Billing Code 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances Registration: Pharmacore, Inc.

[Docket No. DEA-392]

ACTION: Notice of registration.

SUMMARY: Pharmacore, Inc. applied to be registered as a manufacturer of certain basic classes of controlled substances. The Drug Enforcement Administration (DEA) grants Pharmacore, Inc. registration as a manufacturer of those controlled substances.

SUPPLEMENTARY INFORMATION:

By notice dated April 14, 2015, and published in the *Federal Register* on April 22, 2015, 80 FR 22554, Pharmacore, Inc., 4180 Mendenhall Oaks Parkway, High Point, North Carolina 27265 applied to be registered as a manufacturer of certain basic classes of controlled substances. No comments or objections were submitted for this notice.

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Pharmacore, Inc. to manufacture the basic classes of controlled substances 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. 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:

Controlled Substance	Schedule
Oxymorphone (9652)	II
Noroxymorphone (9668)	II

The company plans to manufacture the listed controlled substance as an active pharmaceutical ingredient (API) for clinical trials.

Dated: July 29, 2015

Joseph T. Rannazzisi,
Deputy Assistant Administrator.

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