Written submissions on the public interest must be filed no later than by close of business, eight calendar days after the date of publication of this notice in the **Federal Register**. There will be further opportunities for comment on the public interest after the issuance of any final initial determination in this investigation. Any written submissions on other issues must also be filed by no later than the close of business, eight calendar days after publication of this notice in the **Federal Register**. Complainant may file replies to any written submissions no later than three calendar days after the date on which any initial submissions were due, notwithstanding § 201.14(a) of the Commission's Rules of Practice and Procedure. No other submissions will be accepted, unless requested by the Commission. Any submissions and replies filed in response to this Notice are limited to five (5) pages in length, inclusive of attachments.

Persons filing written submissions must file the original document electronically on or before the deadlines stated above. Submissions should refer to the docket number (''Docket No. 3866'') in a prominent place on the cover page and/or the first page. (*See* Handbook for Electronic Filing Procedures, Electronic Filing Procedures).[1] Please note the Secretary's Office will accept only electronic filings during this time. Filings must be made through the Commission's Electronic Document Information System (EDIS, *https://edis.usitc.gov*.) No in-person paper-based filings or paper copies of any electronic filings will be accepted until further notice. Persons with questions regarding filing should contact the Secretary at *EDIS3Help@usitc.gov*.

Any person desiring to submit a document to the Commission in confidence must request confidential treatment. All such requests should be directed to the Secretary to the Commission and must include a full statement of the reasons why the Commission should grant such treatment. *See* 19 CFR 201.6. Documents for which confidential treatment by the Commission is properly sought will be treated accordingly. All information, including confidential business information and documents for which confidential treatment is properly sought, submitted to the Commission for purposes of this Investigation may be disclosed to and used: (i) by the Commission, its employees and Offices, and contract personnel (a) for developing or maintaining the records of this or a related proceeding, or (b) in internal investigations, audits, reviews, and evaluations relating to the programs, personnel, and operations of the Commission including under 5 U.S.C. Appendix 3; or (ii) by U.S. government employees and contract personnel,[2] solely for cybersecurity purposes. All nonconfidential written submissions will be available for public inspection at the Office of the Secretary and on EDIS.[3]

This action is taken under the authority of section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and of §§ 201.10 and 210.8(c) of the Commission's Rules of Practice and Procedure (19 CFR 201.10, 210.8(c)).

By order of the Commission.

Issued: December 16, 2025.

Lisa R. Barton,

Secretary to the Commission.

[FR Doc. 2025–23426 Filed 12–18–25; 8:45 am]

BILLING CODE 7020–02–P

JUDICIAL CONFERENCE OF THE UNITED STATES

Advisory Committee on Civil Rules; Hearing of the Judicial Conference

AGENCY: Judicial Conference of the United States.

ACTION: Advisory Committee on Civil Rules; notice of cancellation of open hearing.

SUMMARY: The following public hearing on proposed amendments to the Federal Rules of Civil Procedure has been canceled: Civil Rules Hearing on January 13, 2026.

DATES: January 13, 2026.

FOR FURTHER INFORMATION CONTACT: Carolyn A. Dubay, Esq., Chief Counsel, Rules Committee Staff, Administrative Office of the U.S. Courts, Thurgood Marshall Federal Judiciary Building, One Columbus Circle NE, Suite 7–300, Washington, DC 20544, Phone (202) 502–1820, *RulesCommittee_Secretary@ao.uscourts.gov.*

SUPPLEMENTARY INFORMATION: The announcement for this hearing was previously published in the **Federal Register** on July 14, 2025 at 90 FR 31242.

(Authority: 28 U.S.C. 2073.)

Dated: December 17, 2025.

Shelly L. Cox,

Management Analyst, Rules Committee Staff.

[FR Doc. 2025–23482 Filed 12–18–25; 8:45 am]

BILLING CODE 2210–55–P

DEPARTMENT OF JUSTICE

Antitrust Division

Notice Pursuant to the Defense Production Act of 1950

AGENCY: Antitrust Division, U.S. Department of Justice.

ACTION: Notice of review of voluntary agreement.

SUMMARY: Notice is hereby given pursuant to section 708 of the Defense Production Act of 1950 (''DPA''), that the Assistant Attorney General finds, with respect to the Implementing Voluntary Agreements Under the Defense Production Act (''Voluntary Agreement'') proposed by the Department of Energy (''DOE''), that the purposes of section 708(c)(1) of the DPA may not reasonably be achieved through a voluntary agreement having less anticompetitive effects or without any voluntary agreement. Given this finding, the proposed Voluntary Agreement may become effective following the publication of this notice.

