the ESA by conducting an intra-Service section 7 consultation on anticipated ITP actions. We will use the results of this consultation, in combination with the above findings, in our final analysis to determine whether to issue the ITP. If we determine that all requirements are met, we will issue an ITP under section 10(a)(1)(B) of the ESA to the applicant for the take of the covered species, incidental to otherwise lawful covered activities. We will make the final permit decision no sooner than 30 days after the date of this notice.

Authority

We provide this notice in accordance with the requirements of section 10 of the ESA and NEPA and their implementing regulations (50 CFR 17.32 and 40 CFR 1506.6, respectively).

Dated: June 20, 2017.

Theresa E. Rabot,

Deputy Regional Director, Pacific Region, U.S. Fish and Wildlife Service, Portland, Oregon.

[FR Doc. 2017–17082 Filed 8–11–17; 8:45 am]

BILLING CODE 4333–15–P

INTERNATIONAL TRADE COMMISSION

[Investigation No. 337–TA–1066]

Certain Recombinant Factor IX Products; Institution of Investigation

AGENCY: U.S. International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that a complaint was filed with the U.S. International Trade Commission on July 7, 2017, under section 337 of the Tariff Act of 1930, as amended, on behalf of Bioverativ Inc. of Waltham, Massachusetts; Bioverativ Therapeutics Inc. of Waltham, Massachusetts; and Bioverativ U.S. LLC of Waltham, Massachusetts. A supplement to the complaint was filed on July 14, 2017. The complaint alleges violations of section 337 based upon the importation into the United States, the sale for importation, and the sale within the United States after importation of certain recombinant Factor IX products by reason of infringement of certain claims of U.S. Patent No. 9,670,475 (''the '475 patent''); U.S. Patent No. 9,623,091 (''the '091 patent''); and U.S. Patent No. 9,629,903 (''the '903 patent''). The complaint further alleges that an industry in the United States exists as required by the applicable Federal Statute.

The complainants request that the Commission institute an investigation and, after the investigation, issue a limited exclusion order and cease and desist orders.

ADDRESSES: The complaint, except for any confidential information contained therein, is available for inspection during official business hours (8:45 a.m. to 5:15 p.m.) in the Office of the Secretary, U.S. International Trade Commission, 500 E Street SW., Room 112, Washington, DC 20436, telephone (202) 205–2000. Hearing impaired individuals are advised that information on this matter can be obtained by contacting the Commission's TDD terminal on (202) 205–1810. Persons with mobility impairments who will need special assistance in gaining access to the Commission should contact the Office of the Secretary at (202) 205–2000. General information concerning the Commission may also be obtained by accessing its internet server at *https://www.usitc.gov.* The public record for this investigation may be viewed on the Commission's electronic docket (EDIS) at *https://edis.usitc.gov.*

FOR FURTHER INFORMATION CONTACT: Pathenia M. Proctor, The Office of Unfair Import Investigations, U.S. International Trade Commission, telephone (202) 205–2560.

SUPPLEMENTARY INFORMATION:

Authority: The authority for institution of this investigation is contained in section 337 of the Tariff Act of 1930, as amended, 19 U.S.C. 1337 and in section 210.10 of the Commission's Rules of Practice and Procedure, 19 CFR 210.10 (2017).

Scope of Investigation: Having considered the complaint, the U.S. International Trade Commission, on August 8, 2017, *ordered that*—

(1) Pursuant to subsection (b) of section 337 of the Tariff Act of 1930, as amended, an investigation be instituted to determine whether there is a violation of subsection (a)(1)(B) of section 337 in the importation into the United States, the sale for importation, or the sale within the United States after importation of certain recombinant Factor IX products by reason of infringement of one or more of claims 1, 2, 4–19, 24, 25, 29, and 34 of the '475 patent; claims 1–7, 11–16, 18, 19, 21, and 23–27 of the '091 patent; and claims 1–10, 13–15, and 17–28 of the '903 patent; and whether an industry in the United States exists as required by subsection (a)(2) of section 337;

