

IN THE UNITED STATES DISTRICT COURT  
FOR THE MIDDLE DISTRICT OF PENNSYLVANIA

TAMMY KITZMILLER, et al :  
 : CASE NO.  
 v. : 4:04-CR-002688  
 :  
 DOVER AREA SCHOOL DISTRICT, :  
 et al : :

TRANSCRIPT OF PROCEEDINGS  
BENCH TRIAL

**MORNING SESSION**

BEFORE: HON. JOHN E. JONES, III  
  
DATE : October 19, 2005  
8:55 a.m.  
  
PLACE : Courtroom No. 2, 9th Floor  
Federal Building  
Harrisburg, Pennsylvania  
  
BY : Wendy C. Yinger, RPR  
U.S. Official Court Reporter

APPEARANCES:

ERIC J. ROTHSCHILD, ESQUIRE  
WITOLD J. WALCZAK, ESQUIRE  
STEPHEN G. HARVEY, ESQUIRE  
RICHARD B. KATSKEE, ESQUIRE  
THOMAS SCHMIDT, ESQUIRE  
For the Plaintiffs

PATRICK T. GILLEN, ESQUIRE  
RICHARD THOMPSON, ESQUIRE  
ROBERT J. MUISE, ESQUIRE  
For the Defendants

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I N D E X T O W I T N E S S E S

FOR THE DEFENDANTS      DIRECT      CROSS      REDIRECT      RECROSS

Michael Behe

By Mr. Rothschild

12

1                   (Whereupon, the following discussion was  
2                   held in chambers:)

3                   THE COURT: All right. What are -- we have  
4 an issue?

5                   MR. SCHMIDT: Your Honor, we wanted to alert  
6 the Court before we used it in cross examination of a  
7 document that we plan to use that Your Honor may regard  
8 as covered by the confidentiality order having to do  
9 with the draft of the successor to Pandas. It's a page  
10 out of that draft.

11                   It's the page that's analogous to the old  
12 page 25 that dealt with sudden -- intelligent design as  
13 it holds the various forms of life began with  
14 distinctive features already intact.

15                   THE COURT: Is this the latest version --

16                   MR. SCHMIDT: This is the --

17                   THE COURT: As yet unpublished --

18                   MR. SCHMIDT: Correct.

19                   THE COURT: -- of Pandas. And you'll have  
20 to refresh my recollection. I didn't have a chance,  
21 after Liz alerted me, to look in the file, but did we  
22 have a confidentiality order in the midst of determining  
23 FTE's motion. Is that what it was for? You'll have to  
24 help me out, because I don't recall.

25                   MR. SCHMIDT: It originally came up because

1 we subpoenaed it from William Dembski --

2 THE COURT: I recall that.

3 MR. SCHMIDT: -- who was the author. And  
4 FTE participated in that.

5 THE COURT: I recall that it was subpoenaed.  
6 I recall that FTE moved to block --

7 MR. SCHMIDT: For a protective order.

8 THE COURT: -- the subpoena. And, of  
9 course, I know that, we all know that Mr. Dembski is not  
10 testifying, and we all know that FTE was not permitted  
11 to intervene. What I don't remember is sequentially  
12 when the protective order came to being in exactly -- I  
13 understand why it came into play, but apparently it was  
14 not self-extinguishing as it related to the litigation.  
15 Is that a fair statement?

16 MR. SCHMIDT: Yes. In fact, it had a  
17 provision in it that said it would continue past the  
18 trial even until publication of the text.

19 THE COURT: So why do you think you're  
20 entitled to open it up?

21 MR. SCHMIDT: Because nothing in the  
22 protective order says that we couldn't use it. It said,  
23 if we did use it, it would be under seal, preserving the  
24 confidentiality of it.

25 So if there is reference to this, as there

1 will be, I wanted the Court to know that we intended to  
2 do that, so that the courtroom could be cleared, and so  
3 that this part of the record could be under seal to the  
4 extent that it's quoting from it.

5 MR. ROTHSCCHILD: Your Honor, I would just  
6 add that, I would actually interpret the protective  
7 order a little more liberally. It certainly doesn't  
8 allow us to publish this widely, and it required any  
9 filings with briefs to be under seal, and any  
10 depositions that they used it as an exhibit to be under  
11 seal.

12 I think this is why we're alerting to you,  
13 that it does not necessarily mean that once we're in  
14 public trial, that it would preclude its use in public,  
15 but we're also amenable to it being done with a closed  
16 courtroom, if that's --

17 THE COURT: Well, do we have -- was a  
18 protective order entered -- and again, you'll have to  
19 refresh my recollection -- pursuant to a stipulation?

20 MR. SCHMIDT: Yes, it was.

21 THE COURT: And the stipulation, who were  
22 the parties to the stipulation? Was FTE a party?

23 MR. SCHMIDT: The Plaintiffs and FTE.

24 MR. GILLEN: Well, actually, weren't we,  
25 Chuck, as well?

1 MR. SCHMIDT: You were as well.

2 MR. GILLEN: Yeah, we were as well.

3 MR. WALCZAK: Your Honor, I think Eric's  
4 interpretation that this may not apply if it's being  
5 used in open court was largely validated when we had  
6 that hearing on FTE's intervention motion.

7 THE COURT: Does somebody have the  
8 stipulation?

9 LAW CLERK: I can get it.

10 THE COURT: Why don't you pull it off.

11 MR. WALCZAK: And while I don't believe we  
12 used design of life there, the other documents had been  
13 produced under seal including, I believe, and Chuck will  
14 correct me if I'm wrong, the FTE, some of the FTE  
15 statements and writings that they had. And some of  
16 those were introduced in court, put into the record.

17 FTE was there, and they had no objection,  
18 and did not seem to differ from our understanding of the  
19 protective order as not extending to things that  
20 happened in open court.

21 THE COURT: Are you seeking to actually  
22 admit a document in -- you're shaking your head no.  
23 You're going to simply question from the text of the  
24 manuscript?

25 MR. SCHMIDT: And read it to them, yes.

1 THE COURT: What's your position?

2 MR. GILLEN: A couple things. Actually, I'm  
3 grateful to you guys for bringing it to my attention.  
4 My recollection is that, it did cover litigation, that  
5 there was some discussion of that. I think what they're  
6 suggesting though, a short passage, so it can be kept  
7 confidential, does what I thought you had in mind,  
8 Judge, which is to protect their property interests.  
9 And I can see that being a way to get rid of the  
10 problems, so to speak.

11 THE COURT: Well, my recollection is that,  
12 FTE's concern was that they obviously had an  
13 intellectual property interest, and they were concerned  
14 that a wholesale release of the manuscript would subject  
15 it to pre-publication criticism, if I recall, that Mr.  
16 Buell was particularly, and justifiably, I thought,  
17 alarmed about.

18 I really wonder, under the circumstances, if  
19 it's a short passage, how much that's going to interfere  
20 with the intellectual property rights. I suppose you  
21 could argue that, that would allow focus and criticism  
22 of that particular passage, but I'm not so sure that  
23 that's really what his concern was. I thought his  
24 concern was a wholesale release of the entire  
25 manuscript, which is really what was threatened when Mr.

1 Dembski testified.

2 MR. GILLEN: And, Judge, I don't represent  
3 FTE.

4 THE COURT: I understand that.

5 MR. GILLEN: So I can't speak.

6 THE COURT: But as a signatory to the  
7 stipulation, I suppose you have Atillaed the hunt.

8 MR. GILLEN: We had an expert at that time  
9 who asked me to move to protect the intellectual  
10 property right because of his fiduciary duty. I made  
11 that motion, and I want -- I do want to preserve what I  
12 can by way of protection of their work product rights.

13 MR. SCHMIDT: As the draftsman of the  
14 stipulation, I must say that I had in mind the far  
15 broader text. The concern that was expressed was that  
16 this would give the NCSE's people, Scott and others, an  
17 opportunity to poison the well before publication.

18 THE COURT: Let me see the passages.

19 MR. SCHMIDT: It's the second paragraph.

20 MR. ROTHSCHILD: We would probably use one  
21 other page to just correlate some other charts.

22 MR. SCHMIDT: In my own mind, I see this as  
23 kind of analogous to the fair use exception and  
24 copyright law. You can take a snippet and use it  
25 without harming the copyright interests.



1 MR. ROTHSCHILD: I do think there's one  
2 other consideration, Judge, for your --

3 THE COURT: Go ahead.

4 MR. ROTHSCHILD: That it may -- the FTE has  
5 counsel in this area, and it may make sense, before  
6 using it, to alert them. I mean, we do intend to use it  
7 today for purposes of impeachment with Professor Behe.

8 THE COURT: That's exactly what I was going  
9 to suggest. Who's counsel?

10 MR. ROTHSCHILD: Leonard Brown, that group.

11 THE COURT: Yeah. Why don't you do this.  
12 Why don't you take time now to, before we get started,  
13 you know, we've been moving at a pretty good pace, and  
14 we haven't had these things happen, and they do happen  
15 in trials. So why don't you take some time and contact  
16 FTE's counsel. I think you want to do it for your own  
17 protection.

18 Obviously, once I rule, I suppose that  
19 you're protected, but you entered into a stipulation,  
20 and I would have some concern --

21 MR. ROTHSCHILD: I think it's just fair.

22 THE COURT: -- about that, and I think you  
23 want to at least give them notice. If we have to  
24 reconvene and get them on at least a conference call and  
25 let them be heard, and that might be better than having

1 you, you know --

2 MR. GILLEN: Me speak for them.

3 THE COURT: Sure. That puts you in a  
4 difficult position. You're signatory as parties, but  
5 you really don't want to put yourself in the position to  
6 speaking for FTE. And then we can hear out FTE. I'm  
7 not sure, you know, given this brief passage, that it  
8 violates the sense of the stipulation to allow  
9 questioning, even in open court.

10 I'm somewhat reluctant to clear the  
11 courtroom for these brief passages because, again, I'll  
12 read the stipulation and the order because they're not  
13 -- I don't recall them instantly. But I thought the  
14 thrust, and you seem to agree with this, is that the  
15 manuscript, as a whole, would be protected. And I  
16 understand. I think we all understood the purpose for  
17 that at the time.

18 MR. ROTHSCHILD: Your Honor, should we  
19 suggest a time -- I mean, do you want to do that at a  
20 lunch break or find out --

21 THE COURT: How much more cross do you have?

22 MR. ROTHSCHILD: It will be inversely  
23 proportional to mentions of the Big Bang, I think.

24 THE COURT: So you're going to go all day.

25 MR. ROTHSCHILD: It could be quite a while.

1           THE COURT: All right. Well, why don't you  
2 get started. Take some time now. Why don't you contact  
3 them. Why don't you see what their availability is. I  
4 mean, I recognize we're catching them flatfooted. See  
5 if they've got somebody that they can get on the phone,  
6 you know, as soon as possible. I just as soon get  
7 started.

8           If you give me a time later this morning,  
9 we'll just recess. If they say, you know, we're  
10 available at 11, or whatever the case may be, then we  
11 can at least get started; 10:30, 11. I'm not suggesting  
12 a time. Just find a time or we can do it as we break  
13 for lunch, if that is more convenient for them. Hard to  
14 believe they wouldn't have somebody that they could get  
15 at some point involved in a phone conversation.

16           Then you can reserve your cross on this  
17 issue until we hear them out at that point. Now if they  
18 tell you they don't care, which I'd be surprised, but if  
19 they tell you that, then we'll take that up at that  
20 time. I suppose they're going to have to likely contact  
21 FTE and find out what.

22           MR. GILLEN: That's what I can foresee. By  
23 the time they get in touch with FTE which, I think, is  
24 in Texas. You guys know better than I do.

25           THE COURT: And there's a time delay.

1 MR. SCHMIDT: One hour.

2 MR. GILLEN: It's just one hour, but Mr.  
3 Buell is rather difficult to reach.

4 MR. SCHMIDT: When he chooses.

5 THE COURT: Well, you know, if they can't  
6 reach him, I'll rule, if I have to, in the absence of  
7 that. But I think at least fair notice to their  
8 counsel, if they can connect with the mothership, and  
9 we'll take it up at that time.

10 (Whereupon, the discussion held in chambers  
11 concluded at 9:05 a.m. and proceedings  
12 reconvened in open court at 9:18 a.m.)

13 THE COURT: All right. Good morning to all.  
14 I apologize for the somewhat late start. We had a  
15 slight issue that we had to handle in chambers with  
16 counsel. And that rapidly resolved, so that we can  
17 commence this morning's session. We will do so. We  
18 will continue cross examination of the witness by Mr.  
19 Rothschild.

20 (Whereupon, MICHAEL BEHE, Ph.D., resumed the  
21 stand, and testimony continued.)

22 **CROSS EXAMINATION (CONTINUED)**

23 BY MR. ROTHSCHILD:

24 Q. Good morning, Professor Behe.

25 A. Good morning, Mr. Rothschild.

1 Q. How are you?

2 A. Fine, thanks.

3 Q. After the Court adjourned yesterday, did you talk  
4 to anybody about your testimony?

5 A. I did not.

6 Q. I'm going to see if we can reach an agreement on  
7 something here. You agree that this is a case about  
8 biology curriculum?

9 A. Yes, I do.

10 Q. Not about physics, a physics curriculum?

11 A. It's not about a physics curriculum, but from my  
12 understanding, many issues that are being discussed here  
13 are particularly relevant to other issues that have come  
14 up in other disciplines of science.

15 Q. This is a case about what's being taught in  
16 biology class not physics class?

17 A. As I said, I agree that it is, but one more time,  
18 I think many things in the history of science are  
19 relevant to this, and they've happened in other  
20 disciplines as well.

21 Q. You've already testified you're not an expert in  
22 physics or astrophysics?

23 A. That's correct.

24 Q. And you might not know this about me, but I'm not  
25 either.

1           A. I'm surprised.

2           Q. So I'm going to propose an agreement. I won't  
3 ask you any questions about the Big Bang, and you won't  
4 answer any questions about the Big Bang. Can we agree  
5 to that, Professor Behe?

6                   MR. MUISE: Objection, Your Honor. He's  
7 trying to limit the testimony of the witness by some  
8 sort of agreement. He's obviously testified and  
9 explained why the relationship of the Big Bang is so  
10 important. He just answered his questions to try to  
11 proffer some prior agreement to the witness that he  
12 can't reference factors of prior testimony in cross  
13 examination. That just seems inappropriate, Your Honor.

14                   THE COURT: What's your answer?

15                   THE WITNESS: No. , I think references to  
16 the Big Bang are extremely appropriate to making clear  
17 why I think these -- making clear my views on these  
18 issues.

19 BY MR. ROTHSCHILD:

20           Q. Fair to say, Professor --

21                   THE COURT: There you go, Mr. Muise.

22 BY MR. ROTHSCHILD:

23           Q. Fair to say, Professor Behe, that over the last  
24 two days of testimony, you've told us everything you  
25 know about the Big Bang that's relevant to the issue of

1 intelligent design and biology?

2 A. Well, I'm not sure. I would have to reserve  
3 judgment.

4 Q. You might have some more?

5 A. Perhaps.

6 Q. Let the record state, I tried.

7 MR. ROTHSCHILD: May I approach the witness,  
8 Your Honor?

9 THE COURT: You may.

10 BY MR. ROTHSCHILD:

11 Q. Professor Behe, I've showed you what we marked as  
12 Plaintiffs' Exhibit 726, and that's an article that was  
13 published in Christianity Today?

14 A. That is correct, yes.

15 Q. It's titled Tulips and Dandelions?

16 A. Yes.

17 Q. And it actually indicates that there was a  
18 debate, and there's actually a back and forth between  
19 you and another writer named Rebecca, I'm sure I'll  
20 butcher this, but Fleastra (phonetic)?

21 A. Fleastra (phonetic). She's a professor of  
22 biology (inaudible) College in California, yes, that's  
23 correct.

24 Q. This is an article you wrote on or about  
25 September or October 1998?

1           A.    Yes, that's correct.

2           Q.    And if you could turn to the second -- this is an  
3 argument that discusses intelligent design?

4           A.    I think it does, but to be perfectly honest, I  
5 have not read this article since it was published seven  
6 years ago.  So I am not entirely clear exactly what I  
7 said in here.  But it certainly is likely to do so.

