

**UNITED STATES OF AMERICA  
BEFORE THE FEDERAL TRADE COMMISSION**

**COMMISSIONERS: Edith Ramirez, Chairwoman  
Julie Brill  
Maureen K. Ohlhausen  
Joshua D. Wright  
Terrell McSweeney**

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<b>In the Matter of</b>		)
		)
<b>ELI LILLY AND COMPANY,</b>		)
<b>a corporation;</b>		)
		)
<b>and</b>		)
		)
<b>NOVARTIS AG,</b>		)
<b>a corporation.</b>		)
_____		)

**Docket C-4500**

**ORDER TO MAINTAIN ASSETS**

The Federal Trade Commission (“Commission”), having initiated an investigation of the proposed acquisition by Respondent Eli Lilly and Company (“Eli Lilly”) of certain assets comprising the animal health division of Respondent Novartis AG (“Novartis”), collectively “Respondents”, and Respondents having been furnished thereafter with a copy of a draft of Complaint that the Bureau of Competition proposed to present to the Commission for its consideration and which, if issued by the Commission, would charge Respondents with violations of Section 7 of the Clayton Act, as amended, 15 U.S.C. § 18, and Section 5 of the Federal Trade Commission Act, as amended, 15 U.S.C. § 45; and

Respondents, their attorneys, and counsel for the Commission having thereafter executed an Agreement Containing Consent Orders (“Consent Agreement”), containing an admission by Respondents of all the jurisdictional facts set forth in the aforesaid draft of Complaint, a statement that the signing of said Consent Agreement is for settlement purposes only and does not constitute an admission by Respondents that the law has been violated as alleged in such Complaint, or that the facts as alleged in such Complaint, other than jurisdictional facts, are true, and waivers and other provisions as required by the Commission’s Rules; and

The Commission having thereafter considered the matter and having determined to accept the executed Consent Agreement and to place such Consent Agreement on the public record for a period of thirty (30) days for the receipt and consideration of public comments, now in further conformity with the procedure described in Commission Rule 2.34, 16 C.F.R. § 2.34, the Commission hereby issues its Complaint, makes the following jurisdictional findings, and issues this Order to Maintain Assets:

1. Respondent Eli Lilly is a corporation organized, existing and doing business under and by virtue of the laws of the State of Indiana, with its headquarters address located at Lilly Corporate Center, Indianapolis, Indiana 46285.
2. Respondent Novartis is a corporation organized, existing and doing business under and by virtue of the laws of the Swiss Confederation with its headquarters address located at Lichtstrasse 35, Basel, Switzerland, CH4056, and the address of its United States subsidiary, Novartis Corporation, located at 230 Park Avenue, New York, New York 10169.
3. The Commission has jurisdiction of the subject matter of this proceeding and of the Respondents, and the proceeding is in the public interest.

## **ORDER**

### **I.**

**IT IS ORDERED** that, as used in this Order to Maintain Assets, the following definitions and the definitions used in the Consent Agreement and the proposed Decision and Order (and, when made final and effective, the Decision and Order), which are incorporated herein by reference and made a part hereof, shall apply:

- A. “Eli Lilly” means the following: Eli Lilly and Company, its directors, officers, employees, agents, representatives, successors, and assigns; and its joint ventures, subsidiaries, divisions, groups and affiliates in each case controlled by Eli Lilly and Company, and the respective directors, officers, employees, agents, representatives, successors, and assigns of each. After the Acquisition, Eli Lilly shall include the Novartis Animal Health Group.
- B. “Novartis” means the following: Novartis AG, its directors, officers, employees, agents, representatives, successors, and assigns; and its joint ventures, subsidiaries, divisions, groups and affiliates in each case controlled by Novartis AG, and the respective directors, officers, employees, agents, representatives, successors, and assigns of each. Novartis shall not include the OTC Joint Venture.

- C. “Respondents” means Eli Lilly and Novartis, individually and collectively; *provided, however*, that from the later to occur of (i) the OTC Joint Venture Date, or (ii) the Closing Date, “Respondents” shall mean Eli Lilly.
- D. “Commission” means the Federal Trade Commission.
- E. “Decision and Order” means the:
1. Proposed Decision and Order contained in the Consent Agreement in this matter until the issuance of a final and effective Decision and Order by the Commission; and
  2. Final Decision and Order issued by the Commission following the issuance and service of a final Decision and Order by the Commission in this matter.
- F. “Canine Health Product Business(es)” means the Business of Respondent Novartis within the Geographic Territory specified in the Decision and Order related to each of the Canine Health Products to the extent that such Business is owned, controlled, or managed by the Respondents and the Assets related to such Business to the extent such Assets are owned by, controlled by, managed by, or licensed to, the Respondents.
- G. “Interim Monitor” means any monitor appointed pursuant to Paragraph III of this Order to Maintain Assets or Paragraph III of the Decision and Order.
- H. “Orders” means the Decision and Order and this Order to Maintain Assets.

