required under paragraphs (i)(1), (i)(2), and (i)(3) of this section must be removed or otherwise obscured.

(j) A tie-tag attached to the container may be used for providing the information required by paragraphs (e)(1)(iii), (e)(2)(ii), and (e)(3), (h), or (i)(1), (i)(2), and (i)(3) of this section.

■ 4. Section 606.122 is amended by:
■ a. Revising the section heading;
■ b. Revising the introductory text;
■ c. Revising paragraphs (e), (f), (m)(2), (m)(3), and (m)(5); and
■ d. Revising the introductory text in paragraphs (k), (l), (m), and (n).

The revisions read as follows:

§ 606.122 Circular of information.

A circular of information must be available for distribution if the product is intended for transfusion. The circular of information must provide adequate directions for use, including the following information:

* * * * *

(e) A statement that the product was prepared from blood that was found negative when tested for communicable disease agents, as required under § 610.40 of this chapter (include each test that was performed).

(f) The statement: ''Warning: The risk of transmitting infectious agents is present. Careful donor selection and available laboratory tests do not eliminate the hazard.''

* * * * *

(k) For Red Blood Cells, the circular of information must contain:

* * * * *

(l) For Platelets, the circular of information must contain:

* * * * *

(m) For Plasma, the circular of information must contain:

(1) * * *

(2) Instructions to thaw the frozen product at a temperature appropriate for the product.

(3) When applicable, instructions to begin administration of the product within a specified time after thawing.

* * * * *

(5) A statement that this product has the same risk of transmitting infectious agents as Whole Blood; other plasma volume expanders without this risk are available for treating hypovolemia.

(n) For Cryoprecipitated AHF, the circular of information must contain:

* * * * *

■ 6. Section 606.170 is amended by revising paragraph (b) to read as follows:

§ 606.170 Adverse reaction file.

* * * * *

(b) When a complication of blood collection or transfusion is confirmed to be fatal, the Director, Office of Compliance and Biologics Quality, CBER, must be notified by telephone, facsimile, express mail, or electronically transmitted mail as soon as possible. A written report of the investigation must be submitted to the Director, Office of Compliance and Biologics Quality, CBER, by mail, facsimile, or electronically transmitted mail (for mailing addresses, see § 600.2 of this chapter), within 7 days after the fatality by the collecting facility in the event of a donor reaction, or by the facility that performed the compatibility tests in the event of a transfusion reaction.

PART 610—GENERAL BIOLOGICAL PRODUCTS STANDARDS

■ 7. The authority citation for 21 CFR part 610 continues to read as follows:

Authority: 21 U.S.C. 321, 331, 351, 352, 353, 355, 360, 360c, 360d, 360h, 360i, 371, 372, 374, 381; 42 U.S.C. 216, 262, 263, 263a, 264.

■ 8. Section 610.40 is amended by revising paragraphs (h)(2)(ii)(B) and (i) to read as follows:

§ 610.40 Test requirements.

* * * * *

(h) * * *

(2) * * *

(ii) * * *

(B) You must appropriately label such blood or blood components as required under § 606.121 of this chapter, and with the ''BIOHAZARD'' legend;

* * * * *

(i) *Syphilis testing.* In addition to the testing otherwise required under this section, you must test by a serological test for syphilis under §§ 640.5(a), 640.14, 640.23(a), 640.33(a), 640.53(a), and 640.65(b)(1) and (b)(2) of this chapter.

PART 640—ADDITIONAL STANDARDS FOR HUMAN BLOOD AND BLOOD PRODUCTS

■ 9. The authority citation for 21 CFR part 640 continues to read as follows:

Authority: 21 U.S.C. 321, 351, 352, 353, 355, 360, 371; 42 U.S.C. 216, 262, 263, 263a, 264.

§ 640.70 [Removed]

■ 10. Section 640.70 is removed.
■ 11. Section 640.74 is amended by revising paragraph (b)(4) to read as follows:

§ 640.74 Modification of Source Plasma.

* * * * *

(b) * * *

(4) The label affixed to each container of Source Plasma Liquid shall contain, in addition to the information required by § 606.121 of this chapter, but excluding § 606.121(e)(5)(ii) of this chapter, the name of the manufacturer of the final blood derivative product for whom it was prepared.

* * * * *

Dated: December 22, 2011.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2011–33554 Filed 12–30–11; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF LABOR

Occupational Safety and Health Administration

29 CFR Part 1915

RIN 1218–AB50

General Working Conditions in Shipyard Employment; Approval of Information Collection Requirements

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

ACTION: Final rule; notice of Office of Management and Budget (OMB) approval of collection of information requirements.

SUMMARY: OSHA is announcing that OMB approved the collection of information requirements contained in the General Working Conditions Standard under the Paperwork Reduction Act of 1995. The OMB approval number is 1218–0259.

DATES: The rule is effective January 3, 2012. The final rule, published May 2, 2011 (76 FR 24576), became effective and enforceable on August 1, 2011, except for the provisions in § 1915.89, which became effective and enforceable on October 31, 2011.

FOR FURTHER INFORMATION CONTACT: Theda Kenney, OSHA, Directorate of Standards and Guidance, U.S. Department of Labor, Room N–3609, 200 Constitution Avenue NW., Washington, DC 20210; telephone (202) 693–2222.

SUPPLEMENTARY INFORMATION: OSHA published a final rule for General Working Conditions in Shipyard Employment on May 2, 2011 (76 FR 24576), updating existing requirements to reflect advances in industry practices and technology, consolidating some general safety and health requirements into one subpart, and providing hazardous energy protection not addressed in the existing standard.

