

the end of FY 2010, FDA estimates that 445 establishments will have been billed for establishment fees, before all decisions on requests for waivers or reductions are made. FDA estimates that a total of 15 establishment fee waivers or reductions will be made for FY 2010. In addition, FDA estimates that another 15 full establishment fees will be exempted this year based on the orphan drug exemption in FDAAA (see section 736(k) of the act). Subtracting 30 establishments (15 waivers plus the estimated 15 establishments under the orphan exemption) from 445 leaves a net of 415 fee-paying establishments. FDA will use 415 for its FY 2011 estimate of establishments paying fees, after taking waivers and reductions into account. The fee per establishment is determined by dividing the adjusted total fee revenue to be derived from establishments (\$206,356,667) by the estimated 415 establishments, for an establishment fee rate for FY 2011 of \$497,200 (rounded to the nearest \$100).

B. Product Fees

At the beginning of FY 2010, the product fee was based on an estimate that 2,380 products would be subject to and would pay product fees. By the end of FY 2010, FDA estimates that 2,460 products will have been billed for product fees, before all decisions on requests for waivers, reductions, or exemptions are made. FDA assumes that there will be about 50 waivers and reductions granted. In addition, FDA estimates that another 25 product fees will be exempted this year based on the orphan drug exemption in FDAAA (see section 736(k) of the act). FDA estimates that 2,385 products will qualify for product fees in FY 2010, after allowing for waivers and reductions, including the orphan drug products eligible under the FDAAA exemption, and will use this number for its FY 2011 estimate. Accordingly, the FY 2011 product fee rate is determined by dividing the adjusted total fee revenue to be derived from product fees (\$206,356,667) by the estimated 2,385 products for a FY 2011 product fee of \$86,520 (rounded to the nearest \$10).

V. Fee Schedule for FY 2011

The fee rates for FY 2011 are set out in table 6 of this document:

TABLE 6—FEE SCHEDULE FOR FY 2011

Fee Category	Fee Rates for FY 2011
Applications	

TABLE 6—FEE SCHEDULE FOR FY 2011—Continued

Fee Category	Fee Rates for FY 2011
Requiring clinical data	\$1,542,000
Not requiring clinical data	\$771,000
Supplements requiring clinical data	\$771,000
Establishments	\$497,200
Products	\$86,520

VI. Fee Payment Options and Procedures

A. Application Fees

The appropriate application fee established in the new fee schedule must be paid for any application or supplement subject to fees under PDUFA that is received after September 30, 2010. Payment must be made in U.S. currency by check, bank draft, or U.S. postal money order payable to the order of the Food and Drug Administration. Please include the user fee identification (ID) number on your check, bank draft, or postal money order. Your payment can be mailed to: Food and Drug Administration, P.O. Box 70963, Charlotte, NC 28272-0963.

If checks are to be sent by a courier that requests a street address, the courier can deliver the checks to: Wells Fargo, Attn: Food and Drug Administration Lockbox 70963, 1525 West WT Harris Blvd., rm. D1113-022, Charlotte, NC 28262. (Note: This Wells Fargo address is for courier delivery only.)

Please make sure that the FDA post office box number (P.O. Box 70963) is written on the check, bank draft, or postal money order.

Wire transfer payment may also be used. Please reference your unique user fee ID number when completing your transfer. The originating financial institution usually charges a wire transfer fee between \$15.00 and \$35.00. Please ask your financial institution about the fee and include it with your payment to ensure that your fee is fully paid. The account information is as follows: New York Federal Reserve Bank, US Dept of Treasury, TREAS NYC, 33 Liberty St., New York, NY 10045, Acct. No.: 75060099, Routing No.: 021030004, SWIFT: FRNYUS33, Beneficiary: FDA, 1350 Piccard Dr., Rockville, MD.

Application fees can also be paid online with an electronic check (ACH). FDA has partnered with the U.S. Department of the Treasury to utilize

Pay.gov, a Web-based payment application, for online electronic payment. The Pay.gov feature is available on the FDA Web site after the user fee ID number is generated.

The tax identification number of the Food and Drug Administration is 53-0196965.

B. Establishment and Product Fees

FDA will issue invoices for establishment and product fees for FY 2011 under the new fee schedule in August 2010. Payment will be due on October 1, 2010. FDA will issue invoices in November 2011 for any products and establishments subject to fees for FY 2011 that qualify for fees after the August 2010 billing.

Dated: July 29, 2010.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2010-19116 Filed 8-3-10; 8:45 am]

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DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

[Docket No. FR-5376-N-78]

Notice of Submission of Proposed Information Collection to OMB Research Plan for an Evaluation of the Section 202 Demonstration Planning Grant (DPG) Program

AGENCY: Office of the Chief Information Officer, HUD.

ACTION: Notice.

SUMMARY: The proposed information collection requirement described below has been submitted to the Office of Management and Budget (OMB) for review, as required by the Paperwork Reduction Act. The Department is soliciting public comments on the subject proposal.

This research is intended to help HUD better understand sponsor perspectives on the effectiveness of the DPG program in assisting Section 202 properties reach initial closing within 18 months of fund reservation. The study will also provide information on sponsor perspectives of the marketing of the DPG program by HUD filed office staff, the DPG application process and the overall administration of the grant program. The respondents are both recipients and non-recipients on the 202 DPG grant.

DATES: *Comments Due Date: September 3, 2010.*

ADDRESSES: Interested persons are invited to submit comments regarding this proposal. Comments should refer to the proposal by name and/or OMB

approval Number (2528–New) and should be sent to: HUD Desk Officer, Office of Management and Budget, New Executive Office Building, Washington, DC 20503; fax: 202–395–5806.

