Billing Code 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances;

Notice of Registration;

GE Healthcare

By Notice dated October 16, 2013, and published in the Federal Register on October 25, 2013, 78 FR 64018, GE Healthcare, 3350 North Ridge Avenue, Arlington Heights, Illinois 60004-1412, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Cocaine (9041), a basic class of controlled substance listed in schedule II.

The company plans to manufacture a radioactive product to diagnose Parkinson's disease 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 GE Healthcare to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. The DEA has investigated GE Healthcare 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, and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic class of controlled substance listed.

Dated: February 19, 2014.

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

[FR Doc. 2014-05484 Filed 03/12/2014 at 8:45 am; Publication Date: 03/13/2014]