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| 8 | UNITED STATES DISTRICT COURT |
| 9 | NORTHERN DISTRICT OF CALIFORNIA |
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| 11 | GENENTECH, INC., et al., |
| 12 |) Plaintiff(s), No. C08-4909 SI (BZ) |
| 13 | v.) |
| 14 |) SIXTH DISCOVERY ORDER SANOFI-AVENTIS DEUTSCHLAND) |
| 15 | GMBH, et al.,) |
| 16 | Defendant(s).)) |
| 17 | Following a telephone hearing at which all parties were |
| 18 | represented by counsel, IT IS HEREBY ORDERED as follows: |
| 19 | 1. Sanofi-Aventis Deutschland GmbH ("Sanofi") requests |
| 20 | detailed discovery of Rituxan® sales in order to prove the |
| 21 | commercial success of its patented process. Ordinarily, a |
| 22 | "patentee asserts that commercial success supports its |
| 23 | contention of nonobviousness" <u>Demaco Corp. v. F. Von</u> |
| 24 | Langsdorff Licensing Ltd., 851 F.2d 1387, 1392 (Fed. Cir. |
| 25 | 1988). The patentee must prove a legally and factually |
| 26 | sufficient connection, or nexus, "between the proven success |
| 27 | and the patented invention" <u>Id.</u> The purpose of |
| 28 | introducing evidence of commercial success is to prove that |

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"the commercial success was of the <u>patented</u> invention itself."
 <u>Id.</u> at 1394. (emphasis added).

In this case, Sanofi seeks detailed sales information of 3 the infringing product in order to prove non-obviousness of 4 5 the patented process. Sanofi has not persuaded me that 6 discovery of sales information of an infringing product is 7 relevant to prove the non-obviousness and commercial success of the patented process. Commercial success of the allegedly 8 infringing product could be based on any number of factors, 9 10 other than infringement. In any event, defendant apparently has information that Genentech has \$2 billion annual sales of 11 12 Rituxan® in the United States, evidence of the product's 13 commercial success. Defendant has not articulated how the 14 other information it seeks, such as profit margins, is 15 relevant to obviousness.

Tech Air, Inc. v. Denso Mfg. Michigan Inc., 192 F.3d 1353 16 17 (Fed. Cir. 1999) on which defendant relies, is 18 distinguishable. First, that case did not address discovery. 19 Second, the plaintiff in that case had already established 20 infringement. Contrary to Sanofi's assertion, Tech Air does 21 not stand for the general proposition that in a bifurcated 22 trial, sales information of an infringing product is discoverable prior to the damages phase. Defendant's motion 23 24 to compel discovery of further sales information is **DENIED**.

Sanofi further requests discovery of all products
 that employ the HCMV enhancer to prove the unexpected results
 of their patented process. As an initial matter, this request
 suffers from the same defect as Sanofi's request for detailed

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sales information of the infringing product. In the ordinary 1 2 case, a "patent challenger makes a prima facie showing of obviousness" and "the owner may rebut [this] based on 3 unexpected results by demonstrating that the claimed invention 4 5 exhibits some superior property or advantage that a person of ordinary skill in the relevant art would have found surprising 6 or unexpected." Proctor & Gambel Co. v. Teva Pharmaceuticals 7 USA, Inc., 566 F.3d 989 (Fed. Cir. 2009) (emphasis added) 8 9 (internal citations omitted).

10 Sanofi has not persuaded me that any unexpected results plaintiffs experienced with any of their products is relevant 11 12 to the issue of obviousness. In any event, to resolve this 13 dispute, Genentech has agreed to produce the requested discovery for Rituxan®, the accused product. If Sanofi cannot 14 15 show that Genentech experienced unexpected results with 16 Rituxan®, it is hard to see how it can succeed by showing 17 unexpected results with products not accused of infringement. 18 If Sanofi finds evidence of unexpected results in the Rituxan® 19 discovery and believes it still needs evidence of other 20 products, it may renew its motion at that time. Given the 21 cost of producing the requested information, calculated at 22 almost a million dollars, and it questionable relevancy, this 23 request is **DENIED** as premature. See FRCP 26(b)(2)(c)(iii).

Genentech shall produce all documents concerning its use of an HCMV enhancer in its production of Rituxan® by /// 27 /// 28 ///

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| 1 | 5:00 p.m. on December 1, 2009. |
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| 2 | Dated: November 19, 2009 |
| 3 | Bernard Zimmerman |
| 4 | United Sta te s Magistrate Judge |
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