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<p style="text-align: center;">UNITED STATES DISTRICT COURT NORTHERN DISTRICT OF CALIFORNIA</p> <p>THE BOARD OF TRUSTEES OF THE LELAND STANFORD JUNIOR UNIVERSITY,</p> <p style="text-align: center;">Plaintiff,</p> <p style="text-align: center;">vs.</p> <p style="text-align: right;">Case Number C-05-04158-MHP</p> <p>ROCHE MOLECULAR SYSTEMS, INC.; ROCHE DIAGNOSTICS CORPORATION; ROCHE DIAGNOSTICS OPERATIONS, INC.; ROCHE DIAGNOSTIC SYSTEMS, INC., Defendants.</p> <hr/> <p style="text-align: center;">HIGHLY CONFIDENTIAL - ATTORNEYS' EYES ONLY DEPOSITION OF ERNEST S. KAWASAKI, PH.D. Washington, D.C. Thursday, August 24, 2006 9:28 a.m.</p> <p>Reported by: Karen Young CSR - NOTARY PUBLIC Job No. 51916</p> <p style="text-align: right;">Page 1</p>	<p style="text-align: center;">1 APPEARANCES</p> <p>2 ON BEHALF OF THE PLAINTIFF:</p> <p>3 MICHELLE S. RHYU, PH.D., ESQUIRE</p> <p>4 COOLEY GODWARD LLP</p> <p>5 Five Palo Alto Square</p> <p>6 3000 El Camino Real</p> <p>7 Palo Alto, California 94306-2155</p> <p>8 rhyums@cooley.com</p> <p>9 (650) 843-5505</p> <p>10</p> <p>11 ON BEHALF OF THE DEFENDANTS:</p> <p>12 ROBERT B. WILSON, ESQUIRE</p> <p>13 QUINN EMANUEL URQUHART OLIVER & HEDGES, LLP</p> <p>14 51 Madison Avenue, 22nd Floor</p> <p>15 New York, New York 10010</p> <p>16 (212) 849-7145</p> <p>17</p> <p>18 JEREMY AARON BURNS, ESQUIRE</p> <p>19 QUINN EMANUEL URQUHART OLIVER & HEDGES, LLP</p> <p>20 555 Twin Dolphin Drive, Suite 560</p> <p>21 Redwood Shores, California 94065</p> <p>22 (650) 801-5017</p> <p>23</p> <p>24</p> <p>25</p> <p style="text-align: right;">Page 3</p>
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<p>1 Deposition of ERNEST S. KAWASAKI, PH.D.,</p> <p>2 held at the offices of:</p> <p>3 COOLEY GODWARD LLP</p> <p>4 The Bowen Building</p> <p>5 875 15th Street, Northwest</p> <p>6 Suite 800</p> <p>7 Washington, D.C. 20005-2221</p> <p>8 (202) 842-7800</p> <p>9</p> <p>10</p> <p>11</p> <p>12 Pursuant to notice, before Karen Young,</p> <p>13 Notary Public of the District of Columbia.</p> <p>14</p> <p>15</p> <p>16</p> <p>17</p> <p>18</p> <p>19</p> <p>20</p> <p>21</p> <p>22</p> <p>23</p> <p>24</p> <p>25</p> <p style="text-align: right;">Page 2</p>	<p style="text-align: center;">1 C O N T E N T S</p> <p>2 EXAMINATION OF ERNEST S. KAWASAKI, PH.D. PAGE</p> <p>3 By Ms. Rhyu 6</p> <p>4 By Mr. Wilson 96</p> <p>5</p> <p>6</p> <p>7</p> <p>8</p> <p>9</p> <p>10</p> <p>11</p> <p>12</p> <p>13</p> <p>14</p> <p>15</p> <p>16</p> <p>17</p> <p>18</p> <p>19</p> <p>20</p> <p>21</p> <p>22</p> <p>23</p> <p>24</p> <p>25</p> <p style="text-align: right;">Page 4</p>
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1 that end of things, so --
 2 Q. Earlier, you mentioned that Dr. Holodniy
 3 must have had some agreement with Merigan's -- through
 4 Merigan? Did I understand you correctly? What was
 5 your testimony relating to an agreement?
 6 MR. WILSON: Objection, vague.
 7 BY MS. RHYU:
 8 Q. Let me withdraw that question. You
 9 mentioned an agreement earlier. Did you ever see any
 10 agreement pertaining to Dr. Holodniy's presence at
 11 Cetus?
 12 A. Can't remember, no.
 13 Q. Did you ever see any agreement between Cetus
 14 and Dr. Merigan?
 15 A. The same thing, I can't remember. I don't
 16 remember.
 17 Q. When you say you can't remember, are you
 18 saying you could have seen such an agreement but you
 19 don't remember?
 20 A. No, I can't remember, I can't remember.
 21 That's all. I can't remember.
 22 Q. Do you have any specific memory of any
 23 conversations you had with Dr. Holodniy?
 24 A. Specific memories? I would say no.
 25 Q. Do you have any general memories of --

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1 Q. Aside from what you said about generally
 2 discussing RT-PCR, do you recall anything else that
 3 you told Dr. Holodniy regarding PCR technology?
 4 A. At this point I can't say exactly. I know
 5 we discussed all these technologies. The exact
 6 technologies, the way to do RT-PCR and that kind of
 7 stuff was done. Now, if you're -- well, that's
 8 basically it.
 9 Q. And what were you about to say?
 10 A. That's why I'm trying to think. I could --
 11 anyway, that's about it. I can't think of anything
 12 else other than that.
 13 Q. Do you recall giving Dr. Holodniy any
 14 reagents?
 15 A. I can't recall exactly, but I probably did.
 16 Q. Why do you say you probably did?
 17 A. Because he was working in the Cetus lab, so
 18 he had to have reagents from Cetus labs, so I assume
 19 everybody provided reagents.
 20 Q. Do you know that he was also working at
 21 Stanford Labs during that time?
 22 A. I knew he was at Stanford. I don't know
 23 exactly what he was doing at Stanford.
 24 Q. So it's possible that he was using reagents
 25 that he obtained at Stanford Labs in his experiments.

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1 A. General memories, yes.
 2 Q. -- discussions that you had with
 3 Dr. Holodniy?
 4 A. Again, a general memory is discussions of
 5 PCR technologies, what we were doing at Cetus, and in
 6 my case, RT-PCR.
 7 Q. What do you remember telling him about
 8 RT-PCR?
 9 A. Well, when I say discussions of RT-PCR
 10 technology, it just means discussing how the system or
 11 protocol works, and these things are all recorded in
 12 my lab book.
 13 Q. Did you tell Dr. Holodniy which experiments
 14 to perform?
 15 A. Again, I couldn't remember if I told him to
 16 do such and such at this point.
 17 Q. Did you supervise Dr. Holodniy's
 18 experiments?
 19 A. Probably "supervised" is too strong a word.
 20 I probably -- I discussed experiments with him back --
 21 but supervising is a different -- a different thing.
 22 Q. And I just want to get to your memory of
 23 what you discussed with him. Aside from --
 24 MR. WILSON: Let her ask the question.
 25 BY MS. RHYU:

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1 MR. WILSON: Objection, asked and answered.
 2 BY MS. RHYU:
 3 Q. Could you give an audible answer please?
 4 A. I can't say whether -- where -- but I'm sure
 5 we provided him with reagents. Whether he used
 6 Stanford reagents too I don't know.
 7 Q. What's your basis for saying that you're
 8 sure that we, I suppose that means Cetus, provided him
 9 reagents?
 10 MR. WILSON: Asked and answered.
 11 A. Are you saying that if I remember exactly
 12 what reagents we gave to him?
 13 Q. I'm just asking if you remember giving him
 14 any reagents.
 15 MR. WILSON: Asked and answered.
 16 A. Since he was working at Cetus, he would most
 17 likely use Cetus reagents because there was work with
 18 Tom Merigan.
 19 Q. You don't particularly remember giving him
 20 any reagents though?
 21 A. No.
 22 Q. And do you know if anyone in your lab gave
 23 him any reagents?
 24 A. I would say no.
 25 Q. Did you give him any equipment?

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1 experiments?
 2 A. I would say --
 3 MR. WILSON: Objection, vague.
 4 A. I couldn't remember something like that.
 5 Q. Do you remember conveying any confidential
 6 information to Dr. Merigan?
 7 MR. WILSON: Calls for a legal conclusion.
 8 A. Can't remember anything like that.
 9 Q. You mentioned earlier that Dr. Holodniy was
 10 in Dr. Merigan's lab. Do you have any knowledge about
 11 whether Dr. Merigan arranged for Dr. Holodniy to work
 12 at Cetus? Do you have any personal knowledge about
 13 that?
 14 A. I can't remember, but he probably did. I
 15 just assumed it.
 16 Q. You assume it, but you have no personal
 17 knowledge about it.
 18 A. I can't -- I can't remember exactly if
 19 someone said he's coming from Merigan's lab, but
 20 someone comes, you find out where they're from.
 21 Q. Do you know Dr. David Schwartz?
 22 A. No, I don't remember him.
 23 Q. How about Dr. David Katzenstein?
 24 A. Same thing. I don't remember them.
 25 Q. And you don't remember giving any reagents

