## IN THE UNITED STATES DISTRICT COURT FOR THE DISTRICT OF DELAWARE

In	re	Sit	aglipt	tin	Phos	phate	('708	&	'921)
Pa	iter	nt L	itigat	ioi	n				

C.A. No. 19-md-2902-RGA

MERCK SHARP & DOHME CORP.,

Plaintiff,

V.

C.A. No. 19-1489-RGA

MYLAN PHARMACEUTICALS INC.,

Defendants.

## PROPOSED SCHEDULING ORDER

This <u>13</u> day of August, 2019, the Court having conducted an initial Rule 16(b) scheduling conference pursuant to Local Rule 16.1(b), and the parties having determined after discussion that the matter cannot be resolved at this juncture by settlement, voluntary mediation, or binding arbitration;

Additionally, on August 8, 2019, the Judicial Panel on Multidistrict Litigation ordered the centralization of this action, the related actions, and *Merck Sharp & Dohme Corp. v. Mylan Pharmaceuticals Inc. et al.*, Case No. 19-cv-101-IMK (N.D. W. Va.), to the U.S. District Court for the District of Delaware for coordinated and consolidated proceedings. *See* Transfer Order, *In re Sitagliptin Phosphate ('708 & '921) Patent Litigation*, MDL No. 2902 (J.P.M.L. Aug. 8, 2019), ECF No. 56.



<sup>&</sup>lt;sup>1</sup> This Order follows substantively identical scheduling orders dated June 28, 2019, in related actions, C.A. Nos. 19-310-RGA, 19-311-RGA, 19-312-RGA, 19-314-RGA, 19-316-RGA, 19-317-RGA, 19-318-RGA, 19-320-RGA, 19-321-RGA, and 19-347-RGA, involving the same products and patents. The parties have agreed that the schedule in the related actions should apply in this action. This Order thus provides for the same due dates as the scheduling orders in the related actions, but has been edited to account for due dates that have already passed.

### IT IS ORDERED that:

- 1. Rule 26(a)(1) Initial Disclosures. The parties have agreed to exchange their initial disclosures pursuant to Federal Rule of Civil Procedure 26(a)(1) and Paragraph 3 of the Delaware Default Standard for Discovery on or before August 20, 2019.
- 2. <u>Joinder of Other Parties and Amendment of Pleadings</u>. All motions to join other parties, and to amend or supplement the pleadings, shall be filed on or before February 14, 2020.

## 3. <u>Discovery</u>.

- a. <u>Fact Discovery Cut Off.</u> All fact discovery in this case shall be initiated so that it will be completed on or before November 20, 2020.
- b. <u>Document Production</u>. Document production shall be substantially complete by February 21, 2020.
- c. <u>Contentions</u>. The parties shall make their initial disclosures under Paragraphs 3 and 4 in accordance with the Court's Default Standard for Discovery, Including Discovery of Electronically Stored Information ("the Default Standard").
  - Plaintiff shall make its disclosures under Paragraph 4(a) of the Default Standard by August 20, 2019.
  - ii. Defendant shall produce its core technical documents under Paragraph

    4(b) of the Default Standard by August 23, 2019. At the same time

    Defendant produces its core technical documents, Defendant shall also

    produce the DMF for the sitagliptin API used in its proposed ANDA

    products, to the extent it is in Defendant's possession, custody, or control,

    or if Defendant able to obtain the DMF pursuant to a contractual right to

    the DMF with the DMF holder. If Defendant not in possession, custody,



- or control of the DMF, and is not able to obtain the DMF pursuant to a contractual right with the DMF holder, the Defendant shall inform Plaintiff of that fact and identify the DMF holder at the same time it produces their core technical documents.
- iii. Plaintiff shall make its disclosure under Paragraph 4(c) of the Default Standard within 30 days after receiving Defendant's disclosure under Paragraph 4(b) of the Default Standard;
- Defendant shall make its disclosures under Paragraph 4(d) of the Default
   Standard within 30 days after receiving Plaintiff's disclosure under
   Paragraph 4(c) of the Default Standard.
- v. The parties shall exchange supplemental infringement and invalidity contentions on October 14, 2020.
- d. Samples. At the same time Defendant produces its core technical documents, Defendant shall also disclose to Plaintiff whether it is able to produce reasonable quantities of unexpired samples (unexpired as of the entry of this Scheduling Order) of its ANDA products and API to the extent such samples are in Defendant's possession, custody, or control, or if Defendant can obtain such samples pursuant to a contractual right with a supplier. On or before August 27, 2019, Defendant shall produce reasonable quantities of unexpired samples (unexpired as of the entry of this Scheduling Order) of the ANDA products and API to the extent such samples are in Defendant's possession, custody, or control, or if Defendants can obtain such samples due to a contractual right with a supplier. To the extent that Defendant is unable to produce such samples on or before August 27, 2019, Defendant shall inform Plaintiff at the same time Defendant produces its core technical documents and confer with Plaintiff as to a



