as of the date each member organization implements riskless principal routing, the member organization have in place systems and controls that allow them to easily match and tie the riskless principal execution on the Exchange to the underlying orders and that they be able to provide this information to the Exchange upon request.

2. Statutory Basis

The basis under the Act for this proposed rule change is the requirement under section 6(b)(5)[6] that an Exchange have rules that are designed to promote just and equitable principles of trade, to remove impediments to and perfect the mechanism of a free and open market and a national market system, and, in general, to protect investors and the public interest.

B. Self-Regulatory Organization's Statement on Burden on Competition

The Exchange does not believe that the proposed rule change will impose any burden on competition that is not necessary or appropriate in furtherance of the purposes of the Act.

C. Self-Regulatory Organization's Statement on Comments on the Proposed Rule Change Received From Members, Participants, or Others

The Exchange has neither solicited nor received written comments on the proposed rule change.

III. Date of Effectiveness of the Proposed Rule Change and Timing for Commission Action

Because the proposed rule change: (i) Does not significantly affect the protection of investors or the public interest; (ii) does not impose any significant burden on competition; and (iii) does not become operative for 30 days after the date of the filing, or such shorter time as the Commission may designate if consistent with the protection of investors and the public interest, the proposed rule change has become effective pursuant to section 19(b)(3)(A) of the Act[7] and Rule 19b–4(f)(6) thereunder.[8]

At any time within 60 days of the filing of the proposed rule change, the Commission may summarily abrogate the rule change if it appears to the Commission that such action is necessary or appropriate in the public interest, for the protection of investors, or otherwise in furtherance of the purposes of the Act.

IV. Solicitation of Comments

Interested persons are invited to submit written data, views and arguments concerning the foregoing, including whether the proposed rule change is consistent with the Act. Comments may be submitted by any of the following methods:

Electronic Comments

- Use the Commission's Internet comment form (*http://www.sec.gov/rules/sro.shtml*); or
- Send an e-mail to *rule-comments@sec.gov*. Please include File Number SR–NYSE–2007–114 on the subject line.

Paper Comments

- Send paper comments in triplicate to Nancy M. Morris, Secretary, Securities and Exchange Commission, 100 F Street, NE., Washington, DC 20549–1090.

All submissions should refer to File Number SR–NYSE–2007–114. This file number should be included on the subject line if e-mail is used. To help the Commission process and review your comments more efficiently, please use only one method. The Commission will post all comments on the Commission's Internet Web site (*http://www.sec.gov/rules/sro.shtml*). Copies of the submission, all subsequent amendments, all written statements with respect to the proposed rule change that are filed with the Commission, and all written communications relating to the proposed rule change between the Commission and any person, other than those that may be withheld from the public in accordance with the provisions of 5 U.S.C. 552, will be available for inspection and copying in the Commission's Public Reference Room, on official business days between the hours of 10 a.m. and 3 p.m. Copies of the filing also will be available for inspection and copying at the principal office of the Exchange. All comments received will be posted without change; the Commission does not edit personal identifying information from submissions. You should submit only information that you wish to make available publicly. All submissions should refer to File Number SR–NYSE–2007–114 and should be submitted on or before January 10, 2008.

For the Commission, by the Division of Trading and Markets, pursuant to delegated authority.[9]

Florence E. Harmon,

Deputy Secretary.

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[6] 15 U.S.C. 78f(b)(5).

[7] 15 U.S.C. 78s(b)(3)(A).

[8] 17 CFR 240.19b–4(f)(6). Pursuant to Rule 19b–4(f)(6)(iii) under the Act, the Exchange is required to give the Commission written notice of its intent to file the proposed rule change, along with a brief description and text of the proposed rule change, at least five business days prior to the date of filing of the proposed rule change, or such shorter time as designated by the Commission. The Exchange has satisfied the five-day pre-filing requirement.

[9] 17 CFR 200.30–3(a)(12).

SMALL BUSINESS ADMINISTRATION

Small Business Size Standards: Waiver of the Nonmanufacturer Rule

AGENCY: U.S. Small Business Administration.