1 A. No, I don't think so. I can't remember if I
 2 already left. What year was this that it came out?
 3 It came out -- anyway, I don't think so.
 4 Q. Sitting here today, do you think you should
 5 have been an author on this publication?
 6 A. No, no.
 7 Q. And you said you reviewed this paper in
 8 preparation for today's deposition. Is there anything
 9 in your review of the paper that you saw that led you
 10 to believe that -- that you had made a contribution to
 11 the work described in this publication?
 12 MR. WILSON: Objection, vague, foundation.
 13 I guess also to the extent that you reviewed this in
 14 connection with preparing for the deposition, don't
 15 reveal any communications with counsel.
 16 THE WITNESS: Uh-huh. So your question was
 17 again?
 18 MS. RHYU: Could you read it back please?
 19 - - -
 20 THE REPORTER: Question: "And you said you
 21 reviewed this paper in preparation for today's
 22 deposition. Is there anything in your review of the
 23 paper that you saw that led you to believe that you
 24 had made a contribution to the work described in this
 25 publication?"

1 to either of them?
 2 A. No.
 3 Q. You don't remember having any discussions
 4 with either of them?
 5 A. At this point I can't remember, no.
 6 Q. I'm handing you what was previously marked
 7 as Exhibit 1. Is this the JID article that you
 8 previously mentioned having reviewed in preparation
 9 for today's deposition?
 10 A. Let's see. Is this the Journal of
 11 Infectious Diseases? Journal of Infectious Disease.
 12 Yes, yes, one I think I was looking at.
 13 Q. When did you first become aware of Exhibit
 14 1?
 15 A. This one? I can't remember, but I probably
 16 remember it when it came out. That was '91.
 17 Q. Did you think it's likely that you saw it
 18 when it first came out?
 19 A. Yes, uh-huh.
 20 Q. And why do you think that?
 21 A. Because I followed a lot of the PCR work
 22 back then, and these things -- and also the AIDS
 23 research.
 24 Q. Do you remember thinking at the time that
 25 you should have been an author on this publication?

1 - - -
 2 MR. WILSON: Same objections and same
 3 caution.
 4 A. In this case, basically it's RT-PCR
 5 technology. That's my contribution, and at the time,
 6 a lot of stuff was happening. RT technology was not a
 7 -- was not a standard thing that everybody could do
 8 easily, so that's the technology, a lot of technology
 9 was developed at Cetus.
 10 Q. And can you point me specifically to what
 11 you're referring to when you say the RT technology?
 12 A. Well, if we look at all the primers that
 13 were used, most of them as far as I can tell just by
 14 glancing at it are Cetus design.
 15 Q. I'm asking about your personal contribution.
 16 A. RT-PCR?
 17 Q. Yes.
 18 A. It's a thing that he -- I'm pretty sure a
 19 lot of the technology was learned at Cetus, and he
 20 refers to a chapter in a book.
 21 Q. So are you referring to, on the first page
 22 of Exhibit 1, second paragraph, RNA extraction,
 23 reverse transcription and amplification of cDNA? Are
 24 you referring to that paragraph?
 25 A. Yes.

1 Q. And the first sentence of that paragraph
2 states, "Total RNA from 200 microliters of serum was
3 extracted using guanidinium thiocyanate and reverse
4 transcribed with MMLD reverse transcriptase by methods
5 previously described." Is that what you're referring
6 to?
7 A. Yes, that's partly it.
8 Q. That's partly it? What more -- what more
9 are you referring to as your contribution to this
10 paper?
11 A. The contribution -- basically the
12 contribution is how to -- the technology of RT-PCR.
13 When you say the contribution, that's what Cetus'
14 contribution was to the technology.
15 Q. And the RT-PCR is the PCR that's described
16 in these references that are cited to here, 7 and 8?
17 A. That's one reference out of a lot of the
18 technology that Cetus had at the time. That doesn't
19 explain everything. The reference is one reference.
20 Q. Is there anything specific in this article
21 that you can say you contributed to the article?
22 MR. WILSON: Objection, asked and answered
23 multiple times.
24 A. I've answered already.
25 Q. So everything that you say -- you've

