Billing Code 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances;

Notice of Registration;

Cedarburg Pharmaceuticals, Inc.

By Notice dated October 16, 2013, and published in the Federal Register on October 25, 2013, 78 FR 64017, Cedarburg Pharmaceuticals, Inc., 870 Badger Circle, Grafton, Wisconsin 53024, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
4-Anilino-N-phenethyl-4-piperidine (8333)	II
Remifentanil (9739)	II
Fentanyl (9801)	II

The company plans to manufacture the listed controlled substances in bulk for distribution to its customers.

No comments or objections have been received. The DEA has considered the factors in 21 U.S.C. 823(a), and determined that the registration of Cedarburg

Pharmaceuticals, Inc. to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. The DEA has investigated Cedarburg Pharmaceuticals, Inc. to ensure that the company's registration is consistent with the public interest. The investigation has included inspection and testing of the company's physical security systems, verification of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to 21 U.S.C. 823(a) and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Dated: February 19, 2014.

Joseph T. Rannazzisi,
Deputy Assistant Administrator,
Office of Diversion Control,
Drug Enforcement Administration.

[FR Doc. 2014-05496 Filed 03/12/2014 at 8:45 am; Publication Date: 03/13/2014]