Industry Classification System (NAICS) code 334517 product number 6525. The public is invited to comment or provide source information to SBA on the proposed waivers of the Nonmanufacturer Rule for this class of NAICS code within 15 days after date of publication in the Federal Business Opportunities.

Dated: November 6, 2007.

Arthur E. Collins, Jr.,

Director for Government Contracting.

[FR Doc. E7–22353 Filed 11–14–07; 8:45 am]

BILLING CODE 8025–01–P

SMALL BUSINESS ADMINISTRATION

Small Business Size Standards: Waiver of the Nonmanufacturer Rule

AGENCY: U.S. Small Business Administration.

ACTION: Notice of intent to waive the Nonmanufacturer Rule for Electromedical and Electrotherapeutic Apparatus Manufacturing.

SUMMARY: The U.S. Small Business Administration (SBA) is considering granting a request for 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. According to the request, no small business manufacturers supply these classes of products to the Federal government. If granted, the waiver would 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 businesses or SBA's 8(a) Business Development Program.

DATES: Comments and source information must be submitted November 30, 2007.

ADDRESSES: You may submit comments and source information to Edith G. Butler, Program Analyst, U.S. Small Business Administration, Office of Government Contracting, 409 3rd Street, SW., Suite 8800, Washington, DC 20416.

FOR FURTHER INFORMATI0N CONTACT: Edith G. 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 a six digit coding system. The coding system is the Office of Management and Budget North American Industry Classification System (NAICS).

The SBA is currently processing a request 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, North American Industry Classification System (NAICS) code 334510 product number 6525.

The public is invited to comment or provide source information to SBA on the proposed waivers of the Nonmanufacturer Rule for this class of NAICS code within 15 days after date of publication in the **Federal Register**.

Dated: November 6, 2007.

Arthur E. Collins, Jr.,

Director for Government Contracting.

[FR Doc. E7–22357 Filed 11–14–07; 8:45 am]

BILLING CODE 8025–01–P

DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

Notice of Passenger Facility Charge (PFC) Approvals and Disapprovals

AGENCY: Federal Aviation Administration (FAA), DOT.

ACTION: Monthly Notice of PFC Approvals and Disapprovals. In October 2007, there were nine applications approved. This notice also includes information on two applications, approved in September 2007, inadvertently left off the September 2007 notice. Additionally, 14 approved amendments to previously approved applications are listed.

SUMMARY: The FAA publishes a monthly notice, as appropriate, of PFC approvals and disapprovals under the provisions of the Aviation Safety and Capacity Expansion Act of 1990 (Title IX of the Omnibus Budget Reconciliation Act of 1990) (Pub. L. 101–508) and Part 158 of the Federal Aviation Regulations (14 CFR part 158). This notice is published pursuant to paragraph (d) of § 158.29.

PFC Applications Approved

Public Agency: City of Phoenix, Arizona.

Application Number: 07–08–C–00–PHX.

Application Type: Impose and use a PFC.

PFC Level: $4.50.

Total PFC Revenue Approved in This Decision: $202,200,000.

Earliest Charge Effective Date: August 1, 2008.

Estimated Charge Expiration Date: August 1, 2010.

Class of Air Carriers Not Required To Collect PFC's

(1) Non-scheduled, on-demand air carriers filing FAA Form 1800–31; (2) commuters or small certificated air carriers filing Department of Transportation Form 298—C T1 or E1 with less than 7,500 annual enplanements at Phoenix Sky Harbor International Airport (PHX); (3) large certificated air carriers filing Research and Special Programs Administration (RSPA) Form T–100 with less than 7,500 annual enplanements at PHX; and (4) foreign air carriers filing RSPA Form T–100(f) with less than 7,500 annual enplanements at PHX.