Reduced List to 37 Variables on Basis of Other Information From Runs
- Second Set of 2048 Vectors Identified 10 Sensitive Variables
- Full Factorial ($2^{10} = 1024$ Vectors) Identified 6 to 8 Sensitive Variables

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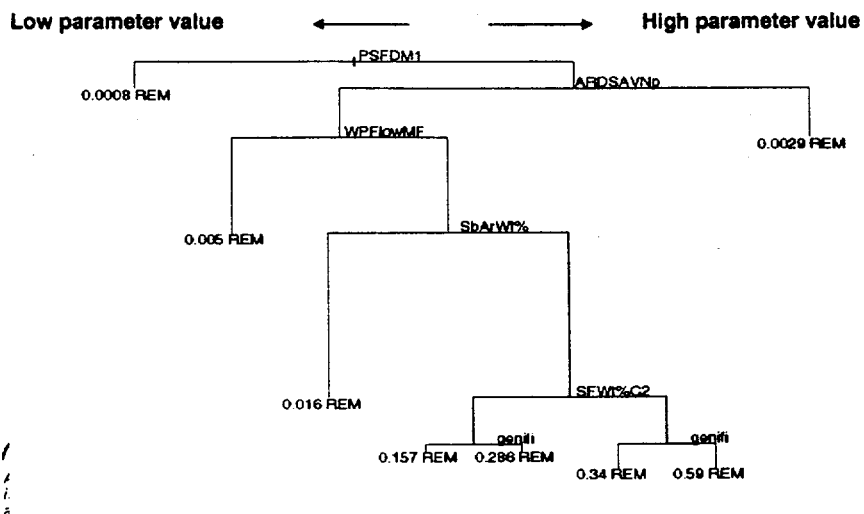
Factorial Design Results- 10k years

Low parameter value ← → High parameter value



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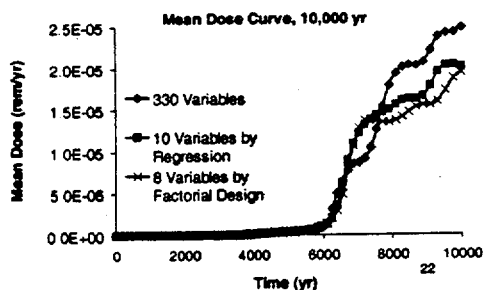
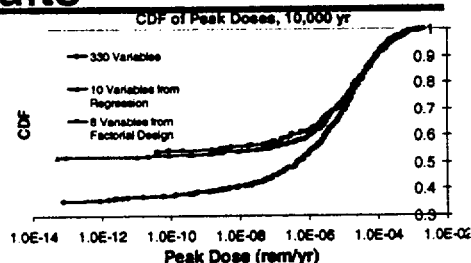
Factorial Design Results-100k years



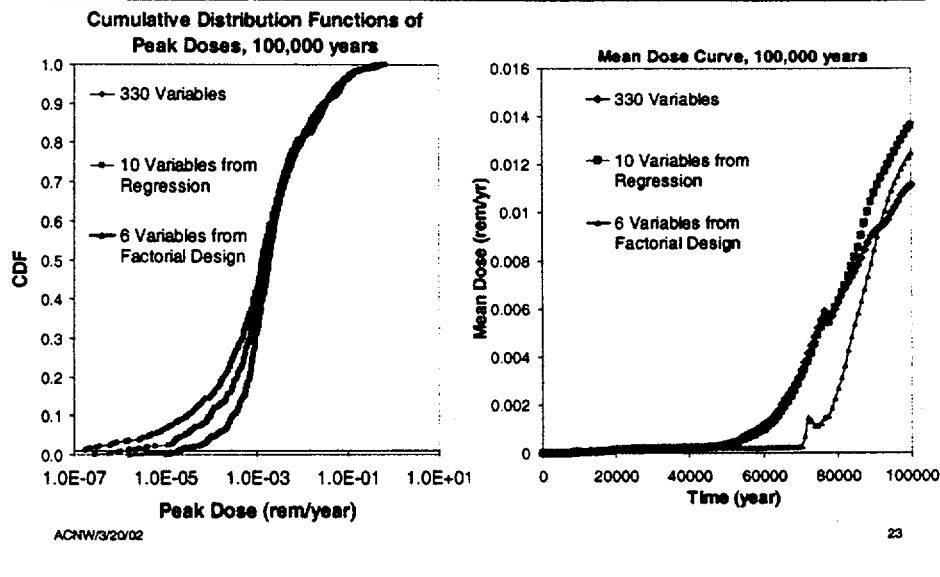
Demonstrating Validity of Sensitivity Results

