

Federal Register Representative (CCR), and must be filed no later than December 3, 2001.

Dated: September 25, 2001.

**Laura M. Nagel,**

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

[FR Doc. 01-24641 Filed 10-2-01; 8:45 am]

BILLING CODE 4410-09-M

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on April 16, 2001, Roche Diagnostics Corporation, Attn: Regulatory Compliance, 9115 Hague Road, Indianapolis, Indiana 46250, made application by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of Alphamethadol (9605), a basic class of controlled substance listed in Schedule I.

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

Any other such applicant and any person who is presently registered with DEA to manufacture such substance may file comments or objections to the issuance of the proposed registration.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistance Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than December 3, 2001.

Dated: September 24, 2001.

**Laura M. Nagel,**

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

[FR Doc. 01-24642 Filed 10-2-01; 8:45 am]

BILLING CODE 4410-09-M

## DEPARTMENT OF LABOR

### Occupational Safety and Health Administration

[Docket No. ICR-1218-0069(2001)]

#### Commercial Diving-Operations Standards (29 CFR part 1910, subpart T); Extension of the Office of Management and Budget's Approval of Information-Collection (Paperwork) Requirements

**AGENCY:** Occupational Safety and Health Administration (OSHA), Labor.

**ACTION:** Request for comments.

**SUMMARY:** OSHA solicits comments concerning its request to increase the total burden-hour estimate for, and to extend OMB approval of, the collection-of-information requirements specified by the Commercial Diving-Operations Standards (29 CFR part 1910, subpart T).<sup>1</sup> These standards specify paperwork requirements for equipment and procedures that expose employees to hazards associated with diving and diving-support operations, and that apply to general industry, construction, ship repairing, shipbuilding, shipbreaking, and longshoring.

**DATES:** Submit written comments on or before December 3, 2001.

**ADDRESSES:** Submit written comments to the Docket Office, Docket No. ICR-1218-0069(2001), OSHA, U.S. Department of Labor, Room N-2625, 200 Constitution Avenue, NW., Washington, DC 20210; telephone (202) 693-2350. Commenters may transmit written comments of 10 pages or less by facsimile to (202) 693-1648.

**FOR FURTHER INFORMATION CONTACT:** Theda Kenney, Directorate of Safety Standards Programs, OSHA, U.S. Department of Labor, Room N-3609, 200 Constitution Avenue, NW., Washington, DC 20210; telephone (202) 693-2222. A copy of the Agency's Information-Collection Request (ICR) supporting the need for the information collections specified by the Commercial-Diving Operations Standards is available for inspection and copying in the Docket Office, or by requesting a copy from Theda Kenney at (202) 693-2222 or Todd Owen at (202) 693-2444. For electronic copies of the ICR, contact OSHA on the Internet at

<sup>1</sup> Based on its assessment of the paperwork requirements contained in these standards, the Agency estimates that the total burden hours increased compared to its previous burden-hour estimate. Under this notice, OSHA is *not* proposing to revise these paperwork requirements in any substantive manner, only to increase the burden hours imposed by the existing paperwork requirements.

<http://www.osha.gov> and select "Information Collection Requests."

#### SUPPLEMENTARY INFORMATION:

##### I. Background

The Department of Labor, as part of its continuing effort to reduce paperwork and respondent (*i.e.*, employer) burden, conducts a preclearance consultation program to provide the public with an opportunity to comment on proposed and continuing information-collection requirements in accordance with the Paperwork Reduction Act of 1995 (PRA-95) (44 U.S.C. 3506(c)(2)(A)). This program ensures that information is in the desired format, reporting burden (time and costs) is minimal, collection instruments are understandable, and OSHA's estimate of the information-collection burden is correct.

The following provisions of the Commercial-Diving Operations Standards (the "Standards") contain paperwork requirements: §§ 1910.401(b); 1910.410(a)(3) and (a)(4); 1910.420(a) and (b); 1910.421(b), (f), and (h); 1910.422(e); 1910.423(b)(1)(ii) through (b)(2), (d), and (e); 1910.430(a), (b)(4), (c)(1)(i), (c)(3)(i), (f)(3)(ii), and (g)(2); and 1910.440(a)(2) and (b). These provisions ensure that employers: Notify OSHA if they deviate from the operational requirements of the Standards; train every diver in cardiopulmonary resuscitation and first aid, and mixed-gas divers (and those who control exposure of divers to mixed-gas breathing conditions) in diving-related physics and physiology; develop and make available to employees a safe-practices manual; maintain a list of emergency telephone or call numbers at the diving location; brief dive-team members on diving-related tasks, safety procedures, hazards, and revisions to operating procedures; display a code flag "A" if diving from a surface other than a vessel in navigable waters; develop and maintain a depth-time profile for each dive; and instruct divers on reporting diving-related illnesses and injuries, and the procedures specified for detecting, treating, and preventing these problems.

The Standards also mandate that employers: Record and maintain diving logs that contain required information; investigate, and provide a written evaluation of, any incident involving decompression sickness; mark diving umbilicals as required; inspect, test, and calibrate specified diving equipment; record modifications, repairs, tests, calibrations, and maintenance performed on any diving equipment; make a record of diving-related injuries and illnesses that result in a diver