violation of section 337 by the eight defaulting respondents and for a recommendation that the Commission issue a general exclusion order (''GEO'') and cease and desist orders (''CDOs''). *See* Complainants' Motion for Summary Determination of Violation and for Recommended Determination on Remedy and Bonding. Skull Shaver accused Yiwu Xingye and Yiwu City of infringing claims 1–3 of the '528 patent and the claim of the '504 design patent. *Id.* at 5. It accused the other respondents of infringing only claim 1 of the '528 patent. *Id.* On June 7, 2021, OUII filed a response in support of Skull Shaver's motion. *See* Commission Investigative Staff's Response to Skull Shaver's Motion for Summary Determination of Violation. No respondent filed a response to Skull Shaver's motion.

On September 23, 2021, OUII filed a notice of supplemental authority concerning the domestic industry requirement. On September 28, 2021, the ALJ issued an order (Order No. 31) ordering certain supplementation of Skull Shaver's domestic industry analysis. On October 14, 2021, Skull Shaver submitted its supplement in response to Order No. 31. No other responses to Order No. 31 were filed. On November 18, 2021, the ALJ granted-in-part Skull Shaver's motion for summary determination as the subject ID.

The ID found that Skull Shaver owns the asserted patents, and that those patents are valid and enforceable. ID at 3. The ID further found that although all respondents imported, sold for importation, or sold within the United States after importation at least one accused article, the only respondents whose articles infringe the asserted patents are Yiwu Xingye and Yiwu City. *Id.* at 3–4. The ID found no infringement as to the other respondents, whose products lack a second recess, *see* ID at 51–52, in view of the ALJ's construction of ''recesses'' as ''indentations that are substantially concave surfaces,'' *id.* at 16 (citation omitted), and Skull Shaver's forfeiture of an infringement theory under the doctrine of equivalents, *id.* at 50 n.7. The ID found that personal jurisdiction is not necessary over each defaulting respondent, but that the defaulting respondents waived any opportunity to contest the allegation that personal jurisdiction exists. *Id.* The ID further found that Skull Shaver meets the technical prong and the economic prong of the domestic industry requirement. *Id.* at 4. As to remedy, the RD found that there is a widespread pattern of unauthorized use of the asserted patents and it is difficult to identify the source of these products; and that a GEO is necessary to prevent circumvention. *Id.* at 4. The RD also recommended issuance of CDOs against the two infringing respondents, who are presumed to maintain domestic inventories. RD at 80–81. The RD recommended a bond rate of one hundred percent (100%) because complete pricing information is not available. RD at 82.

No petitions for review of the ID were filed. The Commission determined to review the ID's findings concerning the economic prong of the domestic industry requirement, and not to review the ID's findings on other issues. Notice, 87 FR 990, 991 (Jan. 7, 2022). The review notice solicited written submissions, including on remedy, the public interest, and bonding from the parties, interested government agencies, and the public. *Id.*

In response to the Commission notice, Skull Shaver and OUII each filed an opening submission and a reply. No other parties filed submissions.

On review, the Commission has determined to affirm the ID's finding that Skull Shaver has satisfied the economic prong of the domestic industry requirement, and the Commission thereby affirms the ID's finding of a violation of section 337.

The Commission finds that the RD's recommended remedy is appropriate for the reasons set forth in the attached opinion. Accordingly, the Commission finds that the appropriate remedy is: (1) A general exclusion order prohibiting the entry of certain electric shavers and components and accessories thereof; and (2) cease and desist orders directed to Yiwu Xingye and Yiwu City. The Commission has determined that the public interest factors enumerated in section 337(d), (f), and/or (g), 19 U.S.C. 1337(d), (f), (g), do not preclude the issuance of the GEO or the CDOs.

The Commission has determined that a bond in the amount of one hundred percent (100%) of the entered value of the subject articles is required during the Presidential review period, 19 U.S.C. 1337(j) for the reasons set forth in the RD and the attached Commission Opinion. The investigation is hereby terminated.