## II.

**IT IS FURTHER ORDERED** that from the date this Order to Maintain Assets becomes final and effective:

- A. Until Respondents fully transfer and deliver the Canine Health Product Assets to an Acquirer, Respondents shall take such actions as are necessary to maintain the full economic viability, marketability and competitiveness of each of the related Canine Health Product Businesses, to minimize any risk of loss of competitive potential for such Canine Health Product Businesses, and to prevent the destruction, removal, wasting, deterioration, or impairment of such Canine Health Product Assets except for ordinary wear and tear. Respondents shall not sell, transfer, encumber or otherwise impair the Canine Health Product Assets (other than in the manner prescribed in the Decision and Order) nor take any action that lessens the full economic viability, marketability or competitiveness of the related Canine Health Product Businesses, *provided, however*, that a Respondent may continue to operate its Business related to any Retained Product in the ordinary course of business.

B. Until Respondents fully transfer and deliver the Canine Health Product Assets to an Acquirer, Respondents shall maintain the operations of the related Canine Health Product Businesses in the regular and ordinary course of business and in accordance with past practice (including regular repair and maintenance of the assets of such business) and/or as may be necessary to preserve the full economic marketability, viability, and competitiveness of such Canine Health Product Businesses and shall use their best efforts to preserve the existing relationships with the following: suppliers; vendors and distributors; High Volume Accounts; end-use customers; Agencies; employees; and others having business relations with each of the respective Canine Health Product Businesses. Respondents' responsibilities shall include, but are not limited to, the following:

1. providing the Canine Health Product Business with sufficient working capital to operate at least at current rates of operation, to meet all capital calls with respect to such business and to carry on, at least at their scheduled pace, all capital projects, business plans and promotional activities for such Canine Health Product Business;
2. continuing, at least at their scheduled pace, any additional expenditures for the Canine Health Product Business authorized prior to the date the Consent Agreement was signed by Respondents including, but not limited to, all research, Development, manufacturing, distribution, marketing and sales expenditures;
3. providing such resources as may be necessary to respond to competition against each of the Canine Health Products and/or to prevent any diminution in sales of each of the Canine Health Products during and after the Acquisition process and prior to the complete transfer and delivery of the related Canine Health Product Assets to an Acquirer; *provided, however*, that a Respondent may continue to operate its Business related to any Retained Product in the ordinary course of business;
4. providing such resources as may be necessary to maintain the competitive strength and positioning of each of the Canine Health Products at the related High Volume Accounts;
5. making available for use by the Canine Health Product Business funds sufficient to perform all routine maintenance and all other maintenance as may be necessary to, and all replacements of, the assets related to such Canine Health Product Business; and
6. providing such support services to the Canine Health Product Business as were being provided to the Canine Health Product Business by Respondents as of the date the Consent Agreement was signed by Respondents.

C. Until Respondents fully transfer and deliver each of the respective Canine Health Product Assets to an Acquirer, Respondents shall maintain a work force that is (i) at least as large in size (as measured in full time equivalents) as, and (ii) comparable in training, and expertise to, what has been associated with the Canine Health Products for the relevant Canine Health Product's last

fiscal year, *provided, however*, that a reduction in employees due the hiring of employees by the Acquirer shall not be considered a reduction in the work force.

