

personnel, solely for cybersecurity purposes. All contract personnel will sign appropriate nondisclosure agreements. All non-confidential written submissions will be available for public inspection at the Office of the Secretary and on EDIS.

The Commission vote for this determination took place on September 9, 2021.

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: September 9, 2021.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2021-19849 Filed 9-14-21; 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 transaction was 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

08/26/2021

20212939	G	Alight, Inc.; Alight Solutions LLC; Aon plc; Aon Hewitt Health Market Insurance Solutions Inc.
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Suzanne Morris,

Chief, Premerger and Division Statistics, Antitrust Division, Department of Justice.

[FR Doc. 2021-19943 Filed 9-14-21; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-893]

Importer of Controlled Substances Application: Cambridge Isotope Laboratories

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Cambridge Isotope Laboratories has applied to be registered as an importer of basic class(es) of controlled substance(s). Refer to **SUPPLEMENTAL INFORMATION** listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before October 15, 2021. Such persons may also file a written request for a hearing on the application on or before October 15, 2021.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701

Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on July 29, 2021, Cambridge Isotope Laboratories, 50 Frontage Road, Andover, Massachusetts 01810-5413, 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
Tetrahydrocannabinols	7370	I
Morphine	9300	II

The company plans to import the listed controlled substances for analytical research. 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.

Brian S. Besser,

Acting Assistant Administrator.

[FR Doc. 2021-19944 Filed 9-14-21; 8:45 am]

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DEPARTMENT OF JUSTICE

Notice of Lodging of Proposed Consent Decree Under the Clean Air Act

On September 9, 2021, the U.S. Department of Justice (DOJ) filed a Complaint and lodged a proposed Consent Decree with the United States District Court for the Eastern District of Michigan in *United States of America and Michigan Department of Environment, Great Lakes, and Energy v. Arbor Hills Energy LLC* No. 5:21-cv-12098.

The proposed Consent Decree resolves several Clean Air Act and State law claims against Arbor Hills Energy LLC (AHE), including for exceedances of permitted SO₂ emissions limits, at AHE's landfill gas-to-energy facility in Northville, Michigan. The AHE facility converts landfill gas (LFG), which is generated by decomposition of waste from an adjacent landfill, into electricity by burning it as fuel in four gas turbines.