As required by the Paperwork Reduction Act of 1995, the **Federal**

Register notice for the General Working Conditions in Shipyard Employment final rule stated that compliance with the collection of information requirements was not required until OMB approved these requirements, and that the Department of Labor would publish a notice in the **Federal Register** announcing that OMB approved and assigned a control number to the requirements. See 76 FR 24695. Under 5 CFR 1320.5(b), an agency may not conduct or sponsor a collection of information unless: (1) The collection of information displays a currently valid OMB control number, and (2) the agency informs those members of the public who must respond to the collection of information that they are not required to respond to the collection of information unless it displays a currently valid OMB control number.

On May 2, 2011, OSHA submitted the General Working Conditions in Shipyard Employment (29 CFR part 1915, subpart F) Information Collection Request for the final rule to OMB for approval in accordance with the Paperwork Reduction Act of 1995 (44 U.S.C. 3501–3520). On October 31, 2011, OMB approved the collections of information contained in the final rule and assigned this collection OMB Control Number 1218–0259.

List of Subjects in 29 CFR Part 1915

Occupational safety and health, reporting, Recordkeeping requirements, Hazards in general working condition in shipyard employment.

Authority and Signature

David Michaels, Ph.D., MPH, Assistant Secretary of Labor for Occupational Safety and Health, directed the preparation of this notice. The authority for this notice is the Paperwork Reduction Act of 1995 (44 U.S.C. 3506 *et seq.*), and Secretary of Labor's Order No. 4–2010 (75 FR 55355).

Signed at Washington, DC, on December 22, 2011.

David Michaels,

Assistant Secretary of Labor for Occupational Safety and Health.

Amendments to Standard

For the reasons stated in the preamble to the final rule, the Occupational Safety and Health Administration amends 29 CFR part 1915 to read as follows:

PART 1915—[AMENDED]

Subpart F—[Amended]

■ Amend § 1915.8, by adding to the table the entries ''1915.83, 1915.87, 1915.88, and 1915.89'' in the proper numerical sequence as follows:

§ 1915.8 OMB control numbers under the Paperwork Reduction Act.

* * * * *

29 CFR citation	OMB control No.
1915.83	1218–0259
1915.87	1218–0259
1915.88	1218–0259
1915.89	1218–0259

* * * * *

[FR Doc. 2011–33260 Filed 12–30–11; 8:45 am]

BILLING CODE 4510–26–P

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 52

[EPA–R02–OAR–2011–0607; FRL–9611–2]

Approval and Promulgation of Air Quality Implementation Plans; State of New Jersey; Regional Haze State Implementation Plan

AGENCY: Environmental Protection Agency (EPA).

ACTION: Final rule.

SUMMARY: The Environmental Protection Agency (EPA) is approving the revision to the New Jersey State Implementation Plan, submitted by the State of New Jersey. The revision addresses Clean Air Act requirements and EPA's rules for states to prevent and remedy future and existing anthropogenic impairment of visibility in mandatory Class I areas through a regional haze program. EPA's approval includes but is not limited to New Jersey's plans to implement Reasonable Progress Goals, Best Available Retrofit Technologies on eligible sources, as well as New Jersey's Subchapter 9, Sulfur in Fuels rule and source-specific SIP revisions.

DATES: *Effective Date:* This rule is effective on February 2, 2012.

ADDRESSES: EPA has established a docket for this action under Docket ID No. EPA–R02–OAR–2011–0607. All documents in the docket are listed on the *www.regulations.gov* Web site. Although listed in the index, some information is not publicly available, *e.g.,* CBI or other information whose disclosure is restricted by statute. Certain other material, such as copyrighted material, is not placed on the Internet and will be publicly available only in hard copy form. Publicly available docket materials are available either electronically through *www.regulations.gov* or in hard copy at the Environmental Protection Agency, Region II Office, Air Programs Branch, 290 Broadway, 25th Floor, New York, New York 10007–1866. This Docket Facility is open from 8:30 a.m. to 4:30 p.m., Monday through Friday, excluding legal holidays. The Docket telephone number is (212) 637–4249.

FOR FURTHER INFORMATION CONTACT: Robert F. Kelly, State Implementation Planning Section, Air Programs Branch, EPA Region 2, 290 Broadway, New York, New York 10007–1866. The telephone number is (212) 637–4249. Mr. Kelly can also be reached via electronic mail at *kelly.bob@epa.gov.*

SUPPLEMENTARY INFORMATION:

Table of Contents

I. What action is EPA taking?

EPA is approving a revision to New Jersey's State Implementation Plan (SIP) submitted on July 28, 2009, that addressed progress toward reducing regional haze for the first implementation period ending in 2018. The initial submittal was supplemented by a December 9, 2010 submittal transmitting New Jersey's adopted regulation Subchapter 9 Sulfur in Fuel, lowering the sulfur content in fuel oil, a March 2, 2011 submittal which included Best Available Retrofit Technologies (BART) determinations and controls, and a December 7, 2011 submittal including Air Pollution Control Operating Permits for sources that require BART reductions, as listed in the regulatory section of this action.

EPA determined that New Jersey's Regional Haze Plan contains the emission reductions needed to achieve New Jersey's share of emission reductions that were determined to be reasonable through the regional planning process. Furthermore, New Jersey's Regional Haze Plan ensures that emissions from the State will not interfere with the Reasonable Progress Goals (RPGs) for neighboring States' Class I areas. Thus, EPA is approving into the SIP the Regional Haze Plan submitted by New Jersey on July 28, 2009 and supplemented on December 9, 2010, March 2, 2011, and December 7, 2011 as satisfying the requirements of the Clean Air Act. EPA is taking this action pursuant to Section 110 of the Act.