

Collection Request. Calculated estimates for an SRS respondent to respond indicate 7 minutes per quarter. The total annual burden hour per respondent is 28 minutes. Total Annual Hour Burden: 7 minutes × 4 quarters = 28 minutes.

6. *An estimate of the total public burden (in hours) associated with the collection:* There are approximately 5,300 hours, annual burden, associated with this information collection.

11,357 respondents × 4 responses/year = 45,428 total annual responses.

45,428 × 7 minutes/60 minutes = 5,300 total annual hour burden.

(This burden estimate does not include the 6,933 NIBRS agencies; the NIBRS burden hours are captured in the NIBRS Information Collection Request.)

If additional information is required contact: Jerri Murray, Department Clearance Officer, United States Department of Justice, Justice Management Division, Policy and Planning Staff, Two Constitution Square, 145 N Street NE., 3E.405, Washington, DC 20530.

Dated: July 9, 2014.

**Jerri Murray,**

*Department Clearance Officer for PRA, U.S. Department of Justice.*

[FR Doc. 2014-16383 Filed 7-11-14; 8:45 am]

**BILLING CODE 4410-02-P**

## DEPARTMENT OF JUSTICE

### Notice of Lodging of Proposed Modification of Amended Consent Decree Under The Clean Air Act

On July 8, 2014, the Department of Justice lodged a proposed Third Amended Consent Decree with the United States District Court for the Eastern District of Wisconsin in the lawsuit entitled *United States and Michigan Department of Environmental Quality, Plaintiffs, and Clean Wisconsin, Sierra Club, and Citizens' Utility Board, Intervenor, v. Wisconsin Electric Power Company*, Civil Action No. 03-c-0371.

Generally, the proposed modifications to the Decree are designed: (1) To accommodate the voluntary decision of the Defendant, Wisconsin Electric Power Company ("WE Energies," "WE" or "Defendant"), to convert all four coal-fired boilers at the Valley Generating Station ("Valley Station"), located in Milwaukee, Wisconsin, from coal to natural gas; and (2) to simplify the process of terminating the Third Amended Decree after December 31, 2015. The coal-to-natural-gas conversion will provide significant emission reductions at the Valley Station, and the

termination-related changes will provide greater finality for the Defendant while also ensuring that the Decree's provisions remain enforceable in the future through federally enforceable state operating permits.

The publication of this notice opens a period for public comment on Proposed Third Amended Consent Decree. Comments should be addressed to the Acting Assistant Attorney General, Environment and Natural Resources Division, and should refer to *United States et al. v. Wisconsin Electric Power Company*, D.J. Ref. No. 90-5-2-1-07493. All comments must be submitted no later than thirty (30) days after the publication date of this notice. Comments may be submitted either by email or by mail:

| <i>To submit comments:</i> | <i>Send them to:</i>  |
|----------------------------|---|
| By email .....             | <i>pubcomment-ees.enrd@usdoj.gov.</i>   |
| By mail .....              | Assistant Attorney General,<br>U.S. DOJ—ENRD, P.O.<br>Box 7611, Washington,<br>DC 20044-7611. |

During the public comment period, the Joint Stipulation to Modify Section XXI of the Amended Consent Decree may be examined and downloaded at this Justice Department Web site: <http://www.usdoj.gov/enrd/ConsentDecrees.html>. We will provide a paper copy of the Proposed Third Amended Consent Decree upon written request and payment of reproduction costs. Please mail your request and payment to: Consent Decree Library, U.S. DOJ—ENRD, P.O. Box 7611, Washington, DC 20044-7611.

Please enclose a check or money order for \$21.75 (25 cents per page reproduction cost) payable to the United States Treasury.

**Thomas P. Carroll,**

*Assistant Section Chief, Environmental Enforcement Section, Environment and Natural Resources Division.*

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA-392]

### Importer of Controlled Substances Registration: Meridian Medical Technologies

**ACTION:** Notice of registration.

**SUMMARY:** Meridian Medical Technologies applied to be registered as

an importer of a certain basic class of narcotic controlled substance. The DEA grants Meridian Medical Technologies registration as an importer of this controlled substance.

**SUPPLEMENTARY INFORMATION:** By notice dated April 21, 2014, and published in the **Federal Register** on April 28, 2014, 79 FR 23374, Meridian Medical Technologies, 2555 Hermelin Drive, St. Louis, Missouri 63144, applied to be registered as an importer of a certain basic class of controlled substance. No comments or objections were submitted for this notice.

The Drug Enforcement Administration (DEA) has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of Meridian Medical Technologies to import the basic class of controlled substance is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated the company's maintenance of effective controls against diversion by inspecting and testing the company's physical security systems, verifying the company's compliance with state and local laws, and reviewing the company's background and history.

Therefore, pursuant to 21 U.S.C. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the above-named company is granted registration as an importer of morphine (9300), a basic class of narcotic controlled substance listed in schedule II.

The company manufactures a product containing morphine in the United States. The company exports this product to customers around the world. The company has been asked to ensure that its product, which is sold to European customers, meets the standards established by the European Pharmacopeia, administered by the Directorate for the Quality of Medicines (EDQM). In order to ensure that its product will meet European specifications, the company seeks to import morphine supplied by EDQM for use as reference standards.

This is the sole purpose for which the company will be authorized by the DEA to import morphine.

Dated: July 7, 2014.

**Joseph T. Rannazzisi,**

*Deputy Assistant Administrator.*

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