reasonable extension of time to produce such samples. For clarity, if Defendant does not have unexpired samples (unexpired as of the entry of this Scheduling Order) of their ANDA Product and API in their possession, custody, or control, and cannot obtain such samples pursuant to a contractual right with a supplier, Defendant shall inform Plaintiff of those facts at the same time it produces its core technical documents.

- e. Requests for Admission. Plaintiff may serve up to 15 requests for admission on the Defendant Groups collectively. To the extent that a request for admission is served on the Defendant Groups collectively, that request for admission shall count as one request for admission even if multiple parties provide a distinct response. The Defendant Groups collectively may jointly serve up to 15 requests for admission on Plaintiff. In addition, each Defendant Group may serve on Plaintiff up to 15 individualized requests for admission, and Plaintiff may serve on each Defendant Group up to 15 individualized requests for admission. Any additional requests for admission may only be served with leave of Court. Any requests for admission directed to the authentication of documents are excluded from the limitations above.
- f. <u>Interrogatories</u>. Plaintiff may serve up to **15** interrogatories on the Defendant Groups collectively. To the extent that an interrogatory is served on the Defendant Groups collectively, that interrogatory shall count as one interrogatory even if multiple parties provide a distinct response. The Defendant Groups collectively may jointly serve up to **15** interrogatories

<sup>&</sup>lt;sup>2</sup> The Defendant Groups (in this action and the related actions) are: (1) Alvogen Pine Brook LLC F/K/A Alvogen Pine Brook, Inc. and Alvogen Malta Operations Ltd.; (2) Anchen Pharmaceuticals, Inc. and Par Pharmaceutical, Inc.; (3) Apotex Inc. and Apotex Corp.; (4) Lupin Limited and Lupin Pharmaceuticals, Inc.; (5) Macleods Pharmaceuticals Limited and Macleods Pharma USA, Inc.; (6) Mylan Pharmaceuticals, Inc.; (7) Sandoz Inc.; (8) Sun Pharma Global FZE and Sun Pharmaceutical Industries Ltd.; (9) Teva Pharmaceuticals USA, Inc.; (10) Torrent Pharmaceuticals Limited and Torrent Pharma Inc.; (11) Watson Laboratories, Inc. and Teva Pharmaceuticals USA, Inc.; (12) Wockhardt Bio AG and Wockhardt USA LLC; and (13) Zydus Pharmaceuticals (USA) Inc. and Cadila Healthcare Ltd.



on Plaintiff. In addition, each Defendant Group may serve on Plaintiff up to 10 individualized interrogatories, and Plaintiff may serve on each Defendant Group up to 10 individualized interrogatories. Any additional interrogatories may only be served with leave of Court.

## g. Depositions.

- <u>Limitation on Hours for Deposition Discovery</u>. Plaintiff is limited to 50 hours of taking fact deposition testimony upon oral examination per Defendant Group, including testimony of former Defendant Group employees.<sup>3</sup> The Defendant Groups collectively are limited to 130 hours of taking fact deposition testimony upon oral examination, including testimony of former Plaintiff employees. Any deposition lasting less than 5 hours will count as 5 hours against the total time of the side taking the deposition. These hour limits on fact depositions may be increased by Court order upon good cause shown. Depositions of inventors of the patents-in-suit who are designated as 30(b)(6) witnesses will be limited to 10 hours per inventor. Depositions of inventors of the patents-in-suit who are not designated as 30(b)(6) witnesses will be limited to 7 hours per inventor. Separate and apart from these hour limits on fact depositions, Plaintiff may depose each witness offered as an expert by a Defendant Group, and the Defendant Groups collectively may depose each witness offered as an expert by Plaintiff. If a deponent testifies wholly or substantially through an interpreter, the party taking the deposition shall be permitted, on a pro rata basis, two hours of deposition time for each hour spent testifying through the interpreter. For clarity, the hour limitations described in this paragraph do not apply to depositions of third-parties or expert witnesses.
- ii. <u>Location of Depositions</u>. The parties shall meet and confer regarding the locations of depositions, taking into account convenience for the deponent.

<sup>&</sup>lt;sup>3</sup> To the extent the same individual is deposed for more than one Defendant Group, there shall be a single deposition.



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