

EXHIBIT C

IN THE UNITED STATES DISTRICT COURT
FOR THE NORTHERN DISTRICT OF WEST VIRGINIA
CLARKSBURG DIVISION

REGENERON PHARMACEUTICALS, INC.,

Plaintiff,

v.

MYLAN PHARMACEUTICALS INC.,

Defendant.

Civil Action No. 1:22-cv-00061-TSK

STIPULATED PROTECTIVE ORDER

Pursuant to Northern District of West Virginia Local Rule 26.05, and upon agreement of the Parties for an order pursuant to Fed. R. Civ. P. 26(c), Plaintiff Regeneron Pharmaceuticals, Inc. (“Regeneron”) and Defendant Mylan Pharmaceuticals Inc. (“Mylan”) (each a “Party” and collectively the “Parties”), hereby stipulate and the Court orders as follows:

I. Definitions.

1. As used in this Protective Order, these terms have the following meanings:
 - (a) “Patents-in-Suit” refers to United States Patents that Regeneron has claimed Mylan infringes as set forth in the Complaint filed on August 2, 2022 or any Amended Complaint that Regeneron files once it becomes the operative pleading in this Action;
 - (b) “Action” means the case captioned *Regeneron Pharmaceuticals, Inc. v. Mylan Pharmaceuticals Inc.*, Case No. 1:22-cv-00061-TSK, which is currently pending in the Northern District of West Virginia;

- (c) “Affiliate” means any Third Party that, directly or indirectly, through one or more intermediaries controls, or is controlled by, or is under common control with, a Party to this Action;
- (d) “CONFIDENTIAL” means information that constitutes, contains, reveals, or reflects trade secrets or other confidential research, development, business, or commercial information within the meaning of Fed. R. Civ. P. 26(c)(1)(G), including but not limited to: scientific and technical information; financial (including pricing and sales information), budgeting and/or accounting information; information about existing and potential customers; marketing and other business strategies, decisions or negotiations; employee compensation, evaluation and other employment information; business plans; manufacturing information; licensing agreements; regulatory information (including non-public correspondence with the United States and foreign patent offices and regulatory agencies); and information that concerns or relates to the trade secrets, processes, operations, style of work, or apparatus, or to the production, sales, shipments, purchases, transfers, identification of customers, inventories, amount or source of any income, profits, losses, or expenditures of any person, firm, partnership, corporation, or other organization; and any other information that is protected from disclosure by the laws of the United States or another country; and includes such confidential and proprietary information about a Third Party, including parents, subsidiaries, and/or other Affiliates; and any other information, including “Personal Data” as

defined below, the disclosure of which would harm the competitive position of the Producing Party if the information becomes known to a person or party other than the Producing Party other than as permitted hereunder;

- (e) “Designated In-house Counsel” means an attorney designated in accordance with Paragraph 6(i);
- (f) “Designating Party” is a Party or Third Party that designates information or items that it produces in Disclosures as “CONFIDENTIAL” or as “OUTSIDE COUNSEL’S EYES ONLY”;
- (g) “Disclosure(s)” means all documents; written discovery requests and responses; deposition transcripts; correspondence between the Parties; pleadings; exhibits; documents and things made available for inspection; expert testimony and reports; biological materials produced by a Party or Third Party in this Action including, but not limited to, any physical samples of cells, polynucleotides, or proteins; all other discovery taken pursuant to the Federal Rules of Civil Procedure, including Third Party discovery pursuant to Federal Rule of Civil Procedure 45; and tangible items and any other information produced or disclosed between the Parties in connection with this Action, including the pre-suit exchanges made pursuant to 42 U.S.C. § 262(*l*). For the sake of clarity, the production of biological samples shall be governed by the terms set forth in Paragraph 31;
- (h) “Document(s)” means all materials within the scope of Federal Rule of Civil Procedure 34(a);

- (i) “Expert” is a person with specialized knowledge or experience in a matter pertinent to the Action who (1) has been retained by a Party or its Outside Counsel to serve as an expert witness or as a consultant in this Action, (2) is not a current employee of a Party, (3) at the time of retention, is not anticipated to become an officer, director, or employee of a Party. Nothing in this Protective Order purports to alter the requirements for offering testimony under Federal Rule of Evidence 703, or to define the term “expert” for purposes other than those addressed in this Protective Order;
- (j) “In-House Counsel” means an in-house attorney who is an employee of a Party to this Action or a Party’s Affiliate;
- (k) “Third Party” is any natural person, partnership, corporation, association, or other legal entity not named as a Party to this Action;
- (l) “Outside Counsel” means the external lawyers retained by the Parties to litigate this Action and who have appeared in this Action or are acting at the direction of a lawyer who has appeared in this Action, including, but not limited to, outside attorneys admitted *pro hac vice* in this Action and paralegals, assistants and other employees of the respective law firms of Outside Counsel;
- (m) “OUTSIDE COUNSEL’S EYES ONLY” means any Disclosure that contains highly sensitive information relevant to the Designating Party’s current or prospective products, scientific or technical information, business information of current or prospective significance and for which production on a confidential basis, even to In-house Counsel, would create a substantial

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