(2) Pursuant to Commission Rule 210.50(b)(1), 19 CFR 210.50(b)(1), the presiding Administrative Law Judge shall take evidence or other information and hear arguments from the parties or other interested persons with respect to the public interest in this investigation, as appropriate, and provide the Commission with findings of fact and a recommended determination on this issue, which shall be limited to the statutory public interest factors set forth in 19 U.S.C. 1337(d)(1), (f)(1), (g)(1);

(3) For the purpose of the investigation so instituted, the following are hereby named as parties upon which this notice of investigation shall be served:

(a) The complainants are: Bioverativ Inc., 225 Second Avenue, Waltham, MA 02451. Bioverativ Therapeutics Inc., 225 Second Avenue, Waltham, MA 02451. Bioverativ U.S. LLC, 225 Second Avenue, Waltham, MA 02451.

(b) The respondents are the following entities alleged to be in violation of section 337, and are the parties upon which the complaint is to be served: CSL Behring LLC, 1020 First Avenue, King of Prussia, PA 19406. CSL Behring GmbH, Emil-von-Behring-Strasse 76, Marburg, Hessen 35041 Germany. CSL Behring Recombinant Facility AG, Wankdorfstrasse 10, Bern, Bern 3014 Switzerland.

(c) The Office of Unfair Import Investigations, U.S. International Trade Commission, 500 E Street SW., Suite 401, Washington, DC 20436; and

(4) For the investigation so instituted, the Chief Administrative Law Judge, U.S. International Trade Commission, shall designate the presiding Administrative Law Judge.

Responses to the complaint and the notice of investigation must be submitted by the named respondents in accordance with section 210.13 of the Commission's Rules of Practice and Procedure, 19 CFR 210.13. Pursuant to 19 CFR 201.16(e) and 210.13(a), such responses will be considered by the Commission if received not later than 20 days after the date of service by the Commission of the complaint and the notice of investigation. Extensions of time for submitting responses to the complaint and the notice of investigation will not be granted unless good cause therefor is shown.

Failure of a respondent to file a timely response to each allegation in the complaint and in this notice may be deemed to constitute a waiver of the right to appear and contest the allegations of the complaint and this notice, and to authorize the administrative law judge and the Commission, without further notice to the respondent, to find the facts to be as alleged in the complaint and this notice and to enter an initial determination and a final determination containing such findings, and may result in the issuance of an exclusion order or a cease

and desist order or both directed against the respondent.

By order of the Commission.

Issued: August 8, 2017.

Lisa R. Barton,

Secretary to the Commission.

[FR Doc. 2017–17058 Filed 8–11–17; 8:45 am]

BILLING CODE P

INTERNATIONAL TRADE COMMISSION

[Investigation No. 337–TA–1065]

Certain Mobile Electronic Devices and Radio Frequency and Processing Components Thereof; Institution of Investigation

AGENCY: U.S. International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that a complaint was filed with the U.S. International Trade Commission on July 7, 2017, under section 337 of the Tariff Act of 1930, as amended, on behalf of Qualcomm Incorporated of San Diego, California. A supplement was filed on July 7, 2017. The complaint, as supplemented, alleges violations of section 337 based upon the importation into the United States, the sale for importation, and the sale within the United States after importation of certain mobile electronic devices and radio frequency and processing components thereof by reason of infringement of U.S. Patent No. 8,633,936 (''the '936 patent''); U.S. Patent No. 8,698,558 (''the '558 patent''); U.S. Patent No. 8,487,658 (''the '658 patent''); U.S. Patent No. 8,838,949 (''the '949 patent''); U.S. Patent No. 9,535,490 (''the '490 patent''); and U.S. Patent No. 9,608,675 (''the '675 patent''). The complaint further alleges that an industry in the United States exists as required by the applicable Federal Statute.

The complainant requests that the Commission institute an investigation and, after the investigation, issue a limited exclusion order and a cease and desist order.