ACTION: Notice of Waiver of the Nonmanufacturer Rule for Electromedical and Electrotherapeutic Apparatus Manufacturing.

SUMMARY: The U. S. Small Business Administration (SBA) is granting a waiver of the Nonmanufacturer Rule for Electromedical and Electrotherapeutic Apparatus Manufacturing, Diagnostic equipment, MRI (magnetic resonance imaging) manufacturing; Magnetic resonance imaging (MRI) medical diagnostic equipment manufacturing; Medical ultrasound equipment manufacturing; MRI (magnetic resonance imaging) medical diagnostic equipment manufacturing; Patient monitoring equipment (e.g., intensive care coronary care unit) manufacturing; PET (positron emission equipment tomography) scanners manufacturing; and Positron emission tomography (PET) scanners manufacturing. The basis for a waiver is that no small business manufacturers are supplying this class of product to the Federal government. The effect of a waiver would be to allow otherwise qualified regular dealers to supply the products of any domestic manufacturer on a Federal contract set aside for small businesses; service-disabled veteran-owned small business or SBA's 8(a) Business Development Program.

DATE: This waiver is effective January 4, 2008.

FOR FURTHER INFORMATION CONTACT: Edith Butler, Program Analyst, by telephone at (202) 619–0422; by FAX at (202) 481–1788; or by e-mail at *edith.butler@sba.gov.*

SUPPLEMENTARY INFORMATION: Section 8(a)(17) of the Small Business Act, (Act) 15 U.S.C. 637(a)(17), requires that recipients of Federal contracts set aside for small businesses, service-disabled veteran-owned small businesses, or SBA's 8(a) Business Development Program provide the product of a small

business manufacturer or processor, if the recipient is other than the actual manufacturer or processor of the product. This requirement is commonly referred to as the Nonmanufacturer Rule. The SBA regulations imposing this requirement are found at 13 CFR 121.406(b). Section 8(a)(17)(b)(iv) of the Act authorizes SBA to waive the Nonmanufacturer Rule for any ''class of products'' for which there are no small business manufacturers or processors available to participate in the Federal market.

As implemented in SBA's regulations at 13 CFR 121.1202(c), in order to be considered available to participate in the Federal market for a class of products, a small business manufacturer must have submitted a proposal for a contract solicitation or received a contract from the Federal government within the last 24 months. The SBA defines ''class of products'' based on six digit coding systems. The first coding system is the Office of Management and Budget North American Industry Classification System (NAICS). The second is the Product and Service Code required as a data entry field by the Federal Procurement Data System.

The SBA received a request on October 23, 2007 to waive the Nonmanufacturer Rule for Electromedical and Electrotherapeutic Apparatus Manufacturing, Diagnostic equipment, MRI (magnetic resonance imaging) manufacturing; Magnetic resonance imaging (MRI) medical diagnostic equipment manufacturing; Medical ultrasound equipment manufacturing; MRI (magnetic resonance imaging) medical diagnostic equipment manufacturing; Patient monitoring equipment (e.g., intensive care coronary care unit) manufacturing; PET (positron emission equipment tomography) scanners manufacturing; and Positron emission tomography (PET) scanners manufacturing. In response, on November 15, 2007, SBA published in the **Federal Register** a notice of intent to waive the Nonmanufacturer Rule for Electromedical and Electrotherapeutic Apparatus Manufacturing, Diagnostic equipment, MRI (magnetic resonance imaging) manufacturing; Magnetic resonance imaging (MRI) medical diagnostic equipment manufacturing; Medical ultrasound equipment manufacturing; MRI (magnetic resonance imaging) medical diagnostic equipment manufacturing; Patient monitoring equipment (e.g, intensive care coronary care unit) manufacturing; PET (positron emission equipment tomography) scanners manufacturing; and Positron emission tomography (PET) scanners manufacturing. SBA explained in the notice that it was soliciting comments and sources of small business manufacturers of this class of products.

