supplemented, alleged violations of section 337 of the Tariff Act of 1930, as amended, 19 U.S.C. 1337, in the importation into the United States, the sale for importation, or the sale within the United States after importation of certain core orientation systems, products containing core orientation systems, components thereof, and methods of using the same by reason of infringement of claims 16–18, 22, and 23 of U.S. Patent No. 7,584,055. *Id.* The complaint further alleged that an industry in the United States exists or is in the process of being established. *Id.* The Commission's notice of investigation named as respondents: Boart Longyear Group Ltd. of West Valley City, UT; Boart Longyear Limited of Australia; Boart Longyear Company of West Valley City, UT; Boart Longyear Manufacturing and Distribution Inc. of West Valley City, UT; Longyear TM, Inc. of West Valley City, UT; Globaltech Corporation Pty Ltd. of Australia; Globaltech Pty Ltd. of Australia; Granite Construction Incorporated of Watsonville, CA; and International Directional Services LLC of Chandler, AZ. The Office of Unfair Import Investigations (''OUII'') is also named as a party in this investigation. *Id.* at 19704–05.

On December 14, 2022, the Commission determined not to review an ID (Order No. 28) granting a joint motion to terminate respondent Granite Construction Incorporated from the investigation based on withdrawal of the complaint.

On November 30, 2022, Complainants and the Remaining Respondents jointly moved to terminate and stay the investigation based on a consent order stipulation. The joint motion included a consent order stipulation, a proposed consent order, and a settlement agreement. OUII filed a response supporting the joint motion.

On December 19, 2022, the ALJ issued an ID (Order No. 31) granting the joint motion to terminate the investigation with respect to the Remaining Respondents based on the entry of a consent order. The ID found that the consent order stipulation and proposed consent order conform with Commission Rule 210.21(c)(3) and (4) (19 CFR 210.21(c)(3) and (4)). The ID also found that termination of the investigation with respect to the Remaining Respondents would not be contrary to the public interest. No petitions for review were filed.

The Commission has determined not to review the subject ID and to issue a consent order against respondents Boart Longyear Group Ltd.; Boart Longyear Limited; Boart Longyear Company; Boart Longyear Manufacturing and Distribution Inc.; Longyear TM, Inc.; Globaltech Corporation Pty Ltd.; Globaltech Pty Ltd.; and International Directional Services LLC. The investigation is terminated in its entirety.

The Commission vote for this determination took place on January 17, 2023.

The authority for the Commission's determination is contained in section 337 of the Tariff Act of 1930, as amended, 19 U.S.C. 1337, and in part 210 of the Commission's Rules of Practice and Procedure, 19 CFR part 210.

By order of the Commission.

Issued: January 17, 2023.

Katherine Hiner,

Acting Secretary to the Commission.

[FR Doc. 2023–01105 Filed 1–20–23; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Antitrust Division

Granting of Requests for Early Termination of the Waiting Period Under the Premerger Notification Rules

Section 7A of the Clayton Act, 15 U.S.C. 18a, as added by Title II of the Hart-Scott-Rodino Antitrust Improvements Act of 1976, requires persons contemplating certain mergers or acquisitions to give the Federal Trade Commission and the Assistant Attorney General advance notice and to wait designated periods before consummation of such plans. Section 7A(b)(2) of the Act permits the agencies, in individual cases, to terminate this waiting period prior to its expiration and requires that notice of this action be published in the **Federal Register**. The following transactions were granted early termination—on the date indicated—of the waiting period provided by law and the premerger notification rules. The listing includes the transaction number and the parties to the transaction. The Federal Trade Commission and the Assistant Attorney General for the Antitrust Division of the Department of Justice made the grants. Neither agency intends to take any action with respect to this proposed acquisitions during the applicable waiting period.

EARLY TERMINATION GRANTED

11/04/2022		
20221286	G	Flatlake Privatshftung; Global Tungsten & Powders Corp.; Allan C. Bir, Jr.; Mi-Tech Tungsten Metals, LLC B6 Manufacturing, LLC.
01/05/2023		
20222859	G	Semtech Corporation; 13548597 Canada, Inc.; Sierra Wireless, Inc.

Suzanne Morris,

Deputy Directory of Civil Enforcement Operations, Antitrust Division, Department of Justice.

[FR Doc. 2023–01198 Filed 1–20–23; 8:45 am]

BILLING CODE 4410–11–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1134]

Bulk Manufacturer of Controlled Substances Application: Janssen Pharmaceuticals Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Janssen Pharmaceuticals Inc., has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to Supplementary Information listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may submit electronic comments on or objections to the issuance of the proposed registration on or before March 24, 2023. Such persons may also file a written request for a hearing on the application on or before March 24, 2023.

ADDRESSES: The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to *https://www.regulations.gov* and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public

view on *https://www.regulations.gov.* If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on December 1, 2022, Janssen Pharmaceuticals Inc., 1440 Olympic Drive, Buildings 1–5 & 7–14, Athens, Georgia 30601–1645, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Methylphenidate	1724	II
Hydromorphone	9150	II
Hydrocodone	9193	II
Oripavine	9330	II
Thebaine	9333	II
Tapentadol	9780	II

The company plans to bulk manufacture the listed controlled substances for the internal use intermediates or for sale to its customers. No other activities for these drug codes are authorized for this registration.

Matthew Strait,

Deputy Assistant Administrator.

[FR Doc. 2023–01205 Filed 1–20–23; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1133]

Importer of Controlled Substances Application: Mylan Pharmaceuticals, Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Mylan Pharmaceuticals, Inc. has applied to be registered as an importer of basic class(es) of controlled substance(s). Refer to **SUPPLEMENTARY INFORMATION** listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may submit electronic comments on or objections to the issuance of the proposed registration on or before February 22, 2023. Such persons may also file a written request for a hearing on the application on or before February 22, 2023.

ADDRESSES: The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to *https://www.regulations.gov* and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on *https://www.regulations.gov.* If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment. All requests for a hearing must be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA **Federal Register** Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on December 6, 2022, Mylan Pharmaceuticals, Inc., 2898 Manufacturers Road, Greensboro, North Carolina 27406–4600, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Remifentanil	9739	II

The company plans to import the above listed controlled substance in finished dosage from for commercial distribution to its customers. No other activities for these drug codes are authorized for this registration.

Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under 21 U.S.C. 952(a)(2). Authorization will not extend to the import of Food and Drug Administration-approved or non-approved finished dosage forms for commercial sale.

Matthew Strait,

Deputy Assistant Administrator.

[FR Doc. 2023–01208 Filed 1–20–23; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1129]

Bulk Manufacturer of Controlled Substances Application: Siegfried USA, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Siegfried USA, LLC has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to **SUPPLEMENTARY INFORMATION** listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may submit electronic comments on or objections to the issuance of the proposed registration on or before March 24, 2023. Such persons may also file a written request for a hearing on the application on or before March 24, 2023.

ADDRESSES: The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to *https://www.regulations.gov* and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on *https://www.regulations.gov.* If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on November 8, 2022, Siegfried USA, LLC, 33 Industrial Park Road, Pennsville, New Jersey 08070–3244, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Gamma Hydroxybutyric Acid	2010	I
Noroxymorphone	9145	I
Hydromorphinol	9301	I
Amphetamine	1100	II
Lisdexamfetamine	1205	II
Methylphenidate	1724	II
Amobarbital	2125	II
Pentobarbital	2270	II
Secobarbital	2315	II
Codeine	9050	II
Oxycodone	9143	II