D. Respondents shall:

1. for a period of one (1) year from the Closing Date, provide the Acquirer or its Manufacturing Designee with the opportunity to enter into employment contracts with the Canine Health Product Core Employees. Each of these periods is hereinafter referred to as the “Canine Health Product Core Employee Access Period(s);”
2. not later than the earlier of the following dates: (i) ten (10) days after notice by staff of the Commission to Respondents to provide the Product Employee Information; or (ii) ten (10) days after written request by an Acquirer or Proposed Acquirer, provide the Acquirer or Proposed Acquirer(s) with the Product Employee Information related to the Canine Health Product Core Employees *unless* the Law requires a mandatory notice period prior to the release of such information in which case the information shall be provided not later than ten (10) days after the expiration of the notice period. Failure by Respondents to provide the Product Employee Information for any Canine Health Product Core Employee within the time provided herein shall extend the Canine Health Product Core Employee Access Period(s) with respect to that employee in an amount equal to the delay; *provided, however*, that the provision of such information may be conditioned upon the Acquirer’s or Proposed Acquirer’s written confirmation that it will (i) treat the information as confidential and, more specifically, (ii) use the information solely to consider whether to provide or continue providing to Canine Health Product Core Employees the opportunity to enter into employment contracts during a Canine Health Product Core Employee Access Period, (iii) restrict access to the information to such of the Acquirer’s or Proposed Acquirer’s employees who need such access in connection with the specified and permitted use, and (iv) destroy or return the information without retaining copies at such time as the specified and permitted use ends;
3. during the Canine Health Product Core Employee Access Period, not interfere with the hiring or employing by the Acquirer or its Manufacturing Designee of the Canine Health Product Core Employees, and remove any impediments within the control of Respondents that may deter these employees from accepting employment with the Acquirer or its Manufacturing Designee, including, but not limited to, any noncompete or nondisclosure provision of employment with respect to a Canine Health Product or other contracts with Respondents that would affect the ability or incentive of those individuals to be employed by the Acquirer or its Manufacturing Designee. In addition, Respondents shall not make any counteroffer to such a Canine Health Product Core Employee who has received a written offer of employment from the Acquirer or its Manufacturing Designee;

*provided, however,* that, subject to the conditions of continued employment prescribed in this Order, this Paragraph shall not prohibit Respondents from continuing to employ any Canine Health Product Core Employee under the terms of that employee's employment with Respondents prior to the date of the written offer of employment from the Acquirer or its Manufacturing Designee to that employee;

4. until the Closing Date, provide all Canine Health Product Core Employees with reasonable financial incentives to continue in their positions and to research, Develop, manufacture and/or market the Canine Health Products consistent with past practices and/or as may be necessary to preserve the marketability, viability and competitiveness of the Canine Health Products and to ensure successful execution of the pre-Acquisition plans for the Canine Health Products. Such incentives shall include a continuation of all employee compensation and benefits offered by Respondents until the Closing Date(s) has occurred, including regularly scheduled raises, bonuses, and vesting of pension benefits (as permitted by Law); and
5. for a period of one (1) year from the Closing Date, not, directly or indirectly, solicit or otherwise attempt to induce any employee of the Acquirer or its Manufacturing Designee with any amount of responsibility related to a Canine Health Product ("Canine Health Product Employee") to terminate his or her employment relationship with the Acquirer or its Manufacturing Designee; or hire any Canine Health Product Employee;

*provided, however,* Respondents may hire any former Canine Health Product Employee whose employment has been terminated by the Acquirer or its Manufacturing Designee or who independently applies for employment with a Respondent, as long as that employee was not solicited in violation of the nonsolicitation requirements contained herein;

*provided further, however,* that this Paragraph does not require nor shall be construed to require Respondents to terminate the employment of any employee or to prevent Respondents from continuing to employ the Canine Health Product Core Employees in connection with the Acquisition;

*provided further, however,* that any Respondent may do the following: (i) advertise for employees in newspapers, trade publications or other media not targeted specifically at the Canine Health Product Employees; or (ii) hire a Canine Health Product Employee who contacts any Respondent on his or her own initiative without any direct or indirect solicitation or encouragement from any Respondent.

- E. Respondents, in consultation with the Acquirer, for the purposes of ensuring and orderly marketing and distribution transition, shall:
1. develop and implement a detailed transition plan to ensure that the commencement of the marketing, distribution and sale of the Canine Health Products is not delayed or impaired by the Respondents;
  2. designate employees of Respondents knowledgeable about the marketing, distribution and sale related to each of the Canine Health Products who will be responsible for communicating directly with the Acquirer, and the Interim Monitor (if one has been appointed), for the purposes of assisting in the transfer of the Canine Health Product Business to the Acquirer;
  3. maintain and manage inventory levels of the Canine Health Products in consideration of the marketing and distribution transition;
  4. continue to market, distribute and sell the Canine Health Products until the Acquirer assumes such functions;
  5. allow the Acquirer access at reasonable business hours to all Confidential Business Information and employees who possess or are able to locate such information for the purposes of identifying the books, records, and files that contain such Confidential Business Information pending the completed delivery of such Confidential Business Information to the Acquirer;
  6. provide the Acquirer with a listing of inventory levels (week of supply) for each customer (*i.e.*, retailer, group purchasing organization, wholesaler or distributor) on a regular basis and in a timely manner;
  7. provide the Acquirer with anticipated reorder dates for each customer in a timely manner; and
  8. establish projected time lines for accomplishing all tasks necessary to effect the marketing and distribution transition in an efficient and timely manner.
- F. Pending divestiture of the Canine Health Product Assets, Respondents shall:
1. not use, directly or indirectly, any Confidential Business Information that is exclusively related to Sentinel® Spectrum® (NADA 141-333) or Sentinel® Flavor Tabs® (NADA 141-084) other than as necessary to comply with the following:

- a. the requirements of this Orders;
  - b. Respondents' obligations to the Acquirer under the terms of any related Remedial Agreement; or
  - c. applicable Law;
2. not disclose or convey any Confidential Business Information that is exclusively related to Sentinel® Spectrum® (NADA 141-333) or Sentinel® Flavor Tabs® (NADA 141-084), directly or indirectly, to any Person except (i) the Acquirer, (ii) another Respondent, but solely for the purposes enumerated in Paragraph II.F.1; (iii) other Persons specifically authorized by the Acquirer to receive such information, (iv) the Commission, or (v) the Interim Monitor (if any has been appointed);
  3. not provide, disclose or otherwise make available, directly or indirectly, any Confidential Business Information exclusively related to Sentinel® Spectrum® (NADA 141-333) or Sentinel® Flavor Tabs® (NADA 141-084) to the Respondents' employees associated with the Business related to those Retained Products that are indicated for the prevention or treatment of parasite(s) in any Species; and
  4. institute procedures and requirements to ensure that the above-described employees:
    - a. do not provide, disclose or otherwise make available, directly or indirectly, any Confidential Business Information in contravention of this Order to Maintain Assets; and
    - b. do not solicit, access or use any Confidential Business Information that they are prohibited from receiving for any reason or purpose.
- G. Not later than thirty (30) days from the earlier of (i) the Closing Date or (ii) the date this Order to Maintain Assets is issued by the Commission, Respondents shall provide written notification of the restrictions on the use and disclosure of the Confidential Business Information related to the Canine Health Products by Respondents' personnel to all of their employees who (i) may be in possession of such Confidential Business Information or (ii) may have access to such Confidential Business Information.
- H. Respondents shall give the above-described notification by e-mail with return receipt requested or similar transmission, and keep a file of those receipts for one (1) year after the Closing Date. Respondents shall provide a copy of the notification to the relevant Acquirer. Respondents shall maintain complete records of all such notifications at Respondents' registered office within the United States and shall provide an officer's certification to the Commission stating that the acknowledgment program has been implemented and is being complied with. Respondents shall provide the relevant Acquirer with copies of all certifications, notifications and reminders sent to



Respondents' personnel.

- I. Respondents shall monitor the implementation by its employees and other personnel of all applicable restrictions with respect to Confidential Business Information, and take corrective actions for the failure of such employees and personnel to comply with such restrictions or to furnish the written agreements and acknowledgments required by this Order to Maintain Assets.
- J. The purpose of this Order to Maintain Assets is to maintain the full economic viability, marketability and competitiveness of the Canine Health Product Businesses within the Geographic Territory through their full transfer and delivery to an Acquirer, to minimize any risk of loss of competitive potential for the Canine Health Product Businesses within the Geographic Territory, and to prevent the destruction, removal, wasting, deterioration, or impairment of any of the Canine Health Product Assets except for ordinary wear and tear.

