(collectively, the Agencies) for both affiliated and nonaffiliated institutions. The form name is the Interagency Bank Merger Act Application. The Agencies need the information collected to insure that the proposed transactions are permissible under law and regulation and are consistent with safe and sound banking practices. The Agencies are required, for example, to consider financial and managerial resources, future prospects, convenience and needs of the community, community reