records as required by 21 CFR 1304.04(f)(1) and (g), and he issued prescriptions to obtain controlled substances for office use in violation of 21 CFR 1306.04(b). These are all also violations of state law. Further from 1993 to 1997, Dr. Potter distributed anabolic steriods to a number of individuals for no legitimate medical purpose and outside the scope of professional practice in violation of 21 U.S.C. 841(a)(1).

As to factor three, it is undisputed that Dr. Potter was convicted of 432 felony offenses relating to his unlawful distribution of anabolic steriods.

Regarding factor five, there is no evidence in the investigation file of any other conduct which may threaten the public health and safety.

The Administrator concludes that Dr. Potter's continued registration would be inconsistent with the public interest based on his controlled substance record keeping violations, his unlawful distribution of anabolic steriods, and his conviction of 432 felony offenses. No evidence of explanation or mitigating circumstances was offered by Dr. Potter, or anyone purporting to represent him.

Accordingly, the Administrator of the Drug Enforcement Administration, pursuant to the authority vested in him by 21 U.S.C. 823 and 824, and 28 CFR 0.100(b), hereby orders that DEA Certificate of Registration BP2137847, previously issued to William C. Potter, D.V.M., be, and it hereby is, revoked. The Administrator further orders that any pending applications for renewal of such registration, be, and they hereby are, denied. This order is effective September 18, 2000.

Dated: August 3, 2000.

Donnie R. Marshall,

Administrator.

[FR Doc. 00–21114 Filed 8–17–00; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated April 6, 2000, and published in the **Federal Register** on April 25, 2000, (65 FR 24227), Roche Diagnostics Corporation, 9115 Hague Road, Indianapolis, Indiana 46250, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Lysergic acid diethylamide (7315)	I
Tetrahydrocannabinols (7370)	I
Phencyclidine (7471)	II
Benzoylecgonine (9180)	II
Methadone (9250)	II
Morphine (9300)	II

Roche Diagnostics Corporation plans to manufacture small quantities of the above listed controlled substances for incorporation in drug of abuse detection kits.

DEA has considered the factors in title 21, United States Code, section 823(a) and determined that the registration of Roche Diagnostics Corporation to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Roche Diagnostics Corporation on a regular basis to ensure that the company's continued registration is consistent with the public interest. These investigations have included inspection and testing of the company's physical security systems, audits of the company's records, 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 28 CFR 0.100 and 0.014, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic classes of controlled substances listed above is granted.

Dated: August 1, 2000.

John H. King,

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

[FR Doc. 00–21117 Filed 8–17–00; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated April 6, 2000, and published in the **Federal Register** on April 25, 2000 (65 FR 24227), Roche Diagnostics Corporation, 9115 Hague Road, Indianapolis, Indiana 46250, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Lysergic acid diethylamide (7315)	I
Tetrahydrocannabinols (7370)	I
Cocaine (9041)	II
Methadone (9250)	II
Morphine (9300)	II

The firm plans to import the listed controlled substances for the manufacture of diagnostic products.

No comments or objections have been received. DEA has considered the factors in title 21, United States Code, section 823(a) and determined that the registration of Roche Diagnostics Corporation, is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971, at this time. DEA has investigated Roche Diagnostics Corporation on a regular basis to ensure that the company's continued registration is consistent with the public interest. These investigations have included inspection and testing of the company's physical security systems, audits of the company's records, verification of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to section 1008(a) of the Controlled Substances Import and Export Act and in accordance with title 21, Code of Federal Regulations, section 1311.42, the above firm is granted registration as an importer of the basic classes of controlled substances listed above.

Dated: August 7, 2000.

John H. King,

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

[FR Doc. 00–21120 Filed 8–17–00; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Graham Travers Schuler, M.D.; Denial of Application

On November 19, 1999, the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration (DEA), issued an Order to Show Cause to Graham Travers Schuler, M.D., of Bloomington, Indiana. The Order to Show Cause notified him of an opportunity to show cause as to why DEA should not deny his application for a DEA Certificate of Registration pursuant to 21 U.S.C. 823(f) and 824(a)(3) and (a)(4), for reason that his state controlled substance