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1 testified to is all that you can remember that you've
2 contributed to this article?
3 MR. WILSON: Asked and answered.
4 A. I would say yes.
5 Q. I'm handing you what's been marked as
6 Exhibit 638. It's a single page document bearing the
7 Bates number 505700. Do you recognize this document,
8 Dr. Kawasaki?
9 A. Looks familiar, '88, uh-huh.
10 Q. The document bears the title "Virus
11 Detection From Cell Sups or Serum."
12 A. Uh-huh.
13 Q. And what does this protocol -- what is this
14 protocol? Can you just describe what the protocol is?
15 A. Okay, let's see. As far as I remember, back
16 then, there was discussion of detecting -- detecting
17 HIV as DNA or as HIV as viral particles, so the way to
18 detect as viral particles, you have to separate cells
19 from viral particles, okay. So if you take a blood
20 sample, there's a huge number of cells. You have to
21 isolate the plasma or serum before you can study viral
22 particles, okay? So this protocol was set up -- I set
23 up -- it actually was an HIV at the time. It was
24 Moloney leukemia virus, and it's a mouse retrovirus
25 which infects -- which is -- infects mouse cell lines,

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1 and there are certain cell lines that produce a large
2 number of viral particles. So I developed a protocol
3 to -- to isolate the viral particles or extract RNA
4 from viral particles to detect retroviral sequences.
5 Q. Did you ever use this protocol for detecting
6 retroviral sequences from HIV?
7 A. Can't remember, but I would say no.
8 Q. Do you know if anyone in your lab ever did
9 that?
10 A. Not in my lab, but I can't remember, so
11 anyway --
12 Q. If you compare this protocol to the protocol
13 for RNA extraction in Exhibit 1, would you agree that
14 the RNA extraction protocol in Exhibit 1 is different
15 from the protocol in Exhibit 638?
16 MR. WILSON: Objection. The documents speak
17 for themselves. Also, are you asking him to make the
18 comparison sitting here today or are you asking for
19 his knowledge?
20 BY MS. RHYU:
21 Q. Can you answer the question, Dr. Kawasaki?
22 And I'm referring you to that paragraph that we were
23 looking at earlier, the RNA extraction, reverse
24 transcription and amplification of cDNA on the first
25 page of Exhibit 1.

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1 MR. WILSON: Okay, in that case calls for
2 expert testimony and speculation.
3 A. Are you saying to say whether it's different
4 or it's just a variation of a theme? RNA extraction
5 -- there are a lot of different RNA extraction
6 methods.
7 Q. The RNA extraction method in Exhibit 1 uses
8 guanidinium thiocyanate, correct?
9 MR. WILSON: The document speaks for itself.
10 A. That's what it does, yes.
11 Q. You're agreeing with me?
12 A. Uh-huh.
13 Q. Yes, and the protocol in Exhibit 638 does
14 not use guanidinium thiocyanate, right?
15 MR. WILSON: The document speaks for itself.
16 A. Yes.
17 Q. Yes, that's correct?
18 A. Uh-huh, yes.
19 Q. And I'm handing you what was previously
20 marked as Exhibit 46.
21 MR. WILSON: Do you have a copy, Counsel?
22 BY MS. RHYU:
23 Q. Oh, I'm sorry. Do you recognize Exhibit 36?
24 It's an article that was published in November of 1991
25 in the Journal of Clinical Investigation.

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1 primer that --
 2 A. That -- whatever we used back then, so I'm
 3 just saying in this case it worked better.
 4 Q. In looking at the JID article, Exhibit 1,
 5 you referred to some primers that were described in
 6 the methods of Exhibit 1. Those primers were not
 7 using random hexamers, correct?
 8 A. Let's see.
 9 MR. WILSON: Objection, vague, also
 10 misstates the documents.
 11 A. Actually, I can't say what they're using
 12 here.
 13 Q. Do you see where it says -- it lists
 14 different SK probes?
 15 A. But that was for PCR, not for the --
 16 Q. Is there anything in Exhibit 1 that suggests
 17 that random hexamers were used?
 18 A. That's --
 19 MR. WILSON: Objection, calls for expert
 20 testimony. The document speaks for itself, also
 21 mischaracterizes the document.
 22 A. Okay, well, in this Exhibit 1, the SK
 23 primers are for PCR, and in this statement for reverse
 24 transcriptase, it doesn't say what he uses. He refers
 25 back to this paper. That's all I can say about that.