### **III.**

#### **IT IS FURTHER ORDERED** that:

- A. At any time after Respondents sign the Consent Agreement in this matter, the Commission may appoint a monitor ("Interim Monitor") to assure that Respondents expeditiously comply with all of their obligations and perform all of their responsibilities as required by the Orders and the Remedial Agreements.
- B. The Commission shall select the Interim Monitor, subject to the consent of Respondents, which consent shall not be unreasonably withheld. If Respondents have not opposed, in writing, including the reasons for opposing, the selection of a proposed Interim Monitor within ten (10) days after notice by the staff of the Commission to Respondents of the identity of any proposed Interim Monitor, Respondents shall be deemed to have consented to the selection of the proposed Interim Monitor.
- C. Not later than ten (10) days after the appointment of the Interim Monitor, Respondents shall execute an agreement that, subject to the prior approval of the Commission, confers on the Interim Monitor all the rights and powers necessary to permit the Interim Monitor to monitor Respondents' compliance with the relevant requirements of the Orders in a manner consistent with the purposes of the Orders.
- D. If an Interim Monitor is appointed, Respondents shall consent to the following terms and conditions regarding the powers, duties, authorities, and responsibilities of the Interim Monitor:
  - 1. The Interim Monitor shall have the power and authority to monitor Respondents' compliance with the divestiture and asset maintenance obligations and related requirements of the Orders, and shall exercise such power and authority and carry out the duties and responsibilities of the Interim Monitor in a manner consistent with the

purposes of the Orders and in consultation with the Commission.

2. The Interim Monitor shall act in a fiduciary capacity for the benefit of the Commission.
3. The Interim Monitor shall serve until the date of completion by the Respondents of the divestiture of all Canine Health Product Assets and the transfer and delivery of the related Product Manufacturing Technology in a manner that fully satisfies the requirements of this Order and, with respect to each Canine Health Product, until the earliest of: (i) the date the Acquirer (or the Acquirer's Manufacturing Designee(s)) is approved by the FDA to manufacture that Canine Health Product and able to manufacture that Canine Health Product in commercial quantities, in a manner consistent with cGMP, independently of the Respondents; (ii) the date the Acquirer notifies the Commission and Respondents of its intention to abandon its efforts to manufacture that Canine Health Product; (iii) the date of written notification from staff of the Commission that the Interim Monitor, in consultation with staff of the Commission, has determined that the relevant Acquirer has abandoned its efforts to manufacture that Canine Health Product;

*provided, however,* that, with respect to each Canine Health Product, the Interim Monitor's service shall not exceed four (4) years from the Closing Date *unless* the Commission decides to extend or modify this period as may be necessary or appropriate to accomplish the purposes of the Orders.

- E. Subject to any demonstrated legally recognized privilege, the Interim Monitor shall have full and complete access to Respondents' personnel, books, documents, records kept in the ordinary course of business, facilities and technical information, and such other relevant information as the Interim Monitor may reasonably request, related to Respondents' compliance with its obligations under the Orders, including, but not limited to, its obligations related to the relevant assets. Respondents shall cooperate with any reasonable request of the Interim Monitor and shall take no action to interfere with or impede the Interim Monitor's ability to monitor Respondents' compliance with the Orders.
- F. The Interim Monitor shall serve, without bond or other security, at the expense of Respondents, on such reasonable and customary terms and conditions as the Commission may set. The Interim Monitor shall have authority to employ, at the expense of Respondents, such consultants, accountants, attorneys and other representatives and assistants as are reasonably necessary to carry out the Interim Monitor's duties and responsibilities.
- G. Respondents shall indemnify the Interim Monitor and hold the Interim Monitor harmless against any losses, claims, damages, liabilities, or expenses arising out of, or in connection with, the performance of the Interim Monitor's duties, including all reasonable fees of counsel and other reasonable expenses incurred in connection with the preparations for, or defense of, any claim, whether or not resulting in any liability, except to the extent that such losses, claims, damages,

liabilities, or expenses result from gross negligence, willful or wanton acts, or bad faith by the Interim Monitor.

- H. Respondents shall report to the Interim Monitor in accordance with the requirements of the Orders and as otherwise provided in any agreement approved by the Commission. The Interim Monitor shall evaluate the reports submitted to the Interim Monitor by Respondents, and any reports submitted by each Acquirer with respect to the performance of Respondents' obligations under the Orders or the Remedial Agreement(s). Within thirty (30) days from the date the Interim Monitor receives these reports, the Interim Monitor shall report in writing to the Commission concerning performance by Respondents of their obligations under the Orders; *provided, however*, beginning ninety (90) days after Respondents have filed their final report pursuant to Paragraph VII.C. of the Decision and Order, and ninety (90) days thereafter, the Interim Monitor shall report in writing to the Commission concerning progress by each Acquirer toward obtaining FDA approval to manufacture each Canine Health Product and obtaining the ability to manufacture each Canine Health Product in commercial quantities, in a manner consistent with cGMP, independently of Respondents.
- I. Respondents may require the Interim Monitor and each of the Interim Monitor's consultants, accountants, attorneys and other representatives and assistants to sign a customary confidentiality agreement; *provided, however*, that such agreement shall not restrict the Interim Monitor from providing any information to the Commission.
- J. The Commission may, among other things, require the Interim Monitor and each of the Interim Monitor's consultants, accountants, attorneys and other representatives and assistants to sign an appropriate confidentiality agreement related to Commission materials and information received in connection with the performance of the Interim Monitor's duties.
- K. If the Commission determines that the Interim Monitor has ceased to act or failed to act diligently, the Commission may appoint a substitute Interim Monitor in the same manner as provided in this Paragraph.
- L. The Commission may on its own initiative, or at the request of the Interim Monitor, issue such additional orders or directions as may be necessary or appropriate to assure compliance with the requirements of the Orders.
- M. The Interim Monitor appointed pursuant to this Order to Maintain Assets may be the same person appointed as a Divestiture Trustee pursuant to the relevant provisions of the Decision and Order.

