UNITED STATES PATENT AND TRADEMARK OFFICE BEFORE THE PATENT TRIAL AND APPEAL BOARD

REGENERON PHARMACEUTICALS, INC., Petitioner,

v.

NOVARTIS PHARMA AG, NOVARTIS TECHNOLOGY LLC, NOVARTIS PHARMACEUTICALS CORPORATION, Patent Owner.

Case No. IPR2021-00816 U.S. Patent No. 9,220,631

JOINT MOTION TO SEAL THE FINAL WRITTEN DECISION 37 CFR §§ 42.14 AND 42.54



I. PRECISE RELIEF REQUESTED

Pursuant to 37 C.F.R. §§ 42.14 and 42.54, Petitioner Regeneron

Pharmaceuticals, Inc. ("Petitioner" or "Regeneron") and Patent Owners Novartis

Pharma AG, Novartis Technology LLC, and Novartis Pharmaceuticals Corp.

("Patent Owner" or "Novartis") respectfully move to seal portions of Paper No.

113, the Board's October 25, 2022 Final Written Decision ("Final Written

Decision" or "FWD"), and file a public redacted version of the Final Written

Decision. A proposed public version of the Final Written Decision with redactions agreed-upon by the parties has been filed concurrently as Ex. 1257.

II. REASONS FOR THE REQUESTED RELIEF AND STATEMENT OF FACTS

A. Good Cause Exists for Sealing Confidential Information

The Board will seal documents for good cause. See 37 C.F.R. § 42.54(a); see also Argentum Pharms. LLC v. Alcon Research, Ltd., Paper 27, 2 (2013). "The rules aim to strike a balance between the public's interest in maintaining a complete and understandable file history and the parties' interest in protecting truly sensitive information." Office Patent Trial Practice Guide, 77 Fed. Reg. 48756, 48760 (2012). The public's interest in having access to confidential business information that is only indirectly related to patentability is "minimal." Garmin v.



Cuozzo, IPR2012-00001, Paper 36, 8-9 (2013) (granting a motion to seal an agreement relating to the "commercializ[ation]" of the patent-at-issue).

The information that Petitioner and Patent Owner seek to seal is either Regeneron's confidential research and development information, Novartis's confidential research and development information, Novartis's confidential commercial and financial information, Novartis's confidential product information, or third party confidential information, as explained in more detail below. To the undersigned's knowledge, the information sought to be sealed has not been published or otherwise made public. Public disclosure of Regeneron's, Novartis's or third party confidential information would competitively harm Regeneron's, Novartis's, and/or third parties' business prospects and put these companies at a competitive disadvantage relative to other similarly positioned companies in the same industry. Therefore, good cause exists to seal portions of the Final Written Decision.

B. Good Cause Exists for Sealing Regeneron's Confidential Information

Portions of the Final Written Decision contain confidential information concerning the specifications of Regeneron's product. *See* FWD at 58, 61, 115 (discussing confidential product specifications of EYLEA PFS). This confidential and proprietary information of Regeneron, if publicly disclosed, would substantially harm Regeneron's competitive position in the pharmaceutical



industry. For example, this information, if not sealed, would allow competitors to use such information to improve its products. Therefore, good cause exists to seal Regeneron's confidential information described in the Final Written Decision and file a public redacted version of the Final Written Decision.

C. Good Cause Exists for Sealing Novartis's Confidential Information

Portions of the Final Written Decision contain Novartis's confidential research and development information. See FWD at 40 (discussing Novartis's internal presentations relating to drug development); 42, 46, 52, 97-98 (discussing Novartis's confidential correspondences with third parties regarding product development). The Final Written Decision also contains Novartis's confidential commercial and financial information. See FWD at 44, 89 (discusses confidential sale and revenue figures for Novartis's products); 53, 98-99 (discussing terms of Novartis's confidential commercial contract with third party); 85 (discussing confidential results of marketing studies); 48, 83-84, 85-86 (referencing Novartis employees' confidential testimonies regarding Novartis's marketing strategy and results). Moreover, portions of the Final Written Decision contain Novartis's confidential information relating to product specifications and features. See FWD at 48-50, 75-79, 81-83, 87, 90, 112 (discussing confidential specifications and features of Lucentis PFS).



Such confidential and proprietary research, commercial, and product specification information of Novartis, if publicly disclosed, would substantially harm Novartis's competitive position in the pharmaceutical industry. For example, this information, if not sealed, would provide competitors with valuable information regarding confidential research and development projects, as well as sensitive financial and market information. Such information could also be used by a competitor to improve its products. Therefore, good cause exists to seal Novartis's confidential information and file a public redacted version of the Final Written Decision.

D. Good Cause Exists for Sealing Third Party Confidential Information

Portions of the Final Written Decision contain third party confidential information relating to product specifications. *See* FWD at 45, 56, 64, 72, 91, 92 (discussing third party Bausch's confidential product specification information); 51, 92, 98 (discussing third party Becton Dickinson's confidential product information). The Final Written Decision also contains third party confidential development and research information. *See* FWD at 45-46, 52, 72-73, 94-96 (discussing third party's confidential product development process).

Public disclosure of the proposed redacted information would harm these third parties, because insights into third parties' technical and development information would provide a competitive advantage to such third parties'



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