8           Q.    Do you need to review it for a moment to confirm  
9 that?

10          A.    That would be great.  Thank you.

11                   THE COURT:  Take all the time you need to  
12 read it.

13                   THE WITNESS:  Thank you.  Yes, thank you.  
14 Yes, that's correct.

15 BY MR. ROTHSCHILD:

16          Q.    Matt, could you turn to the second page of this  
17 document?  And Professor Behe, if you would flip to that  
18 page as well.  It will be on your screen as well.  And,  
19 Matt, if you could highlight the question on the bottom  
20 left-hand column, the last paragraph beginning with the  
21 word, what.  And you asked the question in this article,  
22 what does this all mean for a Christian, correct?

23          A.    Yes.

24          Q.    And you said, On the one hand, not much, right?

25          A.    That's correct.



1       Q.   And, Matt, if you could go to the second column,  
2   and the second full paragraph, second full paragraph --  
3   next paragraph. Thank you. Actually highlight those  
4   two. You say, On the other hand, scientific evidence of  
5   design means a lot for Christians for a couple of  
6   reasons. Correct? That's what you wrote?

7       A.   That's correct, yes.

8       Q.   Going down to the next paragraph, one of the  
9   reasons you give is, Christians live in the world with  
10   non-Christians. We want to share the Good News with  
11   those who have not yet grasped it and to defend the  
12   faith against attacks.

13               Materialism is both a weapon that many  
14   antagonists use against Christianity and a stumbling  
15   block to some who would otherwise enter the church. To  
16   the extent that the credibility of materialism is  
17   blunted, the task of showing the reasonableness of the  
18   faith is made easier, although Christianity can live  
19   with a world where physical evidence of God's action is  
20   hard to discern, materialism has a tough time with a  
21   universe that reeks of design. That's what you wrote,  
22   correct?

23       A.   Yes, that's exactly what I wrote.

24       Q.   And that concept of materialism, that's actually  
25   also mentioned in the section on the Wedge strategy that

1 we looked at yesterday, correct?

2 A. I think so, yes.

3 Q. And when you refer to the Good News there, that  
4 was not just the Yankees winning the world series around  
5 this time, correct?

6 A. That's correct. No, that is intended to mean the  
7 Christian gospel. So here, I was explaining, and I was  
8 speaking as a Christian in a magazine that is a  
9 Christian publication. And assuming the assumptions  
10 that Christians have from non-scientific -- from  
11 non-scientific areas, that is historical, theological,  
12 and philosophical principles, why I think, how I think  
13 this impacts Christian concerns.

14 And I emphasize that first paragraph that you  
15 read from, What does all this mean for a Christian? On  
16 the one hand, not much. The faith of Christians rests  
17 on the historical reality of events recorded in the  
18 gospels rather than on the next theory coming out of the  
19 laboratory.

20 By definition, Christians already believe in  
21 design because they believe in a designer. So by that  
22 -- I'm sorry. But just let me make one more point. So  
23 by that paragraph, I was trying to say that, in fact,  
24 design, apparent design in the world is not necessary  
25 for Christian belief.

1       Q. On one hand, it's not -- it doesn't mean a lot.  
2 On the other hand, it means quite a bit?

3       A. On the one hand, it's not necessary. But on the  
4 other hand, it can offer support to a Christian world  
5 view. And if I might refer back to the Big Bang, the  
6 Big Bang was taken by a number of people as evidence for  
7 a theological world view, and Christians have used that  
8 to argue for the plausibility of Christian views.

9               Nonetheless, simply because the Big Bang is  
10 compatible with Christianity, and because it makes some  
11 theistic views seem more plausible, that does not mean  
12 that the Big Bang itself is not a scientific theory.

13               And in the same sense, just because intelligent  
14 design is compatible with Christian views, or because it  
15 makes such views or other theistic views seem more  
16 plausible does not mean that intelligent design itself  
17 is not a scientific theory.

18       Q. I'd like to return to Darwin's Black Box. And  
19 that is where you're making your scientific argument,  
20 correct, Professor Behe?

21       A. That's correct.

22       Q. If you could turn to page 185 of that book. I'd  
23 actually like you to read -- we'll take turns here --  
24 from the last paragraph on 185 beginning, molecular  
25 evolution, and go to the end of the chapter, which is

1 one more paragraph.

2       A. Molecular evolution is not based on scientific  
3 authority. There is no publication in the scientific  
4 literature, in prestigious journals, specialty journals,  
5 or books that describes how molecular evolution of any  
6 real, complex, biochemical system either did occur or  
7 even might have occurred.

8               There are assertions that such evolution  
9 occurred, but absolutely none are supported by pertinent  
10 experiments or calculations. Since no one knows  
11 molecular evolution by direct experience, and since  
12 there is no authority on which to base claims of  
13 knowledge, it can truly be said that, like the  
14 contention that the Eagles will win the Super Bowl this  
15 year, the assertion of Darwinian molecular evolution is  
16 merely bluster.

17               Publish or perish is a proverb that academicians  
18 take seriously. If you do not publish your work for the  
19 rest of the community to evaluate, then you have no  
20 business in academia. And if you don't already have  
21 tenure, you will be banished.

22               But the saying can be applied to theories as  
23 well. If a theory claims to be able to explain some  
24 phenomenon, but does not generate even an attempt at an  
25 explanation, then it should be banished. Despite

1 comparing sequences and mathematical modeling, molecular  
2 evolution has never addressed the question of how  
3 complex structures came to be.

4 In effect, the theory of Darwinian molecular  
5 evolution, has not published, and so it should perish.

6 Q. That was your view in 1996?

7 A. Yes, that's correct.

8 Q. That is still your view today?

9 A. Yes, it is. And if I may elaborate on that?

10 Q. Professor Behe, the answer was yes?

11 A. Well, I want to tell you what my view was.

12 Q. Professor Behe, you understand that your counsel  
13 will have an opportunity to ask follow-up questions  
14 after I'm done with my cross examination?

15 A. Is that correct?

16 Q. That is. Unless the judge rules otherwise, he  
17 will have that chance, so the answer to my question is  
18 yes? That's still your view today?

19 MR. MUISE: Dr. Behe is trying to completely  
20 answer his question. And counsel is attempting to  
21 prevent him from doing so.

22 THE COURT: Well, he's asking him a yes/no  
23 question.

24 MR. MUISE: I don't think it's a question  
25 that can be answered yes no. He has built in assertions

1 that can't just be answered yes or no.

2 THE COURT: If he says he can't answer it  
3 yes or no, then Mr. Rothschild is stuck with that  
4 answer. So you can answer the question as you see fit.

5 THE WITNESS: No, that's not a completely  
6 accurate view.

7 BY MR. ROTHSCHILD:

8 Q. What's changed, Professor Behe?

9 A. That does not go into sufficient detail to  
10 describe my view.

11 Q. I hesitate to ask whether this will involve the  
12 Big Bang, but give us a little more detail.

13 A. The detail is actually simply this, that by these  
14 publications, I mean detailed rigorous accounts for  
15 complex molecular machines, not just either hypothetical  
16 accounts or sequence comparisons or such things.

17 Q. And so with that qualification, that is your  
18 view?

19 A. Yes.

20 Q. Now you have never argued for intelligent design  
21 in a peer reviewed scientific journal, correct?

22 A. No, I argued for it in my book.

23 Q. Not in a peer reviewed scientific journal?

24 A. That's correct.

25 Q. And, in fact, there are no peer reviewed articles

1 by anyone advocating for intelligent design supported by  
2 pertinent experiments or calculations which provide  
3 detailed rigorous accounts of how intelligent design of  
4 any biological system occurred, is that correct?

5 A. That is correct, yes.

6 Q. And it is, in fact, the case that in Darwin's  
7 Black Box, you didn't report any new data or original  
8 research?

9 A. I did not do so, but I did generate an attempt at  
10 an explanation.

11 Q. Now you have written for peer reviewed scientific  
12 journals on subjects other than intelligent design,  
13 correct?

14 A. Yes.

15 Q. And in those articles, you did report original  
16 research and data, at least in many of them, correct?

17 A. Yes.

18 Q. You would agree that there are some journals that  
19 are more difficult than others to get one's research  
20 published in?

21 A. Yes, that's correct.

22 Q. Proceedings of the National Academy of Science?

23 A. Yes.

24 Q. Nature?

25 A. That's correct.

1 Q. Science?

2 A. Yes.

3 Q. Journal of Molecular Biology?

4 A. That's easier than the other ones, but, yes.

5 Q. Still pretty good?

6 A. Yeah. I would take it, sure.

7 Q. In fact, you have taken that for some of these  
8 publications in your non-intelligent design work?

9 A. That's correct.

10 Q. And you've also served as a peer reviewer,  
11 correct?

12 A. Yes.

13 Q. And when you do that, you get a submission from a  
14 scientist, correct? You receive the submission from the  
15 editor?

16 A. From the editor, yes.

17 Q. And you review those submissions carefully?

18 A. Yes, I do.

19 Q. There are some sort of professional expectations  
20 about how peer reviewers do their task?

21 A. Yes, you're supposed to read the manuscripts  
22 carefully and see if you can make suggestions and  
23 criticisms.

24 Q. You look at the experimental results?

25 A. Sure.



1       Q.   You look -- you try to make a determination  
2 whether the techniques were proper?

3       A.   That's correct.

4       Q.   Try to make an assessment about whether  
5 conclusions follow from the data?

6       A.   That's correct.

7       Q.   You analyze whether there are gaps and problems  
8 in the experiment?

9       A.   Yes, that's right.

10      Q.   And on occasions, you've communicated false in  
11 articles that you were peer reviewing, correct?

12      A.   That's correct.

13      Q.   That's happened to you as well?

14      A.   Sure.

15      Q.   All part of the scientific process, right?

16      A.   Yes, that's correct.

17      Q.   Okay. Now you stated on Monday that Darwin's  
18 Black Box was also peer reviewed, right?

19      A.   That's correct.

20      Q.   You would agree that peer review for a book  
21 published in the Trade Press is not as rigorous as the  
22 peer review process for the leading scientific journals,  
23 would you?

24      A.   No, I would not agree with that. The review  
25 process that the book went through is analogous to peer

1 review in the literature, because the manuscript was  
2 sent out to scientists for their careful reading.

3 Furthermore, the book was sent out to more  
4 scientists than typically review a manuscript. In the  
5 typical case, a manuscript that's going to -- that is  
6 submitted for a publication in a scientific journal is  
7 reviewed just by two reviewers. My book was sent out to  
8 five reviewers.

9 Furthermore, they read it more carefully than  
10 most scientists read typical manuscripts that they get  
11 to review because they realized that this was a  
12 controversial topic. So I think, in fact, my book  
13 received much more scrutiny and much more review before  
14 publication than the great majority of scientific  
15 journal articles.

16 Q. Now you selected some of your peer reviewers?

17 A. No, I did not. I gave my editor at the Free  
18 Press suggested names, and he contacted them. Some of  
19 them agreed to review. Some did not.

20 Q. And one of the peer reviewers you mentioned  
21 yesterday was a gentleman named Michael Atchison?

22 A. Yes, I think that's correct.

23 Q. I think you described him as a biochemist at the  
24 Veterinary School at the University of Pennsylvania?

25 A. I believe so, yes.

1 Q. He was not one of the names you suggested,  
2 correct?

3 A. That is correct.

4 Q. In fact, he was selected because he was an  
5 instructor of your editor's wife?

6 A. That's correct. My editor knew one biochemistry  
7 professor, so he asked, through his wife, and so he  
8 asked him to take a look at it as well.

9 Q. And you found out his name later, correct?

10 A. That's right, yes.

11 Q. From your editor?

12 A. No. I think actually Professor Atchison himself  
13 contacted me later after the book came out.

14 MR. ROTHSCHILD: May I approach the witness?

15 THE COURT: You may.

16 BY MR. ROTHSCHILD:

17 Q. Professor Behe, I've shown you an exhibit marked  
18 P-754, and that's an article titled -- or a writing  
19 titled Mustard Seeds by Dr. Michael Atchison?

20 A. Yes.

21 Q. That is a picture of him, correct?

22 A. I think so. I haven't seen him in a few years.

23 Q. It certainly identifies him as the head of  
24 biochemistry in the department of animal biology at the  
25 University of Pennsylvania?

1           A.    Yes, he's the department chair in the vet school.

2           Q.    Professor Behe, I'd like you to look at the first  
3 -- I'm sorry, the last paragraph on the first page, and  
4 I'm going to read this for the record. This is what  
5 Professor Atchison wrote. While I was identifying  
6 myself as a Christian --

7                   MR. MUISE:  Objection, Your Honor. This is  
8 hearsay, and there's been no foundation he even knows  
9 this thing exists. He's reading into the record a  
10 document that he apparently got from somewhere that we  
11 don't have any foundation for. What he's reading into  
12 the record is absolutely hearsay.

13                   MR. ROTHSCHILD:  I'm not proposing to  
14 introduce this into evidence at this point, although  
15 I'll reserve that right. But this is for purposes of  
16 impeachment. I think it's highly relevant.

17                   MR. MUISE:  He hasn't even shown Dr. Behe  
18 even knows anything about this article or where it's  
19 from or any basis for it.

20                   MR. ROTHSCHILD:  I'm going to ask him about  
21 the facts that are stated in this article.

22                   THE COURT:  Why isn't it fair for  
23 impeachment purposes?

24                   MR. MUISE:  It's -- again, Your Honor, I  
25 guess you have to see how this is going to go. I was

1 objecting because he's going to read into the record a  
2 portion of this document that he hasn't even established  
3 that Dr. Behe has any knowledge about.

4 THE COURT: Well, it's not a transcript.

5 MR. MUISE: That's true. It's a document  
6 that was produced out of court.

7 THE COURT: I understand. But to read it  
8 into the record, as you might not with a transcript,  
9 that's not reason alone to not permit it in the  
10 proceedings. I think, given the witness's answer, it's  
11 fair impeachment. Now --

12 MR. MUISE: I mean, impeachment in what  
13 regard? That he doesn't know this guy? He does know  
14 this guy? This guy is a biochemist. What's the  
15 impeachment? My looking at this, it appears that he's  
16 just try to make an attack against Professor Atchison  
17 because he apparently has some religious views, which  
18 apparently is a theme throughout this case.

19 MR. ROTHSCHILD: That is absolutely not the  
20 case, Your Honor. And I think that will become clear as  
21 we go through the document.

22 THE COURT: All right. Inasmuch as this is  
23 a bench trial, I'm going to give Mr. Rothschild some  
24 latitude. I'll overrule the objection.

25 BY MR. ROTHSCHILD:

1       Q. While I was identifying myself as a Christian in  
2 Philadelphia, a biochemist named Michael Behe at Lehigh  
3 University was writing a book on evolution. As a  
4 biochemist, Behe found the evidence for Darwinian  
5 evolution to be very thin.

6           In fact, when he looked at the cell from a  
7 biochemical perspective, he believed there was evidence  
8 of intelligent design. Behe sent his completed  
9 manuscript to the Free Press publishers for  
10 consideration. That is your publisher of Darwin's Black  
11 Box, correct?

12       A. That's right.

13       Q. The editor was not certain that this manuscript  
14 was a good risk for publication. There were clearly  
15 theological issues at hand, and he was under the  
16 impression that these issues would be poorly received by  
17 the scientific community.

18           If the tenets of Darwinian evolution were  
19 completely accepted by science, who would be interested  
20 in buying the book? The next paragraph says, The editor  
21 shared his concerns with his wife. His wife was a  
22 student in my class. Again, this is consistent with  
23 your understanding of Mr. Atchison's -- Dr. Atchison's  
24 involvement?

25       A. Yes. As I said, I think the editor, his wife was

1 in vet school and knew that she was taking biochemistry  
2 and so asked the professor in that class.

3 Q. She advised her husband to give me a call. So  
4 unaware of all this, I received a phone call from the  
5 publisher in New York. We spent approximately ten  
6 minutes on the phone. After hearing a description of  
7 the work, I suggested that the editor should seriously  
8 consider publishing the manuscript.

9 I told him that the origin of life issue was  
10 still up in the air. It sounded like this Behe fellow  
11 might have some good ideas, although I could not be  
12 certain since I had never seen the manuscript. We hung  
13 up, and I never thought about it again, at least until  
14 two years later.

15 And then in the next session titled A Blessing  
16 Years Later, Dr. Atchison writes, After some time,  
17 Behe's book, Darwin's Black Box, the Free Press, 1996,  
18 was published. It became an instant best seller and was  
19 widely acclaimed in the news media.

20 It is currently in its 15th printing and over  
21 40,000 copies have been sold. I heard about it, but  
22 could not remember if this was the same book that I  
23 received the call about from the publisher. Could it  
24 be?

25 In November 1998, I finally met Michael Behe when

1 he visited Penn for a faculty outreach talk. He told me  
2 that, yes, indeed, it was his book that the publisher  
3 called me about. In fact, he said my comments were the  
4 deciding factor in convincing the publisher to go ahead  
5 with the book. Interesting, I thought.

6 You did meet Dr. Atchison, correct?

7 A. Yes, later, I did, yes.

8 Q. And is this your understanding of the kind of  
9 peer review Dr. Atchison did of your book?

10 A. No, it wasn't. I thought he had received a copy  
11 of the manuscript and went through it. So -- but -- so,  
12 yes, I was under a different impression.

13 Q. So he didn't review your manuscript carefully, he  
14 didn't review it at all, correct, Dr. Behe?

15 MR. MUISE: Objection, Your Honor. He has  
16 no personal knowledge. Again, he's using this document  
17 to assert the truth of the document, and Dr. Behe can  
18 only testify as to what his knowledge is.

19 THE COURT: I think that's a fair objection.  
20 You'll have to rephrase. The objection is sustained.

21 BY MR. ROTHSCHILD:

22 Q. You have no basis by which to dispute this  
23 account in this document, correct, Professor Behe?

24 A. My understanding is different from what is given  
25 in this account.



1 Q. And you did see some comments from some of your  
2 other reviewers, is that right?

3 A. That's correct.

4 Q. And they confirmed that you hadn't made any  
5 errors in the biochemistry, correct?

6 A. Yes.

7 Q. You were describing the bacterial flagellum  
8 correctly, its function, its appearance?

9 A. Yes.

10 Q. But they were reluctant or disagreed about  
11 intelligent design, correct?

12 A. Several were, yes, uh-huh.

13 Q. You also explained that, why you don't expect  
14 intelligent design at scientific conferences, correct?

15 A. Yes, that's because I consider it to be a poor  
16 forum for communicating such ideas.

17 Q. That's because typically you would present in the  
18 sort of poster sessions?

19 A. That's correct, yes.

20 Q. That doesn't really provide the opportunity to  
21 discuss it in detail to the audience?

22 A. That's correct, yes.

23 Q. It's difficult to impart understanding to your  
24 fellow scientists in that abbreviated form?

25 A. Yes. And not many come by. A few people wander

1 by, yes.

2 Q. It's not really an amenable way to present it?

3 A. That's right. It's usually brief conversations.

4 Q. You need to really present it in more detail for  
5 scientists to understand it?

6 A. That's why I discuss it in seminars and so on  
7 before scientific audiences, yes.

8 Q. Fair to say that, that rule probably makes even  
9 more sense with high school students, Professor Behe?

10 A. I'm sorry, what rule is that?

11 Q. The rule that you can't just present intelligent  
12 design in an abbreviated fashion?

13 A. Well, you certainly will not get a full  
14 understanding of intelligent design in a brief session.  
15 However, I think, if we're talking about high school  
16 students, such as you mentioned, it certainly might be a  
17 good thing to mention topics to them that they might  
18 consider pursuing in-depth outside the classroom.

19 Q. But an abbreviated statement is not going to give  
20 them a good understanding anymore than it would your  
21 fellow scientists, is that right?

22 A. A brief statement of any complex subject  
23 certainly will not give a person a complete  
24 understanding of it.

25 Q. Speaking of the students, you went through a

1 number of statements regarding evolution that you  
2 described as philosophical and religious, correct?

3 A. You mean, during my testimony yesterday?

4 Q. I think it was Monday, or maybe it was yesterday.  
5 It's hard to keep track. But some statements by  
6 Professor Miller, by Dr. Dawkins, by Peter Singer?

7 A. Yes, I did.

8 Q. And you would characterize those as  
9 non-scientific statements, rather philosophical or  
10 religious or political statements?

11 A. That's correct.

12 Q. Should they be taught to students in a high  
13 school biology class?

14 A. Well, that's an interesting idea. Since a high  
15 school biology class, in my opinion, is not, should not  
16 simply be focused on producing scientists for the next  
17 generation, since most students won't go on to become  
18 scientists, but rather it's for their liberal education,  
19 understanding science, and also understanding science's  
20 role in the world, I think, in fact, it might be  
21 appropriate not to teach this in a sense of saying, here  
22 are things that are true, but to discuss the comments  
23 that have been made about scientific theories that they  
24 are learning in their class to show the students that  
25 science is not something that is confined to the

1 library, but the ideas generated by science have far  
2 reaching ramifications in the opinion of many learned  
3 people, and that, here are some of them. And I think  
4 that's actually an excellent idea for a science  
5 classroom.

6 Q. In biology class?

7 A. In biology class, in physics class, and other  
8 science classes as well.

9 Q. And you definitely agree that students should be  
10 taught that some biochemical systems are intelligently  
11 designed, correct?

12 A. I'm sorry. Could you restate --

13 Q. Your testimony over the last two days stands for  
14 the proposition that students should be told that  
15 biological life has been intelligently designed?

16 A. I'm afraid I don't think I said that. And if I  
17 did, I'm not quite -- well, I'm not sure that I said  
18 that. I didn't say, students should be told that some  
19 biochemical systems are intelligently designed. If I  
20 said that -- it's a good idea to give students a couple  
21 different frameworks where some data has been  
22 interpreted, so that they can see the difference between  
23 fact and theory, fact and interpretation, and so on.

24 I think intelligent design is, in fact, a good  
25 way to do that, yes.

1       Q. Fair to say that, what you're saying is that, one  
2 valid scientific interpretation that should be taught to  
3 students, along with other theories, is that some  
4 aspects of biological life were intelligently designed?

5       A. I'm saying that, in their discussion of these  
6 issues, students can be told that some scientists have  
7 proposed this idea, and here are the reasons that they  
8 propose. Here are the data that they point to. Here is  
9 what other scientists have proposed.

10           They have proposed a different theory. Here is  
11 the data that they point to. Here are the explanations  
12 they give. Here are the responses that they gave to  
13 that first group. Here are the responses that the first  
14 group gave back. The point -- I'm sorry. The point is  
15 to -- is not to instruct students that this view is  
16 correct, as we've heard many times here.

17           We know that theories can be wrong, that no  
18 theory is guaranteed to be true. So the point is to get  
19 them to discuss data from different points of view.

20       Q. So students should be told that one scientific  
21 theory is that some aspects of biological life were  
22 intelligently designed?

23       A. I think it would be good pedagogy to discuss the  
24 fact that some scientists do think that some aspects of  
25 life were intelligently designed, yes.

1 Q. By an intelligent designer?

2 A. Well, intelligently designed, yes, it implies a  
3 designer, yes.

4 Q. So students should be told that there is a  
5 scientific theory or that scientists contend that some  
6 aspects of biological life were intelligently designed  
7 by an intelligent designer, good pedagogy?

8 A. Again, I think you have to look at the context.  
9 There is a tendency for people to think that when you  
10 say, you're going to teach something in the classroom,  
11 that means you're going to present it to students and  
12 tell them that is true.

13 Q. I'm not suggesting that, Professor Behe. My  
14 question was, you think it's good pedagogy --

15 MR. MUISE: Objection, Your Honor. He's  
16 attempting to answer the question.

17 MR. ROTHSCHILD: He's attempting to evade  
18 the question, Your Honor. I'm being very clear. He  
19 helped me correct it, and I corrected it.

20 THE COURT: Let's let him finish the answer.  
21 Finish the answer.

22 THE WITNESS: It's just that -- I'm just  
23 saying that students should be presented different views  
24 for discussion, not in the sense of saying, this is  
25 either valid or not valid, this is true or not true, but

1 just to give different points of view.

2 BY MR. ROTHSCHILD:

3 Q. I understand that. So what you're saying is,  
4 it's good pedagogy to tell students that one scientific  
5 theory about biological life is that some aspects of  
6 biological life were designed by an intelligent  
7 designer?

8 A. I would phrase it differently. I would say, it's  
9 good pedagogy to tell some students that some people  
10 think that this is the case.

11 Q. Fair enough. Is it also good pedagogy to tell  
12 students in biology class, some scientists argue that  
13 there is no intelligent designer?

14 A. I think it would be good pedagogy to point out  
15 that, in fact, the majority view of science is that  
16 random mutation and natural selection without any  
17 apparent design is responsible for what we find in  
18 biology.

19 Q. And included in that statement, it would be good  
20 pedagogy to tell students, those scientists contend  
21 there is no intelligent designer? Is that good  
22 pedagogy, to tell students that scientists think there  
23 is no intelligent designer?

24 A. No, it would not be good pedagogy, because there  
25 are many different ideas tangled together in your

1 statement. Many scientists who think that, for example,  
2 Darwinian processes are correct, nonetheless do think  
3 that there is a designer in a different sense.

4 One is using the word designer here in several  
5 different senses; designer of laws of nature versus  
6 designer of specific aspects of nature, and so on. So I  
7 think your question is a bit ambiguous.

8 Q. Fair to say that my statement, that telling  
9 students there is no intelligent designer, has religious  
10 and philosophical baggage as well as scientific?

11 A. I'm sorry. Would you say that again?

12 Q. Fair to say that the statement I propose, telling  
13 students there is no intelligent designer in science  
14 class, has religious and philosophical aspects?

15 A. Yes. Like many theories, it does.

16 Q. Are there gaps and problems with the theory of  
17 intelligent design?

18 A. Yes.

19 Q. Should students, high school students being made  
20 aware of intelligent design be made aware that there are  
21 gaps and problems in the theory of intelligent design?

22 A. Absolutely.

23 Q. If they are being made aware of intelligent  
24 design, but are not being told there are gaps and  
25 problems in intelligent design, are they being misled,



1 Professor Behe?

2 A. Well, again, they're not receiving full  
3 instruction then in intelligent design. And so you  
4 could, if you had more time, you could certainly go into  
5 those, and I would certainly recommend that you do so.

6 MR. ROTHSCCHILD: May I approach the witness?

7 THE COURT: You may.

8 BY MR. ROTHSCCHILD:

9 Q. Professor Behe, what I've showed you is  
10 Plaintiffs' Exhibit 721. Do you recognize that as the  
11 article you wrote with David Snoke entitled Simulating  
12 Evolution by Gene Duplication of Protein Feature that  
13 Requires Multiple Amino Acid Residues?

14 A. Yes.

15 Q. And you discussed that over the last couple days?

16 A. Yes.

17 Q. Now in this, you described this as a theoretical  
18 paper?

19 A. Yes.

20 Q. You didn't culture organisms?

21 A. No.

22 Q. Or isolate proteins?

23 A. No, this was a computer study.

24 Q. Okay. Like what you criticized Dr. Pennock for  
25 doing?

1       A. I didn't criticize him for doing computer  
2 studies. I criticized his particular model because I  
3 thought it was not -- it had dissimilarities or it had  
4 assumptions built into it that I thought were  
5 inappropriate.

6       Q. It didn't represent what actually happens in  
7 biological life, that's your --

8       A. That's correct, yes.

9       Q. It didn't represent what is actually understood  
10 to happen in the theory of evolution?

11      A. Well, some aspects of it were sort of like what  
12 has happened in evolution, but it was -- it went a  
13 little bit too far afield, in my opinion, for it to be a  
14 useful model.

15      Q. And this study, this computer simulation was  
16 based on gene sequences that were published by other  
17 laboratories or other researchers?

18      A. No, not really, no. It was a -- based  
19 essentially on simply what we know about protein  
20 structure, was not a sequence study.

21      Q. When you say, what we know about protein, that  
22 was based on the work of other researchers?

23      A. Yes, uh-huh.

24      Q. And you studied a particular type of mutation, a  
25 point mutation?

1       A. That's correct.

2       Q. And let me just ask you a few questions, and you  
3 tell me if I'm fairly summarizing the results of your  
4 computer simulation. What you're asking is, how long  
5 will it take to get -- and please follow with me, I'm  
6 trying to do this slowly and methodically -- two or more  
7 specific mutations, in specific locations, in a specific  
8 gene, in a specific population, if the function is not  
9 able to be acted on by natural selection until all the  
10 mutations are in place, if the only form of mutation is  
11 point mutation, and the population of organisms is  
12 asexual?

13       A. I would have to look at that statement closely  
14 because there are so many different aspects to it that I  
15 don't trust myself to sit here and listen to you say  
16 that and form a correct judgment.

17       Q. Anything I said about that sound incorrect?

18       A. If you repeat it again, I'll try.

19       Q. I'd be happy to. Two or more specific mutations?

20       A. Actually, this dealt with one or more.

21       Q. One or more mutations?

22       A. Yes. If you notice, in figure -- if you notice  
23 in figure 3, you look at the x axis, you notice that  
24 there are data points there that start at one. So we  
25 considered models where there were one, two, and more

1 mutations.

2 Q. Fair enough. In specific locations?

3 A. No, that's not correct. We assumed that there  
4 were several locations in the gene that could undergo  
5 these selectable mutations, but we did not designate  
6 where they were.

7 Q. In the specific gene?

8 A. We were considering one gene, yes.

9 Q. In a specific population?

10 A. Yes.

11 Q. Okay. If the function is not able to be acted on  
12 by natural selection until all mutations are in place?

13 A. Yes, that's what's meant by multiple amino acid  
14 residue, multi-residue feature, yes.

15 Q. If the only form of mutation is point mutation?

16 A. Yes, that's a very common type of mutation, which  
17 is probably half or more of the mutations that occur in  
18 an organism.

19 Q. And if the population of organisms is asexual?

20 A. Yes, we did not -- actually, we did not confine  
21 it just to asexuals, but we did not consider  
22 recombination.

23 Q. Are prokaryotes an example of the kind of  
24 organism that you were studying there?

25 A. Again, we weren't studying organisms, but, yeah,

1 they're a good example of what such a model has in mind.

2 Q. And to say this very colloquially, you conclude  
3 that it will take a large population a long time to  
4 evolve a particular function at disulfide bond, right?

5 A. A multi-residue feature. That's correct, that's  
6 correct.

7 Q. And specifically --

8 A. I'm sorry.

9 Q. Go ahead.

10 A. Let me just finish. Depending on -- as we  
11 emphasize in the paper, it depends on the population  
12 size. And, of course, prokaryotes can oftentimes grow  
13 to very large population sizes.

14 Q. And here the conclusion, the calculations you  
15 concluded was that, if you had a population of 10 to the  
16 9th power, that's a population of 1 billion?

17 A. That's correct.

18 Q. To produce a novel protein feature through the  
19 kind of multiple point mutations you're talking about,  
20 it would take 10 to the 8th generations, that's what it  
21 says in the abstract, correct?

22 A. If, in fact, it was -- if, in fact, the  
23 intermediate states were not selectable.

24 Q. Okay.

25 A. And if this is by gene duplication as well.

1       Q.   Okay.  So 10 to the 8th generation, that's 100  
2 million generations?

3       A.   That's correct.

4       Q.   And yesterday, you explained about bacteria, that  
5 10,000 generations would take about two years in the  
6 laboratory, correct?

7       A.   Yes.

8       Q.   So 100 million generations, that would take about  
9 20,000 years?

10      A.   I'm sorry?

11      Q.   100 million generations, which is what you  
12 calculated here, that would take about 20,000 years?

13      A.   Okay, yes.

14      Q.   And those are numbers based on your probability  
15 calculations in this model, correct?

16      A.   Yes.

17      Q.   Now it would be true that, if you waited a little  
18 longer, say, instead of 10 to 9th generations, 10 to the  
19 10th generations, then it would mean that you wouldn't  
20 need as big a population to get the function that you  
21 are studying?

22      A.   That's right.  The more chances you have, the  
23 more likely you are to develop a feature.  And the  
24 chances are affected by the number of organisms.  So if  
25 you have a smaller population time, and more

1 generations, that could be essentially equal to a larger  
2 population size and fewer generations.

3 Q. So, as you said, so if we get more time, we need  
4 less population to get to the same point, and if we had  
5 more population, less time?

6 A. That's correct, yes.

7 Q. Now would you agree that this model has some  
8 limitations?

9 A. Sure.

10 Q. And you, in fact, were quite candid in indicating  
11 that in the paper?

12 A. That's correct.

13 Q. And if we could turn to, what I believe is, page  
14 8 of the document. And if you look in the paragraph  
15 that's actually continued from the previous page that  
16 says, we strongly emphasize. And if you could --

17 A. I'm sorry. What page number is that?

18 Q. It's page 8 in the document. And it's up on the  
19 screen as well.

20 A. Yes, okay. I've got it.

21 Q. Could you read into the record the text to the  
22 end of the paragraph beginning with, we strongly  
23 emphasize?

24 A. We strongly emphasize that results bearing on the  
25 efficiency of this one pathway as a conduit for

1 Darwinian evolution say little or nothing about the  
2 efficiency of other possible pathways. Thus, for  
3 example, the present study that examines the evolution  
4 of MR protein features by point mutation in duplicate  
5 genes does not indicate whether evolution of such  
6 features by other processes, such as recombination or  
7 insertion/deletion mutations, would be more or less  
8 efficient.

9 Q. So it doesn't include recombination, it doesn't  
10 include insertion/deletion of the mutations?

11 A. That's correct.

12 Q. And those are understood as pathways for  
13 Darwinian evolution?

14 A. They are potential pathways, yes.

15 Q. This study didn't involve transposition?

16 A. No, this focuses on a single gene.

17 Q. And transpositions are, they are a kind of  
18 mutation, is that right?

19 A. Yes. They can be, yes.

20 Q. And so that means, this simulation didn't examine  
21 a number of the mechanisms by which evolution actually  
22 operates?

23 A. That is correct, yes.

24 Q. And this paper, let's be clear here, doesn't say  
25 anything about intelligent design?



1           A. Yes, that's correct. It does imply irreducible  
2 complexity but not intelligent design.

3           Q. But it doesn't say it?

4           A. That's correct.

5           Q. And one last other question on your paper. You  
6 concluded, it would take a population size of 10 to the  
7 9th, I think we said that was a billion, 10 to the 8th  
8 generations to evolve this new disulfide bond, that was  
9 your conclusion?

10          A. That was the calculation based on the assumptions  
11 in the paper, yes.

12                   MR. ROTHSCHILD: May I approach the witness,  
13 Your Honor?

14                   THE COURT: You may.

15 BY MR. ROTHSCHILD:

16          Q. What I've marked as Exhibit P-756 is an article  
17 in the journal Science called Exploring Micro--

18          A. Microbial.

19          Q. Thank you -- Diversity, A Vast Below by T.P.  
20 Curtis and W.T. Sloan?

21          A. Yes, that seems to be it.

22          Q. In that first paragraph, he says, There are more  
23 than 10 to the 16 prokaryotes in a ton of soil. Is that  
24 correct, in that first paragraph?

25          A. Yes, that's right.

1 Q. In one ton of soil?

2 A. That's correct.

3 Q. And we have a lot more than one ton of soil on  
4 Earth, correct?

5 A. Yes, we do.

6 Q. And have for some time, correct?

7 A. That's correct, yes.

8 Q. And, in fact, he gives us a good way of comparing  
9 it. It says, as compared to a mere 10 to the 11th stars  
10 in our galaxy?

11 A. Yes, that's what he writes, uh-huh.

12 Q. And 10 to the 16th prokaryotes is 7 orders of  
13 magnitude higher than the population you included in  
14 your calculations, correct?

15 A. No. We considered a wide range of populations,  
16 and we considered a wide range of number of  
17 substitutions that would be -- or point mutations that  
18 would be necessary. You're focusing on two, but perhaps  
19 I can direct your attention again to that figure from  
20 the paper -- excuse me. Let me find it.

21 The best place I think to look is figure 6, which  
22 is on page 10 of the document. Up in the upper  
23 right-hand corner, that figure there.

24 Q. Sure.

25 A. If you look on the bottom, the x axis there, the

1 bottom of the figure that's labeled lambda, it has the  
2 numbers 2, 4, 6, 8, 10, and so on, those are the number  
3 of point mutations that we consider perhaps some  
4 multi-residue feature might entail. As we said in the  
5 paper, forming a new disulfide bond might require as few  
6 as two point mutations.

7 But forming other multi-residue features such as  
8 protein, protein binding sites might require more. And  
9 so the number on the X axis lambda 2, 4, 6, 8, those are  
10 the number of point mutations that we entertained or we  
11 calculated numbers for to see how long such things would  
12 be expected to take under our model.

13 And if you look up at the top axis, the top x  
14 axis labeled N, at the top of the figure. N stands for  
15 population size. Okay. So if you look at the figures  
16 there on the left, it's slanted, and it's not enlarged  
17 yet, so it's hard to see. It says, 10 to the 6th.  
18 That's a million. And then skip a line. These are in  
19 every 10 to the 3rd increments of population size. That  
20 would be 10 to the 9th.

21 The next label is 10 to the 12th, which is a  
22 trillion. The next label is 10 to the 18th, which is  
23 much more. The next label is 10 to the 24th, which is  
24 much, much, much more. The next label, 10 to the 30th,  
25 which, again, is very much more.

1           So, in fact, we considered population sizes from  
2 1000 all the way up to 10 to the 30th, and multi-residue  
3 features from 2, which might involve disulfide bonds, up  
4 to many more, which might be involved in protein,  
5 protein binding sites.

6           Q. 10 to the 30th, that is quite a lot, right?

7           A. Yes. That's roughly what is calculated to be the  
8 bacterial population of the Earth in any one year. And  
9 so over the course of the billion year, 4 billion year  
10 history of the Earth, there would probably be a total of  
11 roughly 10 to the 40th.

12          Q. And so in the case of prokaryotes, which you said  
13 was a good example of what you were studying, 10 to the  
14 16th in one ton of soil?

15          A. Yes.

16          Q. So a few tons of soil, and we've gone past that  
17 10 to the 30th?

18          A. Well, no. In the 10 to the 14th tons of soil.  
19 10 to the 30th is the number that's in the entire world,  
20 according to the best estimates, including the ocean as  
21 well as soil. So -- but I agree with your point, that  
22 there's a lot of bacteria around and certainly more than  
23 10 to the 9th.

24          Q. So just with the prokaryotes, 10 to the 16th, 7  
25 orders of magnitude higher than what you were

1 calculating here?

2 A. That's certainly true, but in our paper, we had  
3 our eye not only on prokaryotes, but also on eukaryotes  
4 as well, which, if you leave out recombination, one can  
5 -- they certainly undergo point mutations. They  
6 certainly have genes and so on. So much of this is also  
7 applicable to eukaryotes.

8 And the populations of eukaryotes and certainly  
9 larger plants and animals are much, much smaller than  
10 populations of bacteria. So we view our results not  
11 just as supplying that, but to giving us some feel for  
12 what can happen in more complex organisms as well.

13 Q. Well, you're not talking about more complex  
14 organisms here, are you?

15 A. I think we do. I think at the end, if I'm not  
16 mistaken, if I remember correctly -- okay, yes. On page  
17 11, the second full paragraph, on page 11. It begins on  
18 the right-hand column, the second full paragraph. It  
19 says, The lack of recombination in our model means it is  
20 most directly applicable to haploid, asexual organisms.  
21 Nonetheless, the results also impinge on the evolution  
22 of diploid sexual organisms.

23 The fact that very large population sizes, 10 to  
24 9th or greater, are required to build even a minimal MR  
25 feature requiring two nucleotide alterations within 10

1 to the 8th generations by the processes described in our  
2 model, and that enormous population sizes are required  
3 for more complex features or shorter times, seems to  
4 indicate that the mechanism of gene duplication and  
5 point mutation alone would be ineffective, at least for  
6 multicellular diploid species, because few multicellular  
7 species reach the required population sizes.

8 Thus, mechanisms in addition to gene duplication  
9 and point mutation may be necessary to explain the  
10 development of MR features in multicellular organisms.

11 So here we were trying to point out that, because  
12 of the results of the calculation, it seems that, when  
13 we're trying to explain MR features in multicelled  
14 organisms, then we're going to have to look to other  
15 processes for that.

16 Q. Okay. So if we exclude some of the processes by  
17 which we understand evolution to occur, it's hard to get  
18 there for multicellular organisms?

19 A. I'm sorry.

20 Q. If we exclude some of the mechanisms by which we  
21 understand evolution to occur, like recombination, it's  
22 hard to get there?

23 A. Yes.

24 Q. And bringing it back to the prokaryotes. We're  
25 in agreement here, the number of prokaryotes in 1 ton of

1 soil are 7 orders of magnitude higher than the  
2 population, you said it would take 10 to the 8th  
3 generations to produce the disulfide bond?

4 A. Yeah, certainly. Yeah, the bacteria are -- can  
5 grow to very large population sizes.

6 Q. So the time would be?

7 A. Much shorter.

8 Q. Much shorter?

9 A. Absolutely.

10 MR. ROTHSCHILD: Your Honor, this would be a  
11 good time to take a break.

12 THE COURT: All right. Why don't we take  
13 our morning recess now, and we will return in about 20  
14 minutes. Thank you.

15 (Whereupon, a recess was taken at 10:16 a.m.

16 and proceedings reconvened at 10:40 a.m.)

17 THE COURT: All right. We resume with Mr.  
18 Rothschild.

19 MR. ROTHSCHILD: Thank you.

20 **CROSS EXAMINATION (CONTINUED)**

21 BY MR. ROTHSCHILD:

22 Q. Professor Behe, I'd like to turn our attention  
23 now to Darwin's Black Box. What you explain in Darwin's  
24 Black Box is that, modern science has been able to  
25 explore life at the molecular level in a way that was

1 not possible with Darwin, is that right?

2 A. That's right.

3 Q. Or actually for sometime after?

4 A. That's correct.

5 Q. And it's that life at the molecular level that  
6 you are referring to when you call it Darwin's Black  
7 Box, something he couldn't look into?

8 A. That's correct.

9 Q. In fact, in the book, you call it the last black  
10 box?

11 A. Is that right? Could you show me where I do  
12 that?

13 Q. Sure.

14 A. I'm sorry.

15 Q. If you could turn to page 13.

16 A. Yes.

17 Q. Okay. And if you look at the paragraph, you  
18 quote from a ditty from Jonathan Swift?

19 A. Yes.

20 Q. And then you say, in the late 20th century, we  
21 are in the flood tide of research on life, and the end  
22 is in sight. The last remaining black box was the cell,  
23 which was opened to reveal molecules, the bedrock of  
24 nature, the last black box, correct?

25 A. I'm sorry. Yes. Okay, the last remaining black



1 box was the cell, yes.

2 Q. Okay. And then you conclude at the end of that  
3 paragraph, that black box now stands open?

4 A. Yes.

5 Q. And I think you've testified, and I think it's  
6 apparent in your book that, science has discovered a  
7 level of complexity that prior generations of scientists  
8 never predicted?

9 A. That's correct.

10 Q. And your conclusion is that, that complexity  
11 provides an insurmountable obstacle to Darwinian  
12 evolution?

13 A. Well, you always try to avoid words like  
14 insurmountable, but it certainly points to severe  
15 problems for it, yes.

16 Q. And you reached the conclusion that certain  
17 biochemical systems could not be produced by natural  
18 selection because they are irreducibly complex?

19 A. Again, you've got to be careful about using  
20 absolutes like could not, but it certainly seems like  
21 they could not.

22 Q. And these systems also have what you describe as  
23 a purposeful arrangement of parts?

24 A. Yes.

25 Q. And, therefore, you concluded they were

1 intelligently designed?

2 A. Yes.

3 Q. And in terms of the structure of the systems, you  
4 base your conclusions on work on the structure and  
5 function of those molecular systems done by other  
6 scientists?

7 A. That's correct.

8 Q. Many other scientists?

9 A. That's correct.

10 Q. And you read a lot of papers that published in  
11 peer review journals describing the structure and  
12 function of the systems that you discuss in the book?

13 A. That's correct.

14 Q. And those scientists in those papers don't argue  
15 that their work supports irreducible complexity as you  
16 define it?

17 A. That's correct.

18 Q. Or intelligent design?

19 A. That's correct.

20 Q. And, in fact, a good number of them would have  
21 actively opposed that?

22 A. And still do.

23 Q. And the -- Matt, if you could pull up page 39,  
24 please, and highlight the bottom paragraph there at the  
25 bottom. This is the place in Darwin's Black Box where

1 you explain what you mean by irreducibly complex?

2 A. Yes.

3 Q. And as you testified, I believe, on Monday, a  
4 scientist named Alan Orr noted an ambiguity in your  
5 definition?

6 A. Yes.

7 Q. And you responded to that?

8 A. Yes.

9 Q. And you tweaked that definition?

10 A. Right.

11 Q. Matt, could you pull up the tweaked definition  
12 that he created? And I have inserted the words which is  
13 necessarily composed to make this paragraph consistent  
14 with the tweaking you described you did in response to  
15 Alan Orr. And I'm going to read that. And I've called  
16 it here the modified definition of irreducible  
17 complexity from Darwin's Black Box.

18 What it says is, By irreducibly complex, I mean a  
19 single system which is necessarily composed of several  
20 well-matched, interacting parts that contribute to the  
21 basic function, wherein the removal of any one of the  
22 parts causes the system to effectively cease  
23 functioning.

24 An irreducibly complex system cannot be produced  
25 directly, that is by continuously improving the initial

1 function which continues to work the same mechanisms by  
2 slight successive modifications of a pre-cursor system,  
3 because any pre-cursor to an irreducibly complex system  
4 that is missing a part is, by definition,  
5 non-functional.

6 An irreducibly complex biological system, if  
7 there is such a thing, would be a powerful challenge to  
8 Darwinian evolution. Since natural selection can only  
9 choose systems that are already working, then if a  
10 biological system cannot be produced gradually, it would  
11 have to arise as an integrated unit in one fell swoop  
12 for natural selection to have anything to act on.

13 So that's the last paragraph on page 39 adding  
14 the words that you did in response to Dr. Orr?

15 A. Yes.

16 Q. And when you say, it would have to arise as an  
17 integrated unit in one fell swoop for natural selection  
18 to have anything to act on, what you're saying is,  
19 whatever the proposed pre-cursor was, would die because  
20 it doesn't have all of its parts?

21 A. No, that's not correct. Die is not -- the  
22 function of a system is not to live, it's to do  
23 something particular. You say that the system did not  
24 work, it did not do its function. For example, the  
25 bacterial flagellum would not work without the necessary

1 parts.

2 Q. And, therefore, there would be no successive  
3 generation because that flagellum would not move on to  
4 the next generation?

5 A. No, that's not right. A bacterium that is  
6 missing a flagellum would certainly go on and continue  
7 to grow. It can reproduce and so on. But the flagellum  
8 doesn't work. And this is from my article, I believe,  
9 in *Biology and Philosophy*, where I responded to  
10 Professor Orr.

11 And in that article, I specifically said that he  
12 had a misconception that irreducible complexity meant  
13 that an organism could not live without this, without  
14 the system that we were talking about. And that's not  
15 what I meant by it.

16 Q. So the organism with half a flagellum or parts of  
17 a flagellum could continue to live in that circumstance,  
18 it just wouldn't have an operating flagellum?

19 A. Sure, yes.

20 Q. Now could you turn again to Exhibit 718, which is  
21 that article, *Reply to my Critics*, that you just  
22 discussed?

23 A. Yes.

24 Q. Okay. On -- could you turn to page 695?

25 A. Yes.

1       Q. And in the first full paragraph, you repeat some  
2 of the text that we just saw from Darwin's Black Box  
3 about why irreducible complex systems are obstacles for  
4 Darwinian explanations?

5       A. Yes.

6       Q. And then you write, However, commentary by Robert  
7 Pennock and others has made me realize that there is a  
8 weakness in that view of irreducible complexity. The  
9 current definition puts the focus on removing a part  
10 from an already functioning system.

11             And then continuing on after footnote 5, you say,  
12 The difficult task facing Darwinian evolution, however,  
13 would not be to remove parts from sophisticated  
14 pre-existing systems, it would be to bring together  
15 components to make a new system in the first place.

16             Thus, there is an asymmetry between my current  
17 definition of irreducible complexity and the task facing  
18 natural selection. I hope to repair this defect in  
19 future work. That's what you wrote, correct?

20       A. Yes.

21       Q. You haven't repaired that defect, have you,  
22 Professor Behe?

23       A. No, I did not judge it serious enough to do so  
24 yet.

25       Q. So the defect you identified was, you were

1 starting with the function and working backwards,  
2 removing parts, correct?

3 A. That's correct, yes.

4 Q. And natural selection is actually operating in  
5 the opposite direction, you start with the pre-cursors  
6 and then develop until you get to the system you're  
7 studying?

8 A. Yes, that would be a more difficult task.

9 Q. That's the asymmetry?

10 A. Yes.

11 Q. And that asymmetry has not been repaired?

12 A. That asymmetry is not really relevant to  
13 biological circumstances. In the sentence that you  
14 skipped over in that paragraph, I talk about what  
15 Professor Pennock discussed in his book in making this  
16 point.

17 If I could just quote from that. He says, Thus,  
18 seeking a counterexample to irreducible complexity  
19 entower a battle. Pennock writes about a part in a  
20 sophisticated chronometer whose origin is simply assumed  
21 which breaks to give a system that he posits can  
22 nonetheless work in a simpler watch in a less demanding  
23 environment.

24 So I viewed Professor Pennock's objection -- of  
25 course, Professor Pennock is a philosopher, and that was

1 an interesting philosophical turn on my discussion, I  
2 thought, but that is not -- that is not -- I did not  
3 consider that to be relevant to biology.

4 Q. Okay. The task facing natural selection, that's  
5 not relevant to biology?

6 A. No, the particular pathway that Professor Pennock  
7 had in mind where one assumes that one has a very  
8 sophisticated pre-existing system whose origin has been  
9 left unexplained and has just postulated, which then  
10 goes on to breakdown and give less sophisticated parts,  
11 that is the part that I don't think is really relevant  
12 to biology.

13 Q. If you start with the system and then break it  
14 down removing parts, that's not relevant to biology?

15 A. Well, that's not the difficult task facing  
16 evolution.

17 Q. Right. And you're not testing the natural -- the  
18 difficult task facing evolution, which starts from the  
19 pre-cursors and moves forward to the system you're  
20 studying. You're going backwards. Isn't that what  
21 irreducible complexity proposes?

22 A. It does not propose that anything goes backwards.  
23 It asks, how do we identify this problem for Darwinian  
24 evolution? And if you can remove a part, and a system  
25 no longer works, then the system needs those parts to



1 work.

2           And so the problem, how you put that together by  
3 numerous successive slight modifications, as Charles  
4 Darwin thought one had to do, is, I think, illustrated  
5 by that.

6       Q. In any event, you have not repaired this  
7 asymmetry?

8       A. That's correct.

9       Q. And that article was written four years ago,  
10 correct?

11      A. Yes.

12      Q. Now you've used the expression, produced  
13 directly. I think that's in the definition. Matt, if  
14 you could pull that back up. And if I understand what  
15 you mean by directly, it means, for example, in the case  
16 of the flagellum, that it has to be steps in which  
17 there's a rotary motor that continues to become the  
18 rotary motor, that is the flagellum?

19      A. Yes. By direct, I mean that it essentially  
20 worked, as the definition says, it works by the same  
21 mechanism, has the same number of parts; essentially,  
22 it's the same thing.

23      Q. Same thing. And then if you could turn to page  
24 40 of Darwin's Black Box. Matt, if you could highlight  
25 the first paragraph. You acknowledge another

1 possibility?

2 A. That's correct.

3 Q. You say, Even if a system is irreducibly complex  
4 and thus could not have been produced directly, however,  
5 one cannot definitively rule out the possibility of an  
6 indirect, circuitous route, right?

7 A. Yes.

8 Q. And by indirect, you mean evolution from a  
9 pre-cursor with a different function than the system  
10 being studied?

11 A. Yes, different function, perhaps different number  
12 of parts, and so on.

13 Q. And one example of that is what's discussed in,  
14 among evolutionary biologists, as the concept of  
15 exaptation, correct?

16 A. Yeah -- well, before I say, yes, I'd just like to  
17 say, the word exaptation is oftentimes used in loose  
18 sense, but, yes, that's generally correct.

19 Q. And that is a concept that people in the field of  
20 evolutionary biology consider to be a valid concept, a  
21 valid description of the way more and more complex  
22 systems get developed?

23 A. Let me say --

24 Q. I'm not asking you to agree with it. I'm asking  
25 you, is that what an evolutionary biologist proposes?

1       A. Again, let me make clear what we're talking about  
2 here. Some evolutionary biologists certainly think that  
3 exaptation is real and that it's important and so on.  
4 But simply saying that this part over here was exapted  
5 from that part over here does not give an explanation of  
6 how random mutation and natural selection could have  
7 gotten it from one state to the other.

8       Q. But it is certainly, exaptation -- for example, a  
9 bird wing developing from some kind of feathered  
10 structure on a dinosaur that didn't necessarily allow  
11 flight, that's what evolutionary biologists propose, and  
12 they call it exaptation?

13       A. That's entirely possible, and that's consistent  
14 with intelligent design, because intelligent design only  
15 focuses on the mechanism of how such a thing would  
16 happen. So the critical point for my argument is, how  
17 such things could develop by random mutation and natural  
18 selection.

19       Q. And again, intelligent design doesn't describe  
20 how it happened?

21       A. That's correct, only to say that intelligence was  
22 involved somewhere in the process.

23       Q. Okay. Now you go on in this passage and say, As  
24 the complexity of an interacting system increases,  
25 though, the likelihood of such an indirect route drops

1 precipitously, and as the number of unexplained  
2 irreducibly complex biological systems increases, our  
3 confidence that Darwinian's criterion of failure has  
4 been met and skyrockets toward the maximum that science  
5 allows?

6           What you're saying there is, you know, it could  
7 happen, I'm not ruling it out, but it's really  
8 improbable?

9           A. Yes, it's improbable.

10          Q. Okay. And you haven't -- and based on that, you  
11 conclude that intelligent design is a much more probable  
12 explanation?

13          A. Not just based on that, based on the purposeful  
14 arrangement of parts.

15          Q. Fair enough. And you haven't actually quantified  
16 this, have you?

17          A. Not explicitly, but as a biochemist who  
18 understands what it takes to, for example, for a protein  
19 to function, for two proteins to bind specifically to  
20 each other, and so on, I rely on my experience of that  
21 in arriving at this conclusion.

22          Q. And you've seen how long it takes for the  
23 prokaryotes to bind?

24          A. 10 to the 16th in one ton of soil, yes, uh-huh.

25          Q. Now just to be clear -- in this passage, you say,

1 irreducibly complex biological systems, right?

2 A. I'm sorry?

3 Q. In this passage, you say, As the number of  
4 unexplained irreducibly complex biological systems  
5 increases, right, that's what it says there?

6 A. Yes. Yes, I do, uh-huh.

7 Q. But you took pains on Monday to communicate to  
8 the Court that when you're talking about irreducible  
9 complexity, you're just talking about it at the  
10 molecular level?

11 A. Yes, that should be biochemical instead of  
12 biological.

13 Q. Fair enough. You don't make claims about  
14 irreducible complexity at the organ level?

15 A. That's correct.

16 Q. Or at the organism level?

17 A. That's correct.

18 Q. In fact, you don't have any expertise or training  
19 in the organ or organism level?

20 A. That's correct, yes.

21 Q. You also have no expertise in paleontology?

22 A. That's correct.

23 Q. Or physics?

24 A. That's correct, too.

25 Q. Sorry. Couldn't resist. We've gone a long time.

1 But you agree that intelligent design, as opposed to  
2 just Michael Behe, is making an argument for intelligent  
3 design far beyond the cellular level, correct?

4 A. I'm sorry?

5 Q. Intelligent design, as a scientific proposition  
6 and the individuals who advocate for it, are arguing for  
7 intelligent design beyond the cellular level?

8 A. Some people certainly do, based not on my  
9 argument but other arguments.

10 Q. So it's not based on your argument?

11 A. Yes.

12 Q. And, for example, in Pandas, that's certainly in  
13 play intelligent design of not just biochemical  
14 structures but higher level forms?

15 A. Well, let me just correct myself. They're not  
16 basing it on my argument in regard to irreducible  
17 complexity, but they are basing it on the purposeful  
18 arrangement of parts, which is certainly what I discuss  
19 in Darwin's Black Box.

20 Q. In Darwin's Black Box, you talk about a  
21 purposeful arrangement of parts, and you actually say,  
22 you know, using that standard, almost anything looks  
23 design, right?

24 A. I don't think I said that.

25 Q. We'll return to that. In any event, in Pandas,

1 there are arguments for intelligent design of higher  
2 level biological life?

3 A. Yes, there are.

4 Q. And we're clear, that's not based on your work?

5 A. It's not based on any concept of irreducible  
6 complexity. It is based on a concept that I discuss in  
7 Darwin's Black Box, the purposeful arrangements of  
8 parts.

9 Q. That purposeful arrangement of parts, that's not  
10 -- you didn't originate that?

11 A. No, I didn't.

12 Q. At least, it goes back to Reverend Paley?

13 A. Yes, it does. Further back than that.

14 Q. Now let's start with the bacterial flagellum.  
15 You've made a point about how complicated and intricate  
16 it is?

17 A. Yes.

18 Q. And it really is. I mean, it looks remarkable.  
19 But a lot of biological life is pretty remarkable?

20 A. That makes me very suspicious.

21 Q. You're suspicious about how remarkable biological  
22 life is?

23 A. No, it makes me suspicious, you know -- that was  
24 a joking way to say that I think much of biological life  
25 may bespeak design.

1 Q. Plants and photosynthesis, that's very  
2 complicated, right?

3 A. Sure is, yes.

4 Q. Just the physical beauty of a flower is amazing?

5 A. Amazing in a different sense. Of course, when  
6 you're talking about physical beauty, now you're  
7 thinking more of an aesthetic and philosophical concept,  
8 yes.

9 Q. The features seem to be arranged in a way that  
10 gives it great attractiveness?

11 A. Well, okay, but you're now speaking of something  
12 that I was not speaking of. When I talked about the  
13 purposeful arrangement of parts, it was for some  
14 function of the system, not necessarily to be perceived  
15 as pretty.

16 Q. Fair enough. The entire human body, that's an  
17 amazing biological structure?

18 A. I'm thinking of examples.

19 Q. Hopefully, not mine.

20 A. Rest assured. Sure. Yes.

21 Q. We're stipulated here. Because we can make an  
22 agreement about that. The human body, in its entirety,  
23 is an amazing biological system?

24 A. Yes, it's amazing, yes, uh-huh.

25 Q. And just my hand?



1       A.   Yes.

2       Q.   Muscles and joints and bones and nerves.  I can  
3 grab things with it.  I can point.

4       A.   Yes, that is certainly a very impressive  
5 biological system.

6       Q.   Is that a purposeful arrangement of parts?

7       A.   Is it a purposeful arrangement of parts?  Yes, I  
8 think it is.

9       Q.   And the physical world, too, the stars and  
10 planets and gravity, also amazing?

11      A.   They are certainly amazing, yes.

12      Q.   And they function in conjunction with each other  
13 to do things, create gravity, light, things like that,  
14 that are pretty remarkable?

15      A.   Gravity is remarkable.  Light is remarkable.  But  
16 you're going to have to be very careful about the sorts  
17 of conclusions you draw from these things, because --  
18 and simply because you don't want to just become  
19 overenthused about the beauty of nature and try to turn  
20 that into an argument.

21      Q.   But it actually -- I mean, it functions.  Light,  
22 I mean, it functions.  And gravity, it functions?

23      A.   Yes.

24      Q.   And interaction of different elements on the  
25 periodic table combine to make substances in the

1 chemical world, things we rely upon for our life and all  
2 of biological life actually relies on, right?

3 A. Yes, that's certainly true.

4 Q. And we don't rule out natural explanation for all  
5 of these amazing phenomena, do we?

6 A. Well, you're going -- I don't rule out natural  
7 explanations for anything, including intelligent design.  
8 Intelligent design does not rule out natural  
9 explanations. However, you're going to have to make  
10 some distinctions between how phenomena work and what  
11 phenomena strike many people as somehow ordered to, or  
12 is necessary for specific purposes such as the existence  
13 of life.

14 Q. It's really a definitional issue?

15 A. I'm sorry. What is a definitional issue?

16 Q. You just described it. I mean, you got to be  
17 careful about how we're talking about how everything has  
18 different functions when we're making assessments about  
19 whether the natural explanations are valid?

20 A. I couldn't --

21 Q. I'll withdraw that, Professor Behe. You made the  
22 claim that scientists who discuss cellular systems are  
23 calling them machines, correct?

24 A. Yes.

25 Q. And you said, they're not comparing them to

1 machines, they're calling them machines?

2 A. Right.

3 Q. One of the scientists you referred to was Dr.  
4 DeRosier?

5 A. Yes.

6 Q. And what you said, what you quoted from his  
7 article was, More so than other motors, the flagellum  
8 resembles a machine designed by a human?

9 A. Yes.

10 Q. So he's not saying, the flagellum is a machine,  
11 he's saying, it resembles a machine?

12 A. No, he's saying, it resembles a machine designed  
13 by a human. There are other machines in the cell that  
14 may not resemble machines designed by humans, but I  
15 think, as many people can see when looking at an  
16 illustration of the bacterial flagellum, this is a  
17 machine that looks like something that a human might  
18 have designed.

19 Q. It looks like it?

20 A. That's what science has to go on; what we can  
21 see, what we can measure, and so on.

22 Q. It resembles it?

23 A. Exactly.

24 Q. Okay. And when you quoted to -- and he's also  
25 saying, you know, other cellular systems don't resemble

1 machines so much, right? More so than other motors, the  
2 flagellum resembles a machine designed by a human?

3 A. He's saying that more other machines in the cell  
4 don't so much resemble machines designed by humans, but  
5 he is certainly not saying that they are not machines,  
6 at least in my reading.

7 And in that issue -- not -- in a previous issue  
8 of Cell, the one that I pointed to earlier, a number of  
9 scientists were discussing molecular machines that do  
10 not resemble things that do not visually resemble  
11 machines that we have in our world.

12 Q. But here he is saying, resembles a machine  
13 designed by a human. That's your point, right?

14 A. That's what' he said.

15 Q. It looks like a machine a human would design?

16 A. It resembles a machine designed by a human, yes.

17 Q. Now the intelligent designer, when he was forming  
18 a bacterial flagellum millions or billions of years ago,  
19 you're not suggesting he was actually modeling his  
20 design after a manmade rotary motor which didn't exist  
21 until the last century?

22 A. I'm sorry. Could you say that again?

23 Q. Yeah. You're talking about things that resemble  
24 machines designed by humans. You're not suggesting that  
25 the intelligent designer, when the -- when he or she or

1 they designed the first bacterial flagellum millions or  
2 billions of years ago, was modeling its design after  
3 manmade rotary motors which didn't exist until the last  
4 century?

5 A. I'm not quite sure how exactly to address this  
6 question. When you're inferring design, you do not ask  
7 yourself whether a designer had some particular, you  
8 know, look in mind. You're asking whether, in the  
9 structure of this system, you see a purposeful  
10 arrangement of parts.

11 And I think, in the case of the bacteria  
12 flagellum, the fact that it does resemble something from  
13 our everyday world is due to the fact that its function  
14 is similar to some things that we find in our everyday  
15 world such as propulsive motors, like outboard motors on  
16 boats, and, therefore, the functional engineering  
17 requirements would be similar for such a machine in the  
18 cell as well as in our everyday world.

19 Q. Another example you gave was, and just to be  
20 clear, Dr. DeRosier is in no way suggesting that his  
21 article has anything to do with intelligent design?

22 A. Not that I know of.

23 Q. Or irreducible complexity?

24 A. Not that I know of.

25 Q. And then you also cited to Bruce Alberts?

1           A.    Yes.

2           Q.    And I think he is or was the head of AAAS?

3           A.    No, he was the head of the National Academy of  
4    Sciences.

5           Q.    Better yet.  And what you quoted from him was,  
6    Why do we call the large protein assemblies that underlie  
7    cell function protein machines?  Precisely because, like  
8    machines invented by humans, these protein assemblies  
9    contain highly coordinated living parts.  He used the  
10   expression, like a machine?

11          A.    Yes, he did.

12          Q.    And I think what we all learned in grade schools,  
13   when you make a comparison, use like, that's called a  
14   simile?

15          A.    It may be, but I think the point that he was  
16   trying to convey is that these things work like the  
17   machines that we have in our everyday world.  And so, in  
18   fact, they are.

19          Q.    Do you watch football, Professor Behe?

20          A.    I do on occasion, yes.

21          Q.    I watched the Notre Dame/USC game last weekend.  
22   It was quite a game?

23                   MR. MUISE:  I might have to interpose an  
24   objection here, Your Honor.

25                   MR. ROTHSCHILD:  I told Mr. Muise his alma

1 mater did themselves proud, despite the final result.

2 BY MR. ROTHSCHILD:

3 Q. And one of the things the announcer said was  
4 about one of the USC offensive linemen is, he's like a  
5 mountain?

6 A. Yes.

7 Q. Now you don't understand it to say, he was made  
8 like a mountain was, not by wind or erosion or physical  
9 processes on land mass?

10 A. No, of course not. People use words like that in  
11 loose senses all the time. But in this particular case,  
12 Dr. Alberts was making a specific comparison to the  
13 physical functioning of these things and liking it to  
14 the physical functioning of machines in our everyday  
15 world.

16 They require a precise arrangement of parts.  
17 They act by transducing energy in order to accomplish  
18 some function and so on.

19 Q. So when the same announcer said, the running back  
20 is like a bulldozer, that was closer?

21 A. No, I think that's silly.

22 Q. I think it is, too, Professor Behe. And you have  
23 never talked to Bruce Alberts about what exactly he  
24 meant when he used the expression, like a machine?

25 A. No, I didn't.

1 Q. That's your interpretation?

2 A. Yes, it is.

3 Q. And that's true for the other articles you cited  
4 about whether biochemical systems are machines as  
5 opposed to being like machines?

6 A. Well, again, I think we're getting into a  
7 semantical distinction -- or just into semantics. If  
8 something acts like a machine, and something has a  
9 function, and so on, then it is a machine.

10 Q. Now you talked at some length on Monday about the  
11 issue of whether the type III secretory system might be  
12 a pre-cursor to the bacterial flagellum, or the reverse,  
13 that it is a descendent of the bacterial flagellum, or  
14 they might have been a common ancestor, right? You  
15 looked at some articles on that subject?

16 A. Yes.

17 Q. The papers that were discussing that, they were  
18 all discussing this complicated issue within the  
19 framework of evolution, correct?

20 A. Sure. Evolution understood as common descent,  
21 yes.

22 Q. None were suggesting intelligent design?

23 A. No, they did not.

24 Q. They were just scientists trying to figure out  
25 whether it was A that evolved into B, or B that evolved



1 into A, or A and B evolving from C?

2 A. That's right. They were taking the mechanism of  
3 natural selection and random mutation for granted. They  
4 were not demonstrating it. They were not making  
5 arguments for it. They were taking it as an assumption.

6 Q. And in terms of what the order is, they have --  
7 they haven't nailed it down yet, right?

8 A. Not only haven't they nailed it down, but they  
9 have proposed completely opposite scenarios whereby one  
10 can't tell which arose first or second or even if they  
11 arose from each other at all.

12 Q. And you don't expect the dialogue to stop there,  
13 do you?

14 A. I don't expect it to, but it may.

15 Q. Okay. But scientists, as they do with many  
16 subjects on which there's disagreement, may continue to  
17 be making arguments and writing papers and submitting  
18 them to peer review journals and doing experiments to  
19 see if they can come up with a consensus answer on the  
20 subject?

21 A. Sure. And they may write books to try to come up  
22 with an answer, too, as well.

23 Q. That's how you get the royalties, right?

24 A. (No response.)

25 Q. You recently visited the University of Minnesota,

1 didn't you?

2 A. Yes.

3 Q. You spoke with a University Professor named James  
4 Kurzinger?

5 A. Yes, I did.

6 Q. He actually asked you whether the type III  
7 secretory system is a subset of the bacterial flagellum,  
8 is that right?

9 A. I don't think he said exactly that, but I'm not  
10 -- we did talk about the flagellum and the type III  
11 secretory system, but I'm not prepared to say exactly  
12 how the conversation went.

13 MR. ROTHSCHILD: May I approach the witness,  
14 Your Honor?

15 THE COURT: You may.

16 BY MR. ROTHSCHILD:

17 Q. And James Kurzinger is a scientist?

18 A. He identified himself as such.

19 Q. And this is -- this Exhibit 724 is an article in  
20 the Minnesota Daily. It's an opinion piece. And it  
21 says, Intelligent Design 101, Short on Science, Long on  
22 Snake Oil. And it goes on to describe --

23 MR. MUISE: I'm objecting that his use of  
24 this document again is hearsay. He doesn't have  
25 recollection of this, of this conversation. I'm not

1 sure if he's going to be using this to try to refresh  
2 his recollection.

3 MR. ROTHSCHILD: It recounts a conversation,  
4 and I am going to ask Professor Behe whether that  
5 conversation occurred.

6 MR. MUISE: He's going to ask him the  
7 conversation, Your Honor, he can't just read --

8 THE COURT: Well, to the extent that you're  
9 going to try to characterize the -- I think you've  
10 appropriately characterized what the exhibit is, Mr.  
11 Rothschild. So why don't you move on to your question.

12 MR. ROTHSCHILD: Okay. He has expressed a  
13 vague recollection of what happened, so I'm going to  
14 read him the passages in here.

15 THE COURT: I understand.

16 MR. ROTHSCHILD: Okay.

17 THE COURT: I understand. I think the  
18 objection went to the fact that you were beginning to  
19 read or extensively characterize --

20 MR. ROTHSCHILD: Fair enough.

21 THE COURT: -- the exhibit.

22 BY MR. ROTHSCHILD:

23 Q. Just for some more foundation. In the first  
24 paragraph, it says, Intelligent design's leading  
25 scientist, Dr. Behe, a professor of biochemistry,

1 visited the U, which I understand to be the University  
2 of Minnesota, last week as a guest of the McLauren  
3 Institute, and that, in fact, did occur?

4 A. Yes, I visited Minnesota as a guest of the  
5 McLauren Institute.

6 Q. And if you could turn to the third page of the  
7 document. And there's some discussion on that third  
8 page about the bacterial flagellum and the type III  
9 secretory system?

10 A. Yes.

11 Q. And Mr. Kurzinger makes his own observation about  
12 the type III secretory system being a subset of the  
13 bacterial flagellum?

14 A. I'm sorry. Could you say that again?

15 Q. In the paragraph that begins, much to Dr. Behe's  
16 distress --

17 MR. MUISE: Objection, Your Honor, that's  
18 hearsay. He's pointing to a paragraph for the truth of  
19 what's in the statement.

20 THE COURT: Well, it's sustained to the  
21 extent that you're going to read it. He can read it and  
22 put it into context.

23 BY MR. ROTHSCHILD:

24 Q. Could you read the paragraph that says, much to  
25 Dr. Behe's distress?

1           A. Out loud, or --

2           Q. Please.

3           A. Okay. This paragraph says, Much to Dr. Behe's  
4 distress, the TTSS is a subset of the bacterial  
5 flagellum. That's right, a part of the supposedly  
6 irreducible bacterial outboard motor has a biological  
7 function.

8           Q. And I'm not going to ask you about whether you  
9 were distressed or not. But the next paragraph then  
10 says that he asked you about this at lunch, correct?

11          A. That's what it says, yes.

12          Q. And you did have lunch that day?

13          A. We had lunch, and I recall a conversation about  
14 this, but again, I don't recall many details.

15          Q. Okay. And according to Dr. Kurzinger, you  
16 acknowledged that the claim that --

17                   MR. MUISE: Objection, Your Honor. He's  
18 referring to an editorial, and he's trying to recount  
19 this as an exact conversation. Dr. Behe doesn't have  
20 recollection of what occurred. This article has no  
21 relevance.

22                   THE COURT: The next paragraph starting  
23 with, when I asked Dr. Behe, I think, is where you're  
24 going.

25                   MR. ROTHSCHILD: Yes.

1                   THE COURT: Why don't you go right to that,  
2 as it's expressed there, instead of trying to paraphrase  
3 it.

4 BY MR. ROTHSCHILD:

5           Q. It says, When I asked Dr. Behe about this at  
6 lunch, he got a bit testy, but acknowledged that the  
7 claim is correct. Paren, I have witnesses. He added  
8 that the bacterial flagellum is still irreducibly  
9 complex in the sense that the subset does not function  
10 as a flagellum.

11                   My question here is, is Mr. -- Dr. Kurzinger's  
12 account that you agreed that the claim that the TTSS is  
13 a subset of the bacterial flagellum, did you agree to  
14 that?

15           A. I don't recall, but I would, if I was going to  
16 answer it very carefully, I would make a lot of  
17 distinctions before saying so.

18           Q. Okay. But you don't recall whether you said that  
19 or not?

20           A. No, I don't.

21           Q. Okay. And then you go on to say that you still  
22 think -- well, I'll leave that. Your argument is that,  
23 even if the type III secretory system is a pre-cursor to  
24 the bacterial flagellum, is a subset, the bacterial  
25 flagellum is still irreducibly complex because that

1 subset does not function as a flagellum?

2 A. That's correct, yes.

3 Q. And, therefore, the bacterial flagellum must have  
4 been intelligently designed?

5 A. Well, again, the argument is that, there is --  
6 that when you see a purposeful arrangement of parts,  
7 that bespeaks design, so, yes.

8 Q. And yesterday, you testified that, that doesn't  
9 mean the bacterial flagellum was necessarily designed,  
10 appeared abruptly in one fell swoop, correct?

11 A. That's correct.

12 Q. Could have been designed slowly?

13 A. That's correct.

14 Q. So under this scenario, at some period of time,  
15 the bacterial flagellum wouldn't have had all of its  
16 parts until the design was completed?

17 A. Could you say that one more time?

18 Q. Yeah. Under this scenario of slow design --  
19 which was what I experienced with my kitchen -- at some  
20 period of time, the bacterial flagellum wouldn't have  
21 had all its parts until the design was completed?

22 A. That's right.

23 Q. And so without all its parts, it wouldn't be  
24 functional?

25 A. That's right. Not as a flagellum, yes.

1 Q. So that is a phenomenon in both intelligent  
2 design and natural selection?

3 A. I'm not quite sure what you mean.

4 Q. In slow design, the bacterial flagellum has some  
5 prior existence, it doesn't have all its parts, right?

6 A. Well, if -- until it has all its parts and it  
7 starts functioning, I guess it's problematic to call it  
8 a flagellum.

9 Q. It has some subset?

10 A. I guess things that will eventually be part of  
11 the flagellum would begin to appear, yes.

12 Q. Just not function like a flagellum?

13 A. Yes, the system would not yet function as a  
14 flagellum.

15 Q. Just like has been suggested for natural  
16 selection?

17 A. I'm sorry.

18 Q. Just like has been suggested for natural  
19 selection?

20 A. I'm not quite sure what you mean.

21 Q. Natural selection also suggests that there was a  
22 subset of parts that would eventually comprise the  
23 bacterial flagellum, but didn't work as the bacterial  
24 flagellum?

25 A. No. Natural selection, if I remember your



1 question correctly, natural selection does not suggest  
2 that. People see that there is a subset of proteins in  
3 the flagellum which share a lot of sequencology with  
4 proteins that act as a type III secretory system.

5 Nobody, nobody has said how natural selection  
6 could get you the type III secretory system, the  
7 flagellum could get you from the -- even if you had the  
8 type III secretory system, nobody has said how you could  
9 get from that to the flagellum. Nobody has said how you  
10 could get from the flagellum to the type III secretory  
11 system.

12 So this is an example again of conflating  
13 different levels of evolution. We see evidence for  
14 common descent, evidence for relationship, but we see  
15 nothing, nothing that bears on the question of random  
16 mutation and natural selection.

17 Q. Let me see if I've got this right. In natural  
18 selection, the argument is that, there was a subset of  
19 parts, right, like the type III secretory system, that  
20 eventually evolved to become the bacterial flagellum,  
21 right? That's the argument?

22 A. I would want more detail. Are you saying that  
23 in --

24 Q. I'm not asking you to agree with the argument,  
25 Professor Behe. I'm just trying to walk us through

1 this. The argument for the evolution of something like  
2 the bacterial flagellum, just to use that as an example,  
3 is that, at sometime it had a subset of proteins, maybe  
4 looking something like the type III secretory system,  
5 and eventually it evolved to become the bacterial  
6 flagellum? That's the argument, right?

7 A. I would have to see the argument written down.  
8 As you characterize it, I'm not quite sure what it is.

9 Q. Okay. But you're not disputing that the theory  
10 of evolution says, at some point we had a subset of  
11 proteins, then we had eventually all the proteins that  
12 make up whatever system we're discussing?

13 A. That sounds okay.

14 Q. Good. In slow design, same thing. At some  
15 point, we had a subset of the proteins, and eventually,  
16 we got to the whole thing?

17 A. That's right. The crucial question -- the only  
18 question is the mechanism.

19 Q. Okay. So in the case of evolution, there is a  
20 mechanism that's been proposed, natural selection?

21 A. Yes.

22 Q. And you've agreed that natural selection  
23 certainly is a phenomena that operates in the natural  
24 world?

25 A. That is correct.

1 Q. Including at the biochemical level?

2 A. That's right.

3 Q. Then we've got slow design, and there we have no  
4 mechanism at all, no description of a mechanism?

5 A. We have no description of a mechanism. We do  
6 infer design though from the purposeful arrangement of  
7 parts.

8 Q. Now yesterday, I asked you some questions about  
9 the designer's abilities. And you said, all we know  
10 about its abilities is that it was capable of making  
11 whatever we have determined is design. That's the only  
12 statement we can make about the designer's abilities?

13 A. Yes.

14 Q. And in terms of the designer's -- as a scientific  
15 statement?

16 A. That's correct.

17 Q. And the only thing we know scientifically about  
18 the designer's motives or desires or needs is that,  
19 according to your argument, the only thing we would know  
20 scientifically about that is that it must have wanted to  
21 make what we have concluded as design?

22 A. Yes, that's right.

23 Q. In fact, the only way we can make the statement  
24 scientifically that a designer exists is that it made  
25 whatever we conclude was design?

1       A. Yes, that's right.

2       Q. I want to ask you exactly, and this question is  
3 particularly about how -- about the flagellum design.  
4 Was the design limited to the original blueprint for the  
5 first bacterial flagellum?

6       A. I'm not sure what you mean by the blueprint for  
7 the flagellum.

8       Q. The plan?

9       A. The plan? Did the plan cause the flagellum to  
10 occur?

11      Q. Is that all of intelligent design? The designer  
12 planned the bacterial flagellum?

13      A. Well, no. The designer would also have to  
14 somehow cause the plan to, you know, go into effect.

15      Q. It would have to make the thing?

16      A. No, it had to -- well, it would have to have  
17 processes by which it would be made.

18      Q. I mean, it's got to actually be constructed.  
19 We're not talking about a bacterial flagellum in the  
20 mind's eye of the designer. It's actually something we  
21 now know physically exists?

22      A. That's right.

23      Q. Had to be created?

24      A. Well, you're using -- in what sense are you using  
25 the word created? Created can mean -- can have several

1 different senses.

2 Q. You're uncomfortable about that word?

3 A. Yes, because it's a loaded word in these  
4 circumstances.

5 Q. Okay. Created can mean the same thing as made,  
6 right?

7 A. We use the word create when we refer to things  
8 that are made by artists and engineers and so on, yes.

9 Q. Okay. In that sense, the designer created the  
10 bacterial flagellum?

11 A. I might say that, it might be a very indirect  
12 process by which such a thing was made. So when you say  
13 that the designer made the flagellum, it is not  
14 necessary to think that somehow the protein parts of  
15 this were somehow immediately brought together. It  
16 might have been a long process.

17 Q. Did the intelligent designer design each and  
18 every protein of the flagellum?

19 A. That is a difficult question to address, and  
20 there's lots and lots of distinctions to make. When you  
21 ask whether the parts of the flagellum themselves  
22 require design, you have to then focus in on those  
23 parts.

24 As I tried to emphasize earlier in my testimony  
25 when we talk about parts, some people have a simple

1 view, picture in their minds something simple, but each  
2 of the parts is itself a very complicated molecular  
3 entity. And as my work with David Snoke shows, that  
4 even getting small changes in pre-existing proteins,  
5 that is parts, is no easy task. So the question --

6 Q. Unless you have a whole ton of soil?

7 A. I'm sorry?

8 Q. Unless you have a whole ton of soil?

9 A. So that's actually an excellent question. Did  
10 those parts themselves also have to be designed? And I  
11 think right now, the question is open.

12 Q. Did the intelligent designer identify -- design  
13 every individual flagellum in every bacteria or just the  
14 first lucky one?

15 A. Well, since organisms, biological organisms can  
16 reproduce, of course, then if one has the genes and the  
17 proteins and information for a flagellum, then by the  
18 normal processes of biological reproduction, more copies  
19 of the -- of that structure can occur.

20 Q. So the answer is, just the first one?

21 A. That's all that would be needed. That's all we  
22 can infer, yes.

23 Q. Now you have this first flagellum, first bacteria  
24 that has a flagellum. And that has -- those -- that  
25 bacteria with flagellums have had mutations in their

1 flagellums?

2 A. Sure. Genes undergo mutations, yes.

3 Q. And did the designer also design every mutation  
4 of the flagellum since its inception?

5 A. No, you can't -- you certainly can't say that.  
6 There is certainly random processes that go on in our  
7 world, or for processes, that for all we can tell,  
8 certainly appear to be random. So there's no -- nothing  
9 that requires us to think that any mutation, any change  
10 that subsequently occurs to this structure either was  
11 intended or -- was intended.

12 Q. Is that a no or an I don't know?

13 A. Can you restate the question?

14 Q. I asked you the question, did the designer design  
15 every mutation of the flagellum since the first one?  
16 And I'm asking you whether the answer is no or, better  
17 phrase, we don't know?

18 A. Well, that's -- that's a very tricky question.  
19 But the proper answer is that, we don't know.

20 Q. Is the information necessary to answer that  
21 question observable?

22 A. The question of whether the designer designed  
23 every single mutation?

24 Q. Since that first lucky flagellum?

25 A. Is it observable? Hum. We can certainly observe

1 mutations, but unless the mutations and changes and so  
2 on further go on to form a purposeful arrangement of  
3 parts, then we cannot deduce simply from their  
4 occurrence that they were designed.

5 Q. There could be multiple designers, correct?

6 A. Yes, I wrote that in Darwin's Black Box.

7 Q. Could even be competing designers?

8 A. That's correct.

9 Q. Are you aware of any irreducibly complex systems  
10 that have just come into existence in the last five  
11 years?

12 A. Biological systems or mechanical systems or in  
13 our everyday world or other ones?

14 Q. No, Professor Behe, biological systems?

15 A. The last five years? You mean, brand new  
16 irreducibly complex systems?

17 Q. Yes.

18 A. I'm sorry. Brand new ones, not ones that are  
19 just --

20 Q. That are still around, that's right?

21 A. -- reproduced? Not that I'm aware of, no.

22 Q. Last 10 years?

23 A. No.

24 Q. 50 years?

25 A. Not that I know of, no.



1 Q. A hundred years?

2 A. All of the structures that I wrote about in  
3 Darwin's Black Box and have considered are much older  
4 than that.

5 Q. So scientifically, we can't even make -- we can't  
6 even state right now that an intelligent designer still  
7 exists, correct?

8 A. That's correct, yes.

9 Q. Is that what you want taught to high school  
10 students?

11 A. What are you referring to by that?

12 Q. That scientific -- after teaching them about  
13 intelligent design, sign -- and telling them that, that  
14 is a scientific proposition, that right now,  
15 scientifically, we can't even tell you that an  
16 intelligent designer exists? Is that what you want  
17 taught to high school students?

18 A. Well, let's make a couple distinctions. First of  
19 all, when I say, when you use the word taught, again, a  
20 lot of people have in mind instructing students that  
21 this is correct.

22 Q. That's not what I mean, Professor Behe.

23 A. Well, I'm sorry. I was unable to figure out  
24 exactly what you meant. If you're asking --

25 Q. Tell them about it, Professor Behe. Make them

1 aware. Give them information.

2 A. Make them aware that some people say that, from  
3 the purposeful arrangement of parts, we can conclude  
4 that something was designed, but many other questions we  
5 can't determine, including whether there were multiple  
6 designers, whether the designer is natural or not,  
7 whether the designer still exist? Yes, I think that  
8 would be a terrific thing to point out to students.

9 It shows the limitations of theories. It shows  
10 that some evidence bears on one topic, but does not bear  
11 on others. I think that would be terrific pedagogy.

12 Q. Right. Okay. You've taken the position in this  
13 courtroom that intelligent design is open to direct  
14 experimental rebuttal, correct?

15 A. Yes.

16 Q. And you stated that very clearly in your article  
17 Reply to my Critics?

18 A. Yes.

19 Q. And the way you said this could be done, and why  
20 don't we turn to that document, which is Exhibit 718.  
21 If you could turn to page 697. Matt, if you could  
22 highlight in the second paragraph the passage that  
23 starts, To falsify such a claim, and go to the bottom of  
24 the paragraph.

25 And you're asking the question here, or stating,

1 intelligent design is open to direct experimental  
2 rebuttal, correct?

3 A. Yes.

4 Q. And you said, To falsify such a claim, a  
5 scientist could go into the laboratory, place a  
6 bacterial species lacking a flagellum under some  
7 selective pressure, for mobility, say, grow it for  
8 10,000 generations, and see if a flagellum, or any  
9 equally complex system, was produced.

10 If that happened, my claims would be neatly  
11 disproven. Now the test you've described, that would  
12 falsify the claim, your claim that the bacterial  
13 flagellum is irreducibly complex in the way you've  
14 described it, and could, in fact, evolve from  
15 pre-cursors, right, if that was successful?

16 A. That would show that my claim that it required  
17 design -- required intelligent design was incorrect.

18 Q. Let's break that down. You have this concept of  
19 irreducible complexity, right?

20 A. Yes.

21 Q. And you stated that the bacterial flagellum is  
22 irreducibly complex, right?

23 A. That's correct.

24 Q. And this test would, if it was successful,  
25 demonstrate that the bacterial flagellum is not

1 irreducibly complex. We can, in fact, put a bacterial  
2 species lacking a flagellum under some selective  
3 pressure, and eventually it's going to get that  
4 flagellum, right?

5 A. Well, just a distinction. It wouldn't  
6 demonstrate that it wasn't irreducibly complex. It  
7 would demonstrate though that random mutation and  
8 natural selection could produce irreducibly complex  
9 systems.

10 Q. Fair enough. It could evolve, and that would  
11 falsify your claim that an irreducibly complex system,  
12 like a bacterial flagellum, could not evolve through  
13 random mutation and natural selection?

14 A. That's right, yes.

15 Q. But that claim that an irreducibly complex system  
16 cannot evolve through random mutation and natural  
17 selection, that's not your whole case for intelligent  
18 design, correct?

19 A. That's right, it's the purposeful arrangement of  
20 parts.

21 Q. And we saw that bacterial flagellum, right? It's  
22 -- I say, it looks like a machine. You say, it is a  
23 machine. Right?

24 A. Yes.

25 Q. And it sure works like one?

1       A. Yes.

2       Q. So it's got a purposeful arrangement of parts  
3 whether it's irreducibly complex or not?

4       A. It is irreducibly complex. The question is  
5 whether an irreducibly complex system can be put  
6 together by random mutation and natural selection.

7       Q. Okay. So my question is, how would you falsify  
8 the claim that a biological system, like the bacterial  
9 flagellum, which is clearly a purposeful arrangement of  
10 parts, is not intelligently designed?

11      A. Well, since it's an inductive argument, since the  
12 purposeful arrangement of parts is an inductive  
13 argument, then in order to falsify an induction, you  
14 have to find an exception to the inductive argument.

15             So if somebody said that, when you see this  
16 purposeful arrangement of parts -- and again, the -- as  
17 I stress, the argument is quantitative, when there is a  
18 certain degree of complexity and so on. If it was shown  
19 that that did not always, did not always bespeak design,  
20 then the induction would not be reliable, and we would  
21 -- so -- and the argument would be, would be defeated.

22      Q. Now you, in fact, have stated that intelligent  
23 design can never be ruled out, correct?

24      A. Yes, that's right.

25      Q. Now let's turn to your test here of whether

1 bacterial flagellum could evolve through random mutation  
2 and natural selection. 10,000 generations, that's your  
3 proposal, correct?

4 A. Right.

5 Q. And it sounds like a lot, but you actually  
6 testified that, that would just take a couple of years,  
7 right?

8 A. Right.

9 Q. And, you know, based on your understanding of  
10 normal laboratory procedures, even the best  
11 laboratories, how much bacteria would be made a part of  
12 that test?

13 A. Oh, probably at the best, 10 to the 10th, 10 to  
14 the 12th, at the outside.

15 Q. Now you haven't tested intelligent design  
16 yourself this way, have you?

17 A. No, I have not.

18 Q. And nobody in the intelligent design movement  
19 has?

20 A. That's correct.

21 Q. And nobody else has?

22 A. I'm sorry?

23 Q. And nobody else has, outside the intelligent  
24 design movement?

25 A. Well, I'm not sure -- I don't think I would agree

1 with that. I think the experiments described by Barry  
2 Hall were actually in an attempt to do exactly that. He  
3 wanted to see if he could, in his laboratory, re-evolve  
4 a lac operon. His first step in that process in the mid  
5 1970's were the experiments that I discussed here  
6 yesterday, knocking out the beta galactosidase gene.

7 His intention was, from things he has written  
8 later, was to see how that would evolve and then knock  
9 out two steps at a time, and eventually see how he could  
10 get really the whole functioning system. But he had  
11 such trouble with just getting that one step to go, and  
12 since he could not knock out anything else, and get it  
13 to re-evolve, he gave up.

14 And so I would count his efforts as a test of  
15 that, and say that the test, you know, that it was, it  
16 did not falsify intelligent design thinking.

17 Q. And I had actually made a blood pact with my  
18 co-counsel not to ask you about the lac operon, but now  
19 I had to violate it.

20 A. Too late.

21 Q. How many years has he done this experiment?

22 A. I think he was working on it for 20 years or so.

23 Q. In any event, that's the lac operon. But for  
24 bacterial flagellum, you're not aware of that test being  
25 done?

1       A.   No.

2       Q.   Certainly not by anybody in the intelligent  
3 design movement?

4       A.   No.

5       Q.   Okay.  So you can't claim that the proposition  
6 that the bacterial flagellum was intelligently designed  
7 is a well-tested proposition?

8       A.   Yes, you can, I'm afraid.  It's well-tested from  
9 the inductive argument.  We can, from our inductive  
10 understanding of whenever we see something that has a  
11 large number of parts, which interacts to fulfill some  
12 function, when we see a purposeful arrangement of parts,  
13 we have always found that to be design.

14               And so, an inductive argument relies on the  
15 validity of the previous instances of what you're  
16 inducing.  So I would say that, that is tested.

17       Q.   Professor Behe, you say right here, here is the  
18 test, here is the test that science should do, grow the  
19 bacterial flagellum in the laboratory.  And that hasn't  
20 been done, correct?

21       A.   That has not been done.  I was advising people  
22 who are skeptical of the induction that, if they want to  
23 essentially come up with persuasive evidence that, in  
24 fact, an alternative process to an intelligent one could  
25 produce the flagellum, then that's what they should do.



1       Q.   So all those other scientists should do that, but  
2   you're not going to?

3       A.   Well, I think I'm persuaded by the evidence that  
4   I cite in my book, that this is a good explanation and  
5   that spending a lot of effort in trying to show how  
6   random mutation and natural selection could produce  
7   complex systems, like Barry Hall tried to do, is likely  
8   to result -- is not real likely to be fruitful, as his  
9   results were not fruitful. So, no, I don't do that in  
10   order to spend my time on other things.

11       Q.   Waste of time for Barry Hall?

12       A.   I'm sorry?

13       Q.   Waste of time for Barrie Hall?

14       A.   No, certainly not a waste of time. It was very  
15   interesting. He thought that he would learn things.  
16   And he did learn things. But they weren't the things  
17   that he started out to learn. He thought that he would  
18   be able to see the evolution of a complex system. And  
19   he learned how difficult that was.

20       Q.   In any event, you have not undertaken the kind of  
21   test you describe here for any of the irreducibly  
22   complex systems you have identified?

23       A.   I have not.

24       Q.   And neither has anybody else in the intelligent  
25   design movement?

1       A. That's -- well, actually, I think some people are  
2 testing, not the bacterial flagellum, but are testing  
3 other things on protein structure, which I would  
4 probably count under that.

5       Q. Count as irreducibly complex systems?

6       A. Well, I wouldn't really call them irreducibly  
7 complex in that sense, but I think bear on the question.

8       Q. Okay. So in terms of irreducibly complex  
9 structures, you haven't done any tests, right?

10      A. That's right.

11      Q. You're not planning on any tests --

12      A. That's right.

13      Q. -- of the type you described here?

14      A. Well, I'm doing my theoretical work with David  
15 Snoke and hope to continue that, so I think that bears  
16 on this question.

17      Q. Bears on it, but it's not testing an irreducibly  
18 complex system in the way you described in this article?

19      A. That's right.

20      Q. And nobody else, you're not aware of anybody else  
21 in the intelligent design movement doing a test of the  
22 type you described here of an irreducibly complex  
23 system?

24      A. No, not yet.

25      Q. Now you talked about how, you know, your proposal

1 here would take approximately two years, right?

2 A. Yes, yes.

3 Q. I'm sorry. I'm pointing to down here, and that's  
4 -- you're not that good a mind reader. Now bacteria had  
5 been on the Earth for billions of years, correct?

6 A. That's right.

7 Q. And the bacterial population that exists in the  
8 world and has ever existed in the world is orders and  
9 orders of magnitude greater than ever could be in one  
10 laboratory experiment?

11 A. That's right. It should be about 10 to the 40th  
12 or so, I would estimate.

13 Q. And I think you said, 10 to the -- what was your  
14 proposal for the laboratory, 10 to the -- you had said  
15 that you had a suggestion for how much we would study in  
16 one laboratory?

17 A. 10 to the 10th and 10 to the 12th, that's  
18 correct.

19 Q. And you talked about selective pressures that the  
20 bacterial flagellum could be exposed to, but a  
21 laboratory could never recreate all the selective  
22 pressures that have existed in the environment for the  
23 last three and a half billion years?

24 A. Well, that's certainly true. But a scientist --  
25 scientists nonetheless try to understand parts of

1 nature, even though nature is very much bigger than a  
2 laboratory. And in many other instances, such as people  
3 investigating origin of life and so on, they nonetheless  
4 try to understand what the proper environment would be  
5 to study, and so they can kind of focus their efforts on  
6 what would be the most promising type of environment,  
7 and so make it more likely to discover something that  
8 was there than just focusing on the whole world.

9 Q. But it's entirely possible that something that  
10 couldn't be produced in the laboratory in two years, or  
11 a hundred years, or even in the laboratory that was in  
12 operation through all of human existence, could be  
13 produced over three and a half billion years? You have  
14 to agree with that, Professor Behe?

15 A. It's entirely possible, but we can only know if  
16 that is the case if we have, if we have experiments to  
17 back it up or calculations to back it up.

18 Q. Experiments and inferences, right?

19 A. That's right.

20 Q. And so you agree, something we couldn't -- that  
21 couldn't happen in two years, much better chance over  
22 three and a half billion years?

23 A. Absolutely.

24 Q. Okay. And that's why the age of the earth is so  
25 important to a scientific theory about biological life,

1 isn't it, Professor Behe?

2 A. It's very important.

3 Q. But intelligent design, that's a who cares,  
4 right? It could be -- the universe could be -- or the  
5 Earth could be billions of years old or 10,000 years  
6 old, and it doesn't matter to intelligent design?

7 A. Intelligent design is not a person, so it doesn't  
8 have feelings like you are describing.

9 Q. It's a movement, right?

10 A. Intelligent design is a scientific theory that  
11 focuses on a particular question. There are many  
12 scientific theories that focus on particular questions  
13 that do not have anything to do with other interesting  
14 questions. The scientific theory of intelligent design  
15 focuses on discerning design, and that's it.

16 Q. Okay. So it doesn't take a position on the age  
17 of the Earth?

18 A. Theories don't take positions.

19 Q. Okay. The intelligent design -- you described  
20 intelligent design as not making any claims about the  
21 age of the Earth, correct?

22 A. That's correct.

23 Q. And, of course, the prospects for evolution of a  
24 function or a system are also greater if the subject  
25 population is greater?

1       A.   That's correct.

2       Q.   And no human laboratory can duplicate the entire  
3 population of any kind of organism, correct?

4       A.   That's correct.

5       Q.   Okay.  And no human laboratory can duplicate all  
6 of the selective pressures that have existed in the  
7 billions of years that bacteria have been around?

8       A.   That's correct.  So we can't rule out all  
9 explanations.  We have to investigate to see what are  
10 likely.

11      Q.   Professor Behe, the tests you proposed here  
12 regarding the bacterial flagellum is like asking Dr.  
13 Padian to grow a bird wing in a laboratory, isn't it?

14      A.   The test that is sufficient for a theory is  
15 proportional to what the theory claims.  I'm no  
16 physicist, but in physics, there have been claims, many  
17 claims that required enormous amounts of effort by the  
18 entire physical community to build large structures,  
19 took many years to do so.

20           And nonetheless, they thought that this effort  
21 was worth it, because they wanted to be sure of the  
22 answer.  In biology, the claim that random mutation and  
23 natural selection can produce systems like the flagellum  
24 or other molecular machines is a very large claim.  And  
25 one can't simply say that because it would be hard to

1 test it, we will just assume it's true.

2           So if somebody wants to be sure or somebody wants  
3 to -- wants to -- wants to respond to a skeptic with  
4 evidence that would convince somebody that was not  
5 already convinced of the theory, then there is no  
6 escaping the fact that you have to show that your theory  
7 can do what you claim for it.

8           Q. And so to do that, what scientists advocating for  
9 the theory of evolution, including natural selection,  
10 have to do is create a laboratory that repeats human  
11 life -- that contains all of human life in deep time?

12           A. I'm sorry. One more time.

13           Q. In order to validate this big claim that the  
14 theory of evolution makes, what you're really saying is,  
15 they've got to create a laboratory that includes all of  
16 biological life and operates over deep time?

17           A. No, I didn't say that at all. I said, if it can  
18 be demonstrated that random mutation and natural  
19 selection can produce complex systems, then intelligent  
20 design would be falsified. One doesn't have to, you  
21 know, re -- show that something of the complexity of a  
22 flagellum would be made.

23           But if one saw that something somewhat less  
24 complex might be made in a reasonable time, then one  
25 might be able to extrapolate. You'd have to pay

1 attention to the details of the system. So it's not,  
2 you know -- you don't need a worldwide laboratory and a  
3 billion years to test this. You can do things like  
4 Barry Hall tried to do.

5 Q. That can't recreate the opportunities that were  
6 there for biological organisms throughout time?

7 A. There are always opportunities for biological  
8 organisms. Biological organisms compete with each  
9 other. If one manages to compete more successfully, it  
10 will -- it will out grow others. And so there is no  
11 reason we can't expect something, like in Barry Hall's  
12 experiments, to show us some new interesting structure.

13 And if that occurred, that would be a real  
14 feather in the cap of people who think Darwinian theory  
15 is correct.

16 Q. Let's move onto the blood clotting cascade. Now  
17 you showed us some slides yesterday, or the day before,  
18 that show that certain organisms maintain a blood  
19 clotting function with less than all the parts that  
20 mammals have, correct?

21 A. That's correct.

22 Q. Okay. But that's not what you said in the blood  
23 clotting section in Pandas. You said, all the parts  
24 have to be, correct?

25 A. No, I didn't.



1 Q. Let's turn to pages 145 -- page 145 in Pandas,  
2 P-11. And this is the section on blood clotting?

3 A. Page 145?

4 Q. Right.

5 A. This is part of it.

6 Q. Right. And if you could turn to page 146.

7 A. Yes.

8 Q. And, Matt, if you could highlight that top  
9 paragraph, that one that continues over. You say, All  
10 of the proteins had to be present simultaneously for the  
11 blood clotting system to function, right?

12 A. That's right, all the proteins I was talking  
13 about.

14 Q. Okay. And then I understand, on Monday, you were  
15 distinguishing that there are different parts of the  
16 pathway, there are different parts of the pathway?

17 A. Yes.

18 Q. And what you said in -- on Monday is that, some  
19 of those parts, we have a harder time understanding than  
20 other parts?

21 A. Right.

22 Q. Okay. And, therefore, you just focus on a subset  
23 of the parts, right?

24 A. Right.

25 Q. Now you've got this whole cascade. You've got a

1 diagram in Pandas. You got a diagram in your book,  
2 Darwin's Black Box. And you show it as a multi-protein  
3 system that includes that -- I think you said, intrinsic  
4 part of the pathway?

5 A. Yes, uh-huh.

6 Q. So that's the whole blood clotting cascade,  
7 correct?

8 A. That's as it's presented in textbooks, yes.

9 Q. And you presented it that way in Darwin's Black  
10 Box?

11 A. Yes, I did. I used that figure, yes.

12 Q. Okay. And you used it that way in Pandas,  
13 correct?

14 A. I used it -- a very similar figure, yes.

15 Q. And one whole system, one whole blood clotting  
16 cascade?

17 A. These are all the proteins that have been  
18 determined to affect blood clotting, yes.

19 Q. Okay. So -- but your claim in court is that, eh,  
20 let's ignore parts of it, some of those parts don't  
21 matter, we're just looking at a subset, right?

22 A. I made proper distinctions about what is required  
23 and about what we don't have sufficient information to  
24 make claims about that, yes.

25 Q. But those other parts never suggested are not

1 part of the blood clotting cascade, right, the intrinsic  
2 pathway?

3 A. Well, I'm afraid I did. I -- well, I quoted a  
4 section of my book showing that I was confining my  
5 argument to the proteins at the end of the pathway.

6 Q. Matt, could you go to page 143 in Pandas so that  
7 we can have the picture of the system. I understand  
8 what you're saying, Professor Behe. You did indeed, in  
9 Darwin's Black Box, define the blood clotting system in  
10 a particular way, right, meaning --

11 A. Yes.

12 Q. And what you called irreducible complex didn't  
13 include, I guess, what's sort of in that top left-hand  
14 corner of the cascade?

15 A. That's correct.

16 Q. But that's not the entire cascade?

17 A. Well, there are many more proteins that affect  
18 blood clotting. But when I was talking about the  
19 concept of irreducible complexity, I wanted to make sure  
20 that we were talking about ones whose function was as  
21 clear as possible, so I limited it to that.

22 Q. You defined the system down more narrowly?

23 A. I'm sorry?

24 Q. You defined the system more narrowly?

25 A. That's right, yes.

1       Q. And so I guess what you're saying is, part of the  
2 system -- part of the blood clotting system that works  
3 in all of our bodies is irreducibly complex, but as it  
4 gets more complicated, it's not irreducibly complex?

5       A. No, I didn't say that. I said that the portion  
6 of the blood clotting system that I was focusing on was  
7 irreducibly complex. There might be components which  
8 affect blood clotting which can or can't be removed and  
9 help or not help but not break the system. But I was  
10 focusing my argument on irreducible complexity on the  
11 proteins I cited in my testimony.

12       Q. You define the system in whatever way is  
13 convenient to the argument?

14       A. I define the system very carefully to make sure  
15 that people understand what I'm talking about. I use  
16 the standard figure of the blood clotting cascade from a  
17 biochemistry textbook, because that's what is understood  
18 as the protein system that affects blood clotting.

19       Q. Now let me just make sure I understand the  
20 argument. What I think you said was, when I looked  
21 at -- the subset of the blood clotting cascade included  
22 fibrinogen, prothrombin, proaccelerin, and activated  
23 Stuart factor. Those are the things you say in Darwin's  
24 Black Box constitute the irreducibly complex system?

25       A. Okay.

1 Q. Is that correct?

2 A. Yes.

3 Q. And could you look on page 145 of Pandas?

4 A. Yes.

5 Q. Okay. And, Matt, could you highlight in the  
6 middle of the first column where it starts, We may try  
7 many smaller sets. You say here, We may try many  
8 smaller sets of components to get started; fibrinogen,  
9 prothrombin, activate the Stuart factor, and  
10 proaccelerin. And then you give some other  
11 alternatives. But then you say, death is nearly always  
12 the certain result, right?

13 A. Yes, I did.

14 Q. Okay. So that's actually saying, those four  
15 parts of the system, if that's all you got, not good  
16 enough?

17 A. Excuse me a second. Let me read this, please.  
18 Yeah, with those four, the system would not work.

19 Q. With those four, the system would not work?

20 A. Yes.

21 Q. Those are the four you just agreed were enough to  
22 make your irreducibly complex system?

23 A. Well, those are the four that I said that, if you  
24 knock them out of the current system, the system would  
25 not function.

1 Q. So here you're saying, just having those four --  
2 you're saying, that's the irreducibly complex system,  
3 and the rest of it we can forget, and now we look at  
4 that irreducibly complex system, and death would be the  
5 certain result?

6 A. I'm -- I'm not -- I'm not -- I'm not  
7 understanding the distinction you're making, sir.

8 Q. Well, we looked at the puffer fish, right?

9 A. Yes.

10 Q. And it was missing some parts of the blood  
11 clotting cascade. But you said, from my argument, that  
12 doesn't matter, because that's not what I'm talking  
13 about, right?

14 A. Yes.

15 Q. You said, what I am talking about is these four  
16 factors here, right? I won't say them again because  
17 I'll just butcher them. Stuart factor and its friends.  
18 You said in your testimony on Monday, those four, those  
19 you need?

20 A. Yes.

21 Q. That's enough. That's irreducibly complex.

22 A. I didn't say, that's enough. I said that we  
23 certainly need those.

24 Q. And now you're saying here, those four, not  
25 enough, they're just -- they're just dead?

1       A. Well, again, I said that they were necessary. I  
2 don't think I said they were sufficient.

3       Q. You didn't identify any other systems?

4       A. Again, I was trying to identify parts which were  
5 certainly necessary, but I don't think I said that I was  
6 describing a minimal system.

7       Q. Could you turn to page 86 in Darwin's Black Box,  
8 and the first continuing paragraph?

9       A. Yes.

10      Q. Okay. And this is the chapter where you're  
11 talking about how the blood clotting cascade is  
12 irreducibly complex?

13      A. Right.

14      Q. And you say, The function of the blood clotting  
15 system is to form a solid barrier at the right time and  
16 place that is able to stop blood flow out of an injured  
17 vessel. The components of the system beyond the fork in  
18 the pathway -- that's the part we don't know so much  
19 about?

20      A. Yes.

21      Q. -- are fibrinogen, prothrombin, Stuart factor,  
22 and proaccelerin, factors that, by themselves, you die  
23 from, right?

24      A. I'm sorry? The factors --

25      Q. The factors that -- it says, The components of

1 the system beyond the fork in the pathway are  
2 fibrinogen, prothrombin, Stuart factor, and  
3 proaccelerin. And those are the factors that, in  
4 Pandas, you say, if that's all you got, you're dead?

5 A. I -- I -- these are the factors which, if you  
6 break them, will cause the clotting system to stop  
7 working.

8 Q. That's the system, right? That's what it says in  
9 Darwin's Black Box? Those four components, that's the  
10 system?

11 A. The total system? Does it say that?

12 Q. It says, the system.

13 A. I'm sorry. Where are you reading from now?

14 Q. Page 86, Professor Behe. We know it's not the  
15 total system. There's a whole lot that we don't know  
16 about, right, and that the puffer fish can do without.  
17 But the system you're talking about, the single system  
18 that's irreducibly complex, that's those four  
19 components, correct?

20 A. No. Again, I said that we should focus our  
21 attention on those, because a lot more is known about  
22 them, and if you remove them, the system will certainly  
23 be broken.

24 Q. Right above what we just read, it says, The blood  
25 clotting system fits the definition of irreducible



1 complexity?

2 A. I'm sorry. Can you tell me exactly where you  
3 are?

4 Q. Yes, the first full sentence on this page.

5 A. That begins, Leaving aside the system before the  
6 fork in the pathway?

7 Q. Yes. Leaving aside the system before the fork in  
8 the pathway, where some details are less well-known, the  
9 blood clotting system fits the definition of irreducible  
10 complexity. So we're leaving aside that stuff before  
11 the fork?

12 A. Okay.

13 Q. We're leaving the stuff aside that we know the  
14 puffer fish can do without. And you're saying, The  
15 blood clotting system fits the definition of irreducible  
16 complexity. That is, it is a single system composed of  
17 several interacting parts that contribute to the basic  
18 function, and where the removal of any one of the parts  
19 causing the system effectively to cease functioning.

20 It talks more about the function. It says, The  
21 components of the system beyond the fork in the pathway  
22 are fibrinogen, prothrombin, Stuart factor, and  
23 proaccelerin. That's your irreducibly complex system,  
24 isn't it, Professor Behe?

25 A. No, it's not. Again, I was confining my

1 discussion to the point after the fork in the pathway  
2 because, as I said in the book, much more is known about  
3 that. But the fork in the pathway is essentially two  
4 different ways to activate the pathway.

5         And while you can do without one way to activate  
6 the pathway, you can't do without both ways to activate  
7 the pathway. Something has to activate it.

8         Q. So you have to have those four, right?

9         A. Yes, those four are needed for the system to  
10 work. But -- and I confined my discussion to them. But  
11 they're not sufficient for a functioning system.

12         Q. You need the stuff before the pathway, too?

13         A. You need some of the stuff, yes.

14         Q. Except for the puffer fish?

15         A. Well, again, like I said, some of the stuff. The  
16 puffer fish itself has the extrinsic pathway, which is  
17 one way to trigger the remaining steps. It's missing  
18 the intrinsic pathway. But nonetheless, it still has  
19 one way to turn the pathway on.

20         Q. It has those four things?

21         A. It does, yes.

22         Q. Which we know, by themselves, cause death?

23         A. By themselves, they would cause the system to  
24 start stop functioning.

25         Q. Sounds like a bigger mistake than Dr. Doolittle

1 made, Professor Behe?

2 A. I'm not sure what you are referring to.

3 Q. Well, you spent a lot of time trashing Dr.  
4 Doolittle and his work, his article in the Boston  
5 Review. Your mistake here is quite a bit more  
6 substantial than misinterpreting a mice study, isn't it?

7 A. I'm not even quite sure what you are referring to  
8 as my mistake.

9 Q. I'll withdraw that question, Professor Behe.  
10 It's surely not your contention that the mistake you  
11 understand Dr. Doolittle to have made basically  
12 invalidates the possibility that the blood clotting  
13 system could have evolved?

14 A. No, of course not. The only point I was making  
15 with that discussion was that he did not know how  
16 Darwinian processes produced it. It was not an argument  
17 saying that -- or it was not -- did not go to the point  
18 of whether or not that could happen.

19 Q. Okay. And that was an article, whether right or  
20 wrong, that was not in a peer reviewed scientific  
21 journal?

22 A. That's correct.

23 Q. Dr. Doolittle, as you showed us, has actually  
24 written quite a bit on the subject of the blood clotting  
25 cascade in peer reviewed scientific journals?

1       A. He certainly has.

2       Q. Including what we saw about the puffer fish?

3       A. That's correct.

4       Q. And by contrast, how many peer reviewed articles  
5 are there explaining the blood clotting -- why the blood  
6 clotting cascade cannot evolve because it is irreducibly  
7 complex in the way you describe?

8       A. Well, I'm going to say that the articles which  
9 elucidate the structure of the blood clotting pathway  
10 are the ones which demonstrate that. I will agree that  
11 there certainly are no arguments or directly to that  
12 point. But as I tried to show in my book, Darwin's  
13 Black Box, that's an implication that can easily be  
14 drawn from those studies.

15       Q. So these are all those other articles based on  
16 the research of other scientists that you interpret  
17 differently than those scientists do?

18       A. That's right. I was proposing a newer idea.

19       Q. Okay. And how many peer reviewed articles are  
20 there in scientific journals discussing the intelligent  
21 design of the blood clotting cascade?

22       A. Well, again, since we infer design by the  
23 purposeful arrangement of parts, then the peer reviewed  
24 articles in science journals that demonstrate that the  
25 blood clotting system is indeed a purposeful arrangement

1 of parts of great complexity and sophistication, there  
2 are probably a large number of those.

3 Q. Again, those are those articles by other  
4 scientists based on experimental research, right?

5 A. They are certainly by other scientists, not by  
6 myself, and they are certainly based on experiments.

7 Q. And none of those articles are arguing that the  
8 blood clotting cascade are intelligently designed -- is  
9 intelligently designed?

10 A. That's correct.

11 Q. And there are no peer reviewed articles arguing  
12 that the blood clotting cascade is intelligently  
13 designed, right, in scientific journals?

14 A. I wrote my argument in a book, so, yes, that's  
15 correct.

16 Q. And before we leave the blood clotting system,  
17 can you just remind the Court the mechanism by which  
18 intelligent design creates the blood clotting system?

19 A. Well, as I mentioned before, intelligent design  
20 does not say, a mechanism, but what it does say is, one  
21 important factor in the production of systems, and that  
22 is that, at some point in the pathway, intelligence was  
23 involved.

24 MR. ROTHSCHILD: This would be a good time  
25 for a break, Your Honor.

1                   THE COURT: All right. Why don't we take  
2 our lunch break at this point, and we will be in recess  
3 until 1:35 this afternoon. We'll resume cross  
4 examination at that time. Thank you.

5                   (Whereupon, a lunch recess was taken at  
6                   12:10 p.m.)

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