

6/29/2007 Varki, Ajit

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**EXHIBIT B**

1 UNITED STATES DISTRICT COURT  
 2 DISTRICT OF MASSACHUSETTS  
 3  
 4  
 5 AMGEN INC., )  
 6 )  
 7 Plaintiff, )  
 8 )  
 9 vs. ) Civil Action  
 10 )  
 11 F. HOFFMANN-LA ROCHE LTD., a Swiss ) No. 05-12237 WGY  
 12 Company, ROCHE DIAGNOSTICS GmbH, )  
 13 a German Company, and HOFFMANN-LA )  
 14 ROCHE, INC., a New Jersey )  
 15 Corporation )  
 16 )  
 17 Defendants. )  
 18 )  
 19 )  
 20 )  
 21 )  
 22 )  
 23 )

**CONFIDENTIAL**  
**REDACTED**

1 APPEARANCE:  
 2  
 3 For the Plaintiff:  
 4 DAY CASEBEER MADRID & BATCHELDER, LLP  
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 13 BY: CHRISTOPHER JAGOE, ESQUIRE  
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18 Also Present:  
 19 Michael Bullerman  
 20 John Arel, Videographer

21 (This Transcript contains testimony designated  
 22 confidential as per Section 5(c) of the Amended  
 23 Protective Order. Please treat the entire  
 24 transcript in accordance with the protective  
 25 order.)

14 DEPOSITION OF AJIT VARKI, M.D.  
 15 FRIDAY, JUNE 29, 2007  
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 23  
 24 REPORTED BY:  
 25 LINDA D. WHITE  
 CSR NO. 12009

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14 DEPOSITION OF AJIT VARKI, M.D.,  
 15 taken on behalf of Defendant, at  
 16 333 West Harbor Drive, Suite Los  
 17 Angeles, San Diego, California,  
 18 8:03 a.m., Friday, June 29, 2007,  
 19 before LINDA D. WHITE, Certified  
 20 Shorthand Reporter Number 12009 for  
 21 the State of California, pursuant  
 22 to Notice.  
 23  
 24  
 25

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1 BY MR. JAGOE:  
 2 Q Okay. Look at your third report. You can  
 3 keep that aside.  
 4 Look at your third report.  
 5 A 3 supplemental. Second supplemental?  
 6 Q Second supplemental report.  
 7 A Should I keep this here? I might get in  
 8 trouble. Close it over there.  
 9 Q Okay.  
 10 A Okay.  
 11 Q And paragraph 25 on Page 11 and 12.  
 12 A 25 on Page 11 and 12. 11 and 12. Sorry.  
 13 Okay.  
 14 Q All right. And at the end of  
 15 paragraph 25, you have six bulletpoints, right?  
 16 A Right.  
 17 Q And you list isoelectric points of  
 18 glycoforms, sulfation, polylactosamine, repeat  
 19 content?  
 20 A Correct.  
 21 Q Old glycan structure, presence of  
 22 N-glycolylneuraminic acid and absence of alpha 2-6  
 23 sialic acid linkages, correct?  
 24 A Correct. N-glycolyl, G-L-C-O-L-Y-L.  
 25 Q Would those six points summarize the

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1 differences between the prior art EPO that you  
 2 analyzed and compounds that fall within the scope of  
 3 claims of the 933 Patent?  
 4 A I think those are the ones that -- yes, I  
 5 think we -- I tried to summarize there what all the  
 6 issues I looked at. I can't think of any other.  
 7 But of course, a lot of these differences are not  
 8 based on what was known in 1983. But subsequent  
 9 analysis of either Miyake or Miyake-like  
 10 preparations.  
 11 Q All of these differences would be  
 12 differences in the carbohydrate component of the  
 13 erythropoietin, right?  
 14 MR. LOEB: Objection. Vague and ambiguous.  
 15 THE WITNESS: Carbohydrates refers to molecules  
 16 that have a particular composition, a CHO  
 17 composition. The sulfate, for example, would not  
 18 fall in the category of carbohydrates. The  
 19 N-glycolyl, I suppose, could, but it becomes a  
 20 semantic issue or technical issue.  
 21 All the differences -- perhaps better to  
 22 put it as all the differences are related to things  
 23 that are components of the glycans that are attached  
 24 to the polypeptide. That is the way of summarizing  
 25 all of the difference I looked at. So everything

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1 beyond the glyco -- peptide -- polypeptide itself.  
 2 BY MR. JAGOE:  
 3 Q Are any of the differences related to the  
 4 folding of the polypeptide?  
 5 A Not directly, but the -- it's well known  
 6 that the glycans would -- could have a dramatic  
 7 effect on the folding of the polypeptide during  
 8 biosynthesis. I believe I addressed that in some  
 9 other places in the reports.  
 10 Q The sulfation difference, the sulfation  
 11 takes place on the carbohydrate as opposed to on the  
 12 amino acid residues; is that right?  
 13 MR. LOEB: Objection. Vague and ambiguous.  
 14 THE WITNESS: The presence of the sulfate  
 15 esters, based on the data I've seen, especially the  
 16 results of using the PNGase F enzyme. The data  
 17 suggests. But again, sulfation is extremely  
 18 difficult to study and has been very poorly studied  
 19 in most of this work. It seems like it's on --  
 20 attached to the sugar chains.  
 21 BY MR. JAGOE:  
 22 Q Okay. Now, the -- let me just ask you.  
 23 So do you agree with the statement that  
 24 urinary EPO and recombinant EPO are the same  
 25 product?

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1 MR. LOEB: Objection. Vague and ambiguous.  
 2 THE WITNESS: Can you define "the same  
 3 product"? Is it the source or is it the production  
 4 method or is it --  
 5 BY MR. JAGOE:  
 6 Q The substances themselves are the same.  
 7 MR. LOEB: Objection. Vague and ambiguous.  
 8 THE WITNESS: The substance -- they're not.  
 9 From everything I've seen, there is a lot of  
 10 differences between them.  
 11 BY MR. JAGOE:  
 12 Q And are you intending to offer an opinion  
 13 that urinary EPO and recombinant EPO can be  
 14 distinguished?  
 15 A If you define recombinant EPO as all the  
 16 recombinant EPOs that have been made to date, and  
 17 all the ones I've seen -- and there seems to be a  
 18 lot of them that have been studied -- yes, I have  
 19 not seen a single recombinant EPO prep that -- well,  
 20 if you just look at the -- the way it's used in --  
 21 in -- you know, looking for EPO doping in sports,  
 22 it's very obvious they never have a problem telling  
 23 them apart right from there.  
 24 And since the differences they're looking  
 25 at directly reflect differences in the glycan

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