SUPPLEMENTARY INFORMATION: Under the DPA, DOE may enter into plans with representatives of private industry for the purpose of improving the efficiency with which private firms contribute to the national defense when conditions exist that may pose a direct threat to the national defense or its preparedness. Such arrangements are generally known as ''voluntary agreements.'' Participants in an existing voluntary agreement may adopt documented methods, known as ''plans of action,'' to implement that voluntary agreement. A defense to actions brought under the antitrust laws is available to each participant acting within the scope of a voluntary agreement and plan of action that has come into force under the DPA.

The DPA requires that each proposed plan of action be reviewed by the Attorney General prior to becoming effective. If, after consulting with the Chair of the Federal Trade Commission, the Attorney General finds that the purposes of the DPA's plans of action provision ''may not reasonably be achieved through a . . . voluntary agreement having less anticompetitive effects or without any . . . voluntary agreement,'' the voluntary agreement may become effective. 50 U.S.C. 4558(f)(1)(B). All functions which the

[1] Handbook for Electronic Filing Procedures: *https://www.usitc.gov/documents/handbook_on_filing_procedures.pdf.*

[2] All contract personnel will sign appropriate nondisclosure agreements.

[3] Electronic Document Information System (EDIS): *https://edis.usitc.gov.*

Attorney General is required or authorized to perform by section 708 of the DPA have been delegated to the Assistant Attorney General, Antitrust Division. 28 CFR 0.40(l).

Executive Order 14,302 ''*Reinvigorating the Nuclear Fuel Base*'', 90 FR 22595 (''E.O. 14,302'') required the Secretary of Energy, in coordination with the Attorney General and the Chairman of the Federal Trade Commission, to utilize authority provided to the President in section 708(c)(1) of the Defense Production Act to seek voluntary agreements with domestic nuclear energy companies to provide for the national defense. The purpose of the proposed Voluntary Agreement is to establish a consortium and plans of action to ensure that the domestic nuclear fuel supply chain capacity is available to enable the continued reliable operation of the Nation's existing and future nuclear reactors. The phases of the domestic nuclear fuel supply chain that will be addressed in the consortium and plans of action include milling, conversion, enrichment, deconversion, fabrication, recycling and reprocessing, end users, and Uranium Fuel Infrastructure Resilience Mechanism (''UFIRM''). The consortium will allow for consultation with domestic nuclear energy companies to discuss and implement methods to enhance the capability to manage spent nuclear fuel to ensure the continued reliable operation of domestic nuclear reactors. DOE has certified that the proposed Voluntary Agreement is necessary to carry out its purpose, as specified in E.O. 14,302.

DOE requested that the Assistant Attorney General, Antitrust Division, pursuant to the Attorney General's delegation of authority under 28 CFR 0.40(i), issue a finding that the proposed Voluntary Agreement satisfies the statutory criteria set forth in 50 U.S.C. 4558(f)(1)(B). The Assistant Attorney General, Antitrust Division, reviewed the proposed Voluntary Agreement and consulted with the Chair of the Federal Trade Commission. On December 12, 2025, by letter to Assistant Secretary for Nuclear Energy Thedore J. Garrish, Gail Slater, Assistant Attorney General, Antitrust Division, issued a finding, pursuant to 50 U.S.C. 4558(f)(1)(B), that the purposes of the DPA's plans of action provision ''may not reasonably be achieved through a . . . plan of action having less anticompetitive effects or without any . . . plan of action.''

Dated: December 16, 2025.

David G.B. Lawrence,

Policy Director, Antitrust Division.

[FR Doc. 2025–23443 Filed 12–18–25; 8:45 am]

BILLING CODE 4410–11–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1614]

Importer of Controlled Substances Application: Aveva Drug Delivery Systems, Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Aveva Drug Delivery Systems, 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 January 20, 2026. Such persons may also file a written request for a hearing on the application on or before January 20, 2026.

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 April 16, 2025, Aveva Drug Delivery Systems, Inc., 3250 Commerce Parkway, Miramar, Florida 33025–3907, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Thebaine	9333	II

The company plans to import the listed controlled substance for analytical purposes only. No other activity for this drug code 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.

Thomas W. Prevoznik,

Deputy Assistant Administrator.

[FR Doc. 2025–23469 Filed 12–18–25; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1612]

Importer of Controlled Substances Application: Bright Green Corporation

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Bright Green Corporation 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 January 20, 2026. Such persons may also file a written request for a hearing on the application on or before January 20, 2026.

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