#### IV.

**IT IS FURTHER ORDERED** that within thirty (30) days after the date this Order to Maintain Assets is issued by the Commission, and every sixty (60) days thereafter until Respondents have fully complied with this Order to Maintain Assets and the Paragraphs that are enumerated in Paragraph VII.C. of the related Decision and Order, Respondents shall submit to the Commission a verified written report setting forth in detail the manner and form in which they intend to comply, are complying, and have complied with the Orders. Respondents shall submit at the same time a copy of their report concerning compliance with the Orders to the Interim Monitor, if any Interim Monitor has been appointed. Respondents shall include in their reports, among other things that are required from time to time, a detailed description of their efforts to comply with the relevant paragraphs of the Orders, including:

- A. a detailed description of all substantive contacts, negotiations, or recommendations related to (i) the divestiture and transfer of all relevant assets and rights, (ii) transitional services being provided by the Respondents to the relevant Acquirer, and (iii) the agreement(s) to Contract Manufacture; and
- B. a detailed description of the timing for the completion of such obligations.

*provided, however,* that, after the Decision and Order in this matter becomes final and effective, the reports due under this Order to Maintain Assets may be consolidated with, and submitted to the Commission at the same time as, the reports required to be submitted by Respondent pursuant to Paragraph VII of the Decision and Order.

#### V.

**IT IS FURTHER ORDERED** that Respondents shall notify the Commission at least thirty (30) days prior to:

- A. any proposed dissolution of a Respondent;
- B. any proposed acquisition, merger or consolidation of a Respondent; or
- C. any other change in a Respondent including, but not limited to, assignment and the creation or dissolution of subsidiaries, if such change might affect compliance obligations arising out of the Orders.

## VI.

**IT IS FURTHER ORDERED** that, for purposes of determining or securing compliance with this Order, and subject to any legally recognized privilege, and upon written request and upon five (5) days' notice to any Respondent made to its principal United States offices, registered office of its United States subsidiary, or its headquarters address, that Respondent shall, without restraint or interference, permit any duly authorized representative of the Commission:

- A. access, during business office hours of the Respondent and in the presence of counsel, to all facilities and access to inspect and copy all books, ledgers, accounts, correspondence, memoranda and all other records and documents in the possession or under the control of the Respondent related to compliance with this Order, which copying services shall be provided by the Respondent at the request of the authorized representative(s) of the Commission and at the expense of the Respondent; and
- B. to interview officers, directors, or employees of the Respondent, who may have counsel present, regarding such matters.

## VII.

**IT IS FURTHER ORDERED** that this Order to Maintain Assets shall terminate on the later of:

- A. three (3) days after the Commission withdraws its acceptance of the Consent Agreement pursuant to the provisions of Commission Rule 2.34, 16 C.F.R. § 2.34;
- B. the day after the divestiture of all of the Canine Health Product Assets, as required by and described in the Decision and Order, has been completed;
- C. the day after the Product Manufacturing Technology related to each Canine Health Product has been provided to the Acquirer in a manner consistent with the Technology Transfer Standards and the Interim Monitor, in consultation with Commission staff and the Acquirer(s), notifies the Commission that all assignments, conveyances, deliveries, grants, licenses, transactions, transfers and other transitions related to the provision of the Product Manufacturing Technology are complete; or
- D. the day the Commission otherwise directs that this Order to Maintain Assets is terminated.

By the Commission.

SEAL:  
ISSUED: December 19, 2014

Donald S. Clark  
Secretary