Exhibit B

David Cousineau /DC/US/KSFHH 03/27/2007 05:45 PM To "Mathie, BeLinda I." <belinda.mathie@kattenlaw.com>

CC

bcc Roche Email Repository@KSFHHNotes
Subject Re: Summary of DaVita/Roche CTSAs

Belinda,

Thank you for sending me this document Roche does not have a problem with you producing the document as is.

Best, David

David L. Cousineau Kaye Scholer LLP 901 Fifteenth St, NW Washington, D.C. 20005-2327 Phone: 202/682-3617

Phone: 202/682-3617 Fax: 202/414-0344

"Mathie, BeLinda I." <belinda.mathie@kattenlaw.com>



"Mathie, BeLinda I." <belinda.mathie@kattenlaw .com> 03/27/2007 02:54 PM

To <DCousineau@kayescholer.com>

cc "Kemnitz, Christian T." <christian.kemnitz@kattenlaw.com>

Subject Summary of DaVita/Roche CTSAs

David,

Attached please find for your information the current version of the document we intend to produce to Amgen summarizing DaVita's clinical trial service agreements with Roche. You will note that the two protocols you believe are non -responsive, ML20336 and ML20338 are not listed. Chris will give you a call to discuss.

BeLinda I. Mathie Litigation Associate Katten Muchin Rosenman LLP 525 West Monroe Chicago, Illinois 60661 (312) 902-5283 (direct) (312) 577-4482 (fax)

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(New) Summary of Clinical Trial Services Agreements Executed by DaVita with Roche.DOC

Summary of Clinical Trial Services Agreements/SMO Clinical Trial Agreements Executed by DaVita with Roche

Protocol No.	Effective Date	Date Executed	Maximum Number of Patients	Expected Duration (Best Efforts Completion Date)	Exclusionary Provisions
Protocol# BA17284	7/20/04	7/29/04	6	8/25/05	None
Rollover Study of 7/20/04 Agreement (Protocol #BH18387)	5/9/05	7/16/05	7	December 2006	None
Protocol# BA17284	9/7/04	10/7/04	6	8/25/05	None
Rollover Study of 9/7/04 Agreement (Protocol #BH18387)	7/15/05	7/19/05	3	December 2006	None

David Cousineau /DC/US/KSFHH 03/29/2007 10:48 AM

- To "Fishman, Deborah" <dfishman@daycasebeer.com>
- Kallus" <Kallus@fr.com>; "Krista Carter" bcc Roche Email Repository@KSFHHNotes

Subject RE: Amgen/Roche: Fresenius's Document Production to Amgen

Dear Deborah,

We have explained our position on these points numerous times in correspondence with you and in briefs filed with the court. I do not see the point in belaboring this further.

I learned this morning that of the documents related to clinical trials Mark Hebert sent us, only one trial is ongoing. I passed this information on to Mark this morning and told him that Roche does not object to production of the documents related to the completed clinical trials. For the reasons set out in our prior correspondence, in briefs to the court, and in the court's orders, we maintain our objections to production of information related to the one ongoing clinical trial. We will similarly maintain this objection during the Fresenius deposition on Friday.

Your suspicions about Roche's motives are incorrect. We are simply trying to maintain the integrity of the court-imposed discovery limits. Furthermore, your selective quotation of the Model Rules is disingenuous. First, as Judge Young ruled, the information requested in your subpoena is not relevant (as required by your quote) to this case. Second, the model rule only pertains to instances where the attorney does not "reasonably believe] that the person's interests will not be adversely affected by refraining from giving such information." Because Fresenius is a third-party whom Amgen is pressuring to produce irrelevant information, it is hard to imagine how Roche's communications to Fresenius's lawyer about the scope of relevant discovery adversely affects Fresenius's interests.

Thank you, David

David L. Cousineau Kaye Scholer LLP 901 Fifteenth St, NW Washington, D.C. 20005-2327 Phone: 202/682-3617

Fax: 202/414-0344

"Fishman, Deborah" <dfishman@daycasebeer.com>



"Fishman, Deborah" <dfishman @daycasebeer.com> 03/28/2007 10:28 PM

To <dcousineau@kayescholer.com>; <jbrew@kayescholer.com>

Subject RE: Amgen/Roche: Fresenius's Document Production to Amgen

Dear David,