ADDRESSES: The complaint, except for any confidential information contained therein, is available for inspection during official business hours (8:45 a.m. to 5:15 p.m.) in the Office of the Secretary, U.S. International Trade Commission, 500 E Street SW., Room 112, Washington, DC 20436, telephone (202) 205–2000. Hearing impaired individuals are advised that information on this matter can be obtained by contacting the Commission's TDD terminal on (202) 205–1810. Persons with mobility impairments who will need special assistance in gaining access to the Commission should contact the Office of the Secretary at (202) 205–2000. General information concerning the Commission may also be obtained by accessing its internet server at *https://www.usitc.gov.* The public record for this investigation may be viewed on the Commission's electronic docket (EDIS) at *https://edis.usitc.gov.*

FOR FURTHER INFORMATION CONTACT: Pathenia M. Proctor, The Office of Unfair Import Investigations, U.S. International Trade Commission, telephone (202) 205–2560.

SUPPLEMENTARY INFORMATION:

Authority: The authority for institution of this investigation is contained in section 337 of the Tariff Act of 1930, as amended, 19 U.S.C. 1337 and in section 210.10 of the Commission's Rules of Practice and Procedure, 19 CFR 210.10 (2017).

Scope of Investigation: Having considered the complaint, the U.S. International Trade Commission, on August 8, 2017, *ordered that*—

(1) Pursuant to subsection (b) of section 337 of the Tariff Act of 1930, as amended, an investigation be instituted to determine whether there is a violation of subsection (a)(1)(B) of section 337 in the importation into the United States, the sale for importation, or the sale within the United States after importation of certain mobile electronic devices and radio frequency and processing components thereof by reason of infringement of one or more of claims 1–27, 29, 38, 49, 55–60, 67, and 68 of the '936 patent; claims 1 and 6–20 of the '558 patent; claims 9, 10, 12, 14, and 20–22 of the '658 patent; claims 1–8, 10–14, 16, 20, and 22 of the '949 patent; claims 1–6, 8, 10, 16, 17, and 31 of the '490 patent; and claims 1–3 and 7–14 of the '675 patent; and whether an industry in the United States exists as required by subsection (a)(2) of section 337;

(2) Pursuant to Commission Rule 210.50(b)(1), 19 CFR 210.50(b)(1), the presiding Administrative Law Judge shall take evidence or other information and hear arguments from the parties or other interested persons with respect to the public interest in this investigation, as appropriate, and provide the Commission with findings of fact and a recommended determination on this issue, which shall be limited to the statutory public interest factors set forth in 19 U.S.C. 1337(d)(1), (f)(1), (g)(1);

(3) For the purpose of the investigation so instituted, the following are hereby named as parties upon which this notice of investigation shall be served:

(a) The complainant is: Qualcomm Incorporated, 5775 Morehouse Drive, San Diego, CA 92121.

(b) The respondent is the following entity alleged to be in violation of section 337, and is the party upon which the complaint is to be served: Apple Inc., 1 Infinite Loop, Cupertino, CA 95014.

(c) The Office of Unfair Import Investigations, U.S. International Trade Commission, 500 E Street SW., Suite 401, Washington, DC 20436; and

(4) For the investigation so instituted, the Chief Administrative Law Judge, U.S. International Trade Commission, shall designate the presiding Administrative Law Judge.

Responses to the complaint and the notice of investigation must be submitted by the named respondent in accordance with section 210.13 of the Commission's Rules of Practice and Procedure, 19 CFR 210.13. Pursuant to 19 CFR 201.16(e) and 210.13(a), such responses will be considered by the Commission if received not later than 20 days after the date of service by the Commission of the complaint and the notice of investigation. Extensions of time for submitting responses to the complaint and the notice of investigation will not be granted unless good cause therefor is shown.

Failure of the respondent to file a timely response to each allegation in the complaint and in this notice may be deemed to constitute a waiver of the right to appear and contest the allegations of the complaint and this notice, and to authorize the administrative law judge and the Commission, without further notice to the respondent, to find the facts to be as alleged in the complaint and this notice and to enter an initial determination and a final determination containing such findings, and may result in the issuance of an exclusion order or a cease and desist order or both directed against the respondent.

By order of the Commission.

Issued: August 8, 2017.

Lisa R. Barton,

Secretary to the Commission.

[FR Doc. 2017–17057 Filed 8–11–17; 8:45 am]

BILLING CODE 7020–02–P