1 nucleated cells are created.
 2 Q. Those are the white cells?
 3 A. Yeah, the white cells, and then do it that
 4 way. You get a different -- slightly different cell
 5 population by fractionating versus lysis, but that's
 6 another technical issue
 7 Q. But both of these procedures involve
 8 preparation of DNA from cells.
 9 A. Uh-huh, yes.
 10 Q. Dr. Kawasaki, I'm handing you what was
 11 previously marked as Exhibit 15. It's U.S. patent
 12 number 5,968,730, issued to Thomas Merigan, David
 13 Katzenstein and Mark Holodniy. Can you just let me
 14 know if you've seen that document before?
 15 A. No.
 16 Q. You have not seen it?
 17 A. I have not seen this.
 18 Q. And I'm handing you what's previously marked
 19 as Exhibit 16. It's U.S. patent number 6,503,705.
 20 Have you seen this patent before?
 21 A. No, no, not at all.
 22 MS. RHYU: If we could just take a short
 23 five-minute break, I can see if there's anything more
 24 I need to cover.
 25 MR. WILSON: Sure.

1 Q. I'm handing you what's been marked as
 2 Exhibit 641. This is another chapter from the PCR
 3 protocols book. It's chapter 18 entitled "Sample
 4 Preparation From Blood Cells and Other Fluids," and
 5 you are the sole author of this chapter. Did you
 6 write chapter 18 for the PCR protocols book?
 7 A. Yes.
 8 Q. And did you in writing this chapter intend
 9 to disseminate information about sample preparation
 10 for use in PCR amplifications?
 11 A. Yes, that was the --
 12 Q. Can you turn to page 148?
 13 A. 148, yeah.
 14 Q. There's a section in the middle of the page
 15 entitled "Preparation of DNA From White Blood Cells or
 16 Whole Blood." Can you just explain the difference
 17 between fractionated cells and whole blood?
 18 A. Let's see. Which way did I do this? Okay,
 19 so fractionated cells are when you take blood and spin
 20 it in a -- in a column, it's called ficoll-hypaque,
 21 and what it does is fractionate the red cells from the
 22 white cells, and so you can isolate the white cells,
 23 and from there, you can isolate DNA or RNA. If you
 24 take whole blood without fractionation, you can just
 25 lyse the red cells and then just spin out the -- the

1 MS. RHYU: Thanks.
 2 (Recessed at 12:19 p.m.)
 3 (Reconvened at 12:27 p.m.)
 4 BY MS. RHYU:
 5 Q. Did anyone outside of Cetus ask you about
 6 how to perform reverse transcription PCR reactions
 7 prior to the publication of the protocols in the PCR
 8 protocols book?
 9 A. Can't remember, but I would say probably.
 10 Q. And what was your practice when people asked
 11 you about the reverse transcriptase PCR procedure?
 12 MR. WILSON: Objection, vague.
 13 A. What was my reaction?
 14 Q. What was your practice? Did you help them?
 15 MR. WILSON: Objection, vague and
 16 foundation.
 17 A. I can't remember something like that because
 18 it's just -- I would say I don't know. I can't
 19 remember the practice.
 20 Q. I guess what I'm getting at is when outside
 21 scientists approached you with a request for help in
 22 their RT-PCR reactions, was it your practice to help
 23 them or not?
 24 MR. WILSON: Objection, vague, hypothetical,
 25 foundation.

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9 I, ERNEST S. KAWASAKI, PH.D., do hereby declare under
10 penalty of perjury that I have read the foregoing
11 transcript; that I have made any corrections as appear
12 noted, in ink, initialed by me, or attached hereto; that
13 my testimony as contained herein, as corrected, is true
14 and correct.
15 EXECUTED this _____ day of _____,
16 20____, at _____,
(City) (State)

17
18
19
20 _____
ERNEST S. KAWASAKI, PH.D.

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22
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24
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1 CERTIFICATE OF SHORTHAND REPORTER - NOTARY PUBLIC
2 I, Karen Young, the officer before whom the
3 forgoing deposition was taken, do hereby certify that
4 the forgoing transcript is a true and correct record
5 of the testimony given; that said testimony was taken
6 by me stenographically and thereafter reduced to
7 typewriting under my supervision; and that I am
8 neither counsel for or related to, nor employed by any
9 of the parties to this case and have no interest,
10 financial or otherwise, in its outcome.
11 IN WITNESS WHEREOF, I have hereunto set my
12 hand and affixed my notarial seal this ____ day of
13 _____,
14 _____
15 _____
16 _____
17 NOTARY PUBLIC IN AND FOR
18 THE DISTRICT OF COLUMBIA
19
20 My commission expires:
21 July 31, 2009
22
23
24
25

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