In response to this notice, a comment was received from an interested party, however, no small business manufacturing sources were discovered. SBA has determined that there are no small business manufacturers of this class of products, and is therefore granting the waiver of the Nonmanufacturer Rule for Electromedical and Electrotherapeutic Apparatus Manufacturing, Diagnostic equipment, MRI (magnetic resonance imaging) manufacturing; Magnetic resonance imaging (MRI) medical diagnostic equipment manufacturing; Medical ultrasound equipment manufacturing; MRI (magnetic resonance imaging) medical diagnostic equipment manufacturing; Patient monitoring equipment (e.g., intensive care coronary care unit) manufacturing; PET (positron emission equipment tomography) scanners manufacturing; and Positron emission tomography (PET) scanners manufacturing, NAICS 334510.

Authority: 15 U.S.C. 637(a)(17).

Dated: December 13, 2007.

Arthur E. Collins, Jr.,

Director, Office of Government Contracting.

[FR Doc. E7–24716 Filed 12–19–07; 8:45 am]

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DEPARTMENT OF STATE

[Public Notice 6038]

Culturally Significant Objects Imported for Exhibition Determinations: ''Color Chart: Reinventing Color''

SUMMARY: Notice is hereby given of the following determinations: Pursuant to the authority vested in me by the Act of October 19, 1965 (79 Stat. 985; 22 U.S.C. 2459), Executive Order 12047 of March 27, 1978, the Foreign Affairs Reform and Restructuring Act of 1998 (112 Stat. 2681, *et seq.*; 22 U.S.C. 6501 note, *et seq.*), Delegation of Authority No. 234 of October 1, 1999, Delegation of Authority No. 236 of October 19, 1999, as amended, and Delegation of Authority No. 257 of April 15, 2003 [68 FR 19875], I hereby determine that the objects to be included in the exhibition ''Color Chart: Reinventing Color'', imported from abroad for temporary exhibition within the United States, are of cultural significance. The objects are imported pursuant to loan agreements with the foreign owners or custodians. I also determine that the exhibition or display of the exhibit objects at the Museum of Modern Art, New York, NY, from on or about March 2, 2008, until on or about May 12, 2008, and at possible additional exhibitions or venues yet to be determined, is in the national interest. Public Notice of these Determinations is ordered to be published in the **Federal Register**.

FOR FURTHER INFORMATION CONTACT: For further information, including a list of the exhibit objects, contact Richard Lahne, Attorney-Adviser, Office of the Legal Adviser, U.S. Department of State (telephone: 202/453–8058). The address is U.S. Department of State, SA–44, 301 4th Street, SW., Room 700, Washington, DC 20547–0001.

Dated: December 13, 2007.

C. Miller Crouch,

Principal Deputy Assistant Secretary for Educational and Cultural Affairs, Department of State.

[FR Doc. E7–24731 Filed 12–19–07; 8:45 am]

BILLING CODE 4710–05–P

DEPARTMENT OF STATE

[Public Notice 6037]

Culturally Significant Objects Imported for Exhibition Determinations: ''The Color of Life''

SUMMARY: Notice is hereby given of the following determinations: Pursuant to the authority vested in me by the Act of October 19, 1965 (79 Stat. 985; 22 U.S.C. 2459), Executive Order 12047 of March 27, 1978, the Foreign Affairs Reform and Restructuring Act of 1998 (112 Stat. 2681, *et seq.*; 22 U.S.C. 6501 note, *et seq.*), Delegation of Authority No. 234 of October 1, 1999, Delegation of Authority No. 236 of October 19, 1999, as amended, and Delegation of Authority No. 257 of April 15, 2003 [68 FR 19875], I hereby determine that the objects to be included in the exhibition ''The Color of Life'', imported from abroad for temporary exhibition within the United States, are of cultural significance. The objects are imported pursuant to loan agreements with the foreign owners or custodians. I also determine that the exhibition or display of the exhibit objects at The J. Paul Getty Museum at the Getty Villa, Malibu, California, from on or about March 6, 2008, until on or about June 23, 2008, and at possible additional exhibitions or venues yet to be determined, is in the national interest. Public Notice of these Determinations is ordered to be published in the **Federal Register**.

FOR FURTHER INFORMATION CONTACT: For further information, including a list of