FOR FURTHER INFORMATION CONTACT:

Leroy McKinney Jr., Reports Management Officer, QDAM, Department of Housing and Urban Development, 451 Seventh Street, SW., Washington, DC 20410; e-mail Leroy McKinney Jr. at Leroy.McKinneyJr@hud.gov or telephone (202) 402–5564. This is not a toll-free number. Copies of available documents submitted to OMB may be obtained from Mr. McKinney.

SUPPLEMENTARY INFORMATION: This notice informs the public that the Department of Housing and Urban Development has submitted to OMB a request for approval of the Information collection described below. This notice

is soliciting comments from members of the public and affecting agencies concerning the proposed collection of information to: (1) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; (2) Evaluate the accuracy of the agency’s estimate of the burden of the proposed collection of information; (3) Enhance the quality, utility, and clarity of the information to be collected; and (4) Minimize the burden of the collection of information on those who are to respond; including through the use of appropriate automated collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

This notice also lists the following information:

Title of Proposal: Research Plan for an Evaluation of the Section 202 DPG Program.

OMB Approval Number: 2528–New.

Form Numbers: None.

Description of the Need For the Information and Its Proposed Use: This research is intended to help HUD better understand sponsor perspectives on the effectiveness of the DPG program in assisting Section 202 properties reach initial closing within 18 months of fund reservation. The study will also provide information on sponsor perspectives of the marketing of the DPG program by HUD filed office staff, the DPG application process and the overall administration of the grant program. The respondents are both recipients and non-recipients on the 202 DPG grant.

Frequency of Submission: On-occasion.

	Number of respondents	Annual responses	×	Hours per response	=	Burden hours
Reporting Burden	100	1		1.01		101

Total Estimated Burden Hours: 101.
Status: New Collection.

Authority: Section 3507 of the Paperwork Reduction Act of 1995, 44 U.S.C. 35, as amended.

Dated: July 28, 2010.

Leroy McKinney, Jr.,
Departmental Reports Management Officer,
Office of the Chief Information Officer.

[FR Doc. 2010–19081 Filed 8–3–10; 8:45 am]

BILLING CODE 4210–67–P

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

[Docket No. FR–5430–N–01]

Proposed Fair Market Rents for the Housing Choice Voucher Program and Moderate Rehabilitation Single Room Occupancy Program Fiscal Year 2011

AGENCY: Office of the Assistant Secretary for Policy Development and Research, HUD.

ACTION: Notice of Proposed Fiscal Year (FY) 2011 Fair Market Rents (FMRs).

SUMMARY: Section 8(c)(1) of the United States Housing Act of 1937 (USHA) requires the Secretary to publish FMRs periodically, but not less than annually, adjusted to be effective on October 1 of each year. Today’s notice proposes FMRs for FY 2011 to be used to determine payment standard amounts for the Housing Choice Voucher (HCV) program, to determine initial renewal

rents for some expiring project-based Section 8 contracts, and to determine initial rents for Housing Assistance Payment (HAP) contracts in the Moderate Rehabilitation Single Room Occupancy program. Other programs may require use of FMRs for other purposes.

The proposed FY 2011 FMR areas are based on current Office of Management and Budget (OMB) metropolitan area definitions and include HUD modifications that were first used in the determination of FY 2006 FMR areas. Changes to the OMB metropolitan area definitions through December 2009 are incorporated. The principal city for three metropolitan areas changed, which resulted in a code change. In Alaska, there was a name change for a non-metropolitan borough and two boroughs in Alaska were divided to make four new boroughs.¹ Proposed FY 2011 FMRs are based on 2000 Census data updated using more current survey data. For FY 2011, FY 2010 FMRs are updated using 2008 American Community Survey (ACS) data, and more recent Consumer Price Index (CPI)

¹ The three metropolitan areas are: North Port-Bradenton-Sarasota, FL MSA, Crestview-Fort Walton Beach-Destin, FL MSA, and Steubenville-Weirton, OH-WV MSA. In Alaska, Prince of Wales-Ketchikan Census Area, AK is changed to Prince of Wales-Hyder Census Area, AK; the Alaskan borough of Skagway-Hoonah-Angoon is divided into Skagway and Hoonah-Angoon boroughs; and the Alaskan borough of Wrangell-Petersburg is divided into Wrangell and Petersburg boroughs.

rent and utility indexes. HUD continues to use ACS data in different ways based upon the number of two-bedroom standard-quality and recent-mover sample cases that are available in the FMR area or its Core-Based Statistical Area (CBSA).

This notice also proposes Small Area FMRs for the Dallas, TX HUD Metro FMR Area in accordance with a **Federal Register** Notice published May 18, 2010, (75 FR27808) announcing a Small Area FMR Demonstration project.

DATES: *Comment Due Date:* 30 days after publication.

ADDRESSES: Interested persons are invited to submit comments regarding HUD’s estimates of the FMRs, as published in this notice, to the Office of General Counsel, Rules Docket Clerk, Department of Housing and Urban Development, 451 Seventh Street, SW., Room 10276, Washington, DC 20410–0001. Communications should refer to the above docket number and title and should contain the information specified in the “Request for Comments” section.

Submission of Hard Copy Comments. To ensure that the information is fully considered by all of the reviewers, each commenter who is submitting hard copy comments, by mail or hand delivery, is requested to submit two copies of its comments to the address above, one addressed to the attention of the Rules Docket Clerk and the other addressed to the attention of Economic and Market