While temporary remote operating procedures are in place in response to COVID–19, the Office of the Secretary is not able to serve parties that have not retained counsel or otherwise provided a point of contact for electronic service. Accordingly, pursuant to Commission Rules 201.16(a) and 210.7(a)(1) (19 CFR 201.16(a), 210.7(a)(1)), the Commission orders that the Complainant complete service for any party without a method of electronic service noted on the attached Certificate of Service and shall file proof of service on the Electronic Document Information System (EDIS).

The Commission's orders and opinion were delivered to the President and the United States Trade Representative on the day of their issuance.

The Commission vote for these determinations took place on March 17, 2022.

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: March 17, 2022.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2022–06122 Filed 3–22–22; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–985]

Bulk Manufacturer of Controlled Substances Application: Johnson Matthey Pharmaceutical Materials, Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Johnson Matthey Pharmaceutical Materials, 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 May 23, 2022. Such persons may also file a written request for a hearing on the application on or before May 23, 2022.

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 February 28, 2022, Johnson Matthey Pharmaceutical Materials, Inc., 2003 Nolte Drive, West Deptford, New Jersey 08066–1742, 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
Marihuana	7360	I
Tetrahydrocannabinols	7370	I
Noroxymorphone	9145	I
Difenoxin	9168	I
Amphetamine	1100	II
Methamphetamine	1105	II
Lisdexamfetamine	1205	II
Methylphenidate	1724	II
Nabilone	7379	II
ANPP (4-Anilino-N-phenethyl-4-piperidine).	8333	II
Norfentanyl (N-phenyl-N-(piperidin-4-yl) propionamide).	8366	II
Cocaine	9041	II
Codeine	9050	II
Dihydrocodeine	9120	II
Oxycodone	9143	II
Hydromorphone	9150	II
Diphenoxylate	9170	II
Ecgonine	9180	II
Hydrocodone	9193	II
Levorphanol	9220	II
Meperidine	9230	II
Methadone	9250	II
Methadone intermediate	9254	II
Morphine	9300	II
Thebaine	9333	II
Opium tincture	9630	II
Oxymorphone	9652	II
Noroxymorphone	9668	II
Alfentanil	9737	II
Remifentanil	9739	II
Sufentanil	9740	II
Tapentadol	9780	II
Fentanyl	9801	II

The company plans to bulk manufacture the listed controlled substances for the internal use intermediates or for sale to its customers. In reference to drug codes 7360 (Marihuana), and 7370 (Tetrahydrocannabinols), the company plans to bulk manufacture these drugs as synthetic. No other activities for these drug codes are authorized for this registration.

Matthew J. Strait,

Deputy Assistant Administrator.

[FR Doc. 2022–06159 Filed 3–22–22; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–979]

Importer of Controlled Substances Application: Sharp Clinical Services, Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Sharp Clinical Services, 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 April 22, 2022. Such persons may also file a written request for a hearing on the application on or before April 22, 2022.

ADDRESSES: The DEA 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 February 2, 2022, Sharp Clinical Services, Inc. 2400 Baglyos Circle, Bethlehem, Pennsylvania 18020–8024, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Gamma Hydroxybutyric Acid.	2010	I
3,4-Methylenedioxymethamphetamine.	7405	I

The company plans to import the listed control substances for clinical trials. No other activity for these drug codes is 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 J. Strait,

Deputy Assistant Administrator.

[FR Doc. 2022–06161 Filed 3–22–22; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–984]

Bulk Manufacturer of Controlled Substances Application: Siemens Healthcare Diagnostics, Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Siemens Healthcare Diagnostics, 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 May 23, 2022. Such persons may also file a written request for a hearing on the application on or before May 23, 2022.

ADDRESSES: The Drug Enforcement Administration (DEA) requires that all comments be submitted electronically