- Reconstruct Cumulative Distribution Functions of Peak Doses and Mean Dose Curve From Full Set of 330 Variables and Reduced Sets
- 500 Vectors With Reproduced Sampling of Each Variable
- Results Show Most Uncertainty Captured by Reduced Sets of Variables

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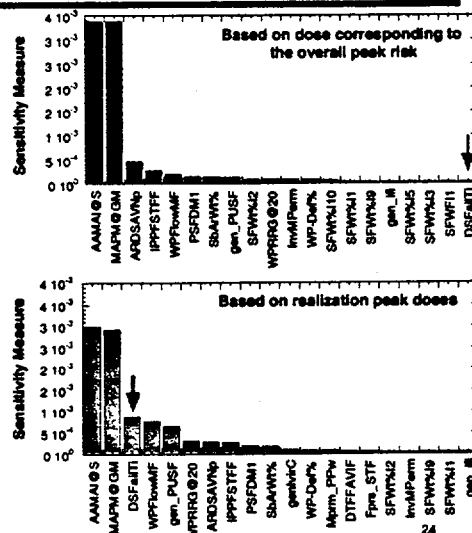


Demonstrating Validity of Sensitivity Results (continued)



Sensitivity Measures

- Two Options:
 - (1) Dose from each realization at time of peak of mean
 - (2) Peak dose from each realization
- Rankings Substantially Different
 - No change in rank only for the top two parameters
 - Four variables from Option 2 do not appear in results from Option 1 and the vice versa
- Option 2 Recommended Because the Sensitivity is Greater (beyond first two variables)



Treatment of Data Variability in Performance Assessment Modeling

- Experimental Data Uncertain Either Because of Lack of Knowledge (Epistemic) or Variability (Aleatory)
- Differences Between Epistemic and Aleatory Uncertainty Often Blurred
- Example: Treatment of Corrosion-rate Data for Waste Packages, and Its Effects on Dose

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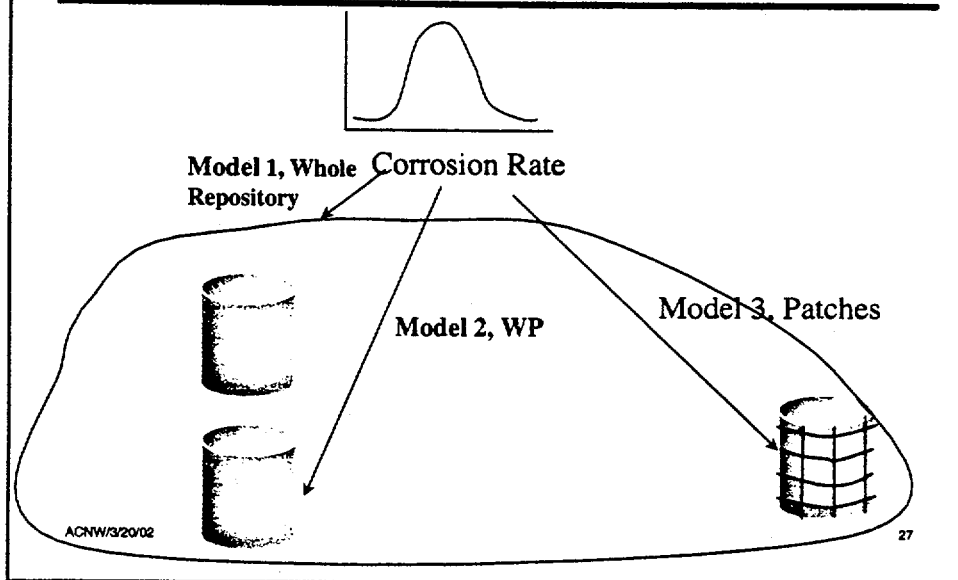
Treatment of Corrosion Rate Data

- NRC Model (TPA 4.1) Represents All Variability by Few "Representative" Waste Packages
- DOE Total System Performance Assessment Uses "Patch Failure" Model That Allows Significant Spatial Variability
- Corrosion Data Could Be:
 - A fixed but uncertain rate (epistemic), or
 - A spatially varying rate due to material and environmental variability (aleatory)

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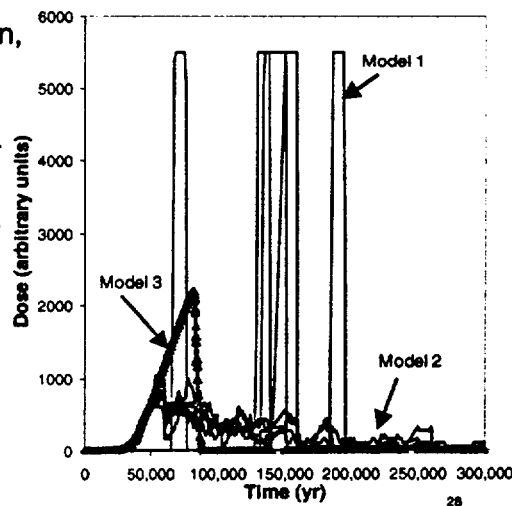
26

Demonstration Models- Three Conceptual Models

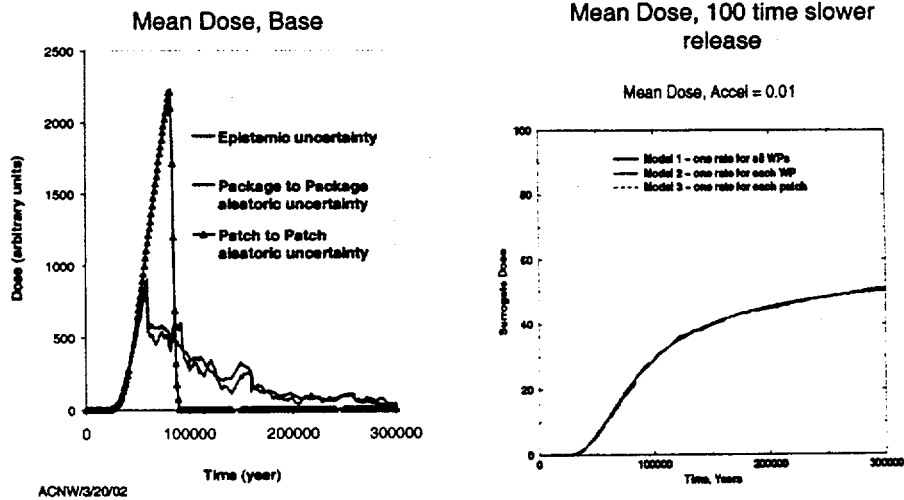


Results of Model Studies

- Doses From Model 1 Have High Peaks Within Realization, but at Widely Different Times
- Model 3 Doses Have Smaller Peaks, but Were Less Different Among Realizations
- Doses Averaged Over All Realizations (Peak-of-mean Approach) Sensitive to Uniformity Among Random Runs

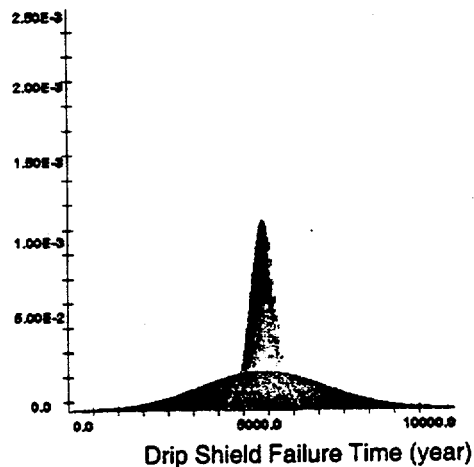


Results of Model Studies (continued)



Risk Dilution

- From Inappropriate Use of Wide Distribution Range
 - May reduce peak mean dose
 - Caused by factors that affect timing of dose
- Example:
 - Drip shield failure times with identical mean, but different ranges
 - Narrow Range: 0.051 mrem/year at 6,410 years
 - Wide range: 0.039 mrem/year at 10,000 years



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Preliminary Insights from Sensitivity Analyses

- **Parametric Sensitivities for 10,000 years**
 - Factors controlling water/fuel contact dominate performance
 - Most dose from low-retardation and long-half-life radionuclides
- **Parametric Sensitivities for 100,000 years**
 - Waste packages usually fail; corrosion parameters not necessarily sensitive
 - Fuel/water contact still important
 - Retardation of neptunium important
- **Barrier Sensitivity**
 - Both natural barrier and engineered barriers make substantial contributions

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Additional Work in Progress

- **Neural Networks for Statistical Sensitivity**
- **New Sensitivity Measures Consistent With Peak Expected Dose**
- **Efficient Distributional Sensitivity Methods**
- **Formal Analysis of Barrier Performance**
 - 29 of 64 analyses completed
 - Tree structure and Analysis of Variance to show dominant sequences of barriers

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Conclusions

- **Parametric Sensitivities Provide Useful Risk Insights**
 - *Ad hoc* sensitivity ranking appears to work well
 - Factorial design shows great promise in clearly defining parameter sensitivity and variable interactions
- **Distributional Sensitivity Appears to be an Effective Approach to Identify Impact of the Choice of Parameter Distribution Shape and Mean**
- **Inappropriate Parameter Range may Lead to Risk Dilution**
- **Treatment of Uncertainty as Lack of Knowledge (Epistemic) or Variability (Aleatory) can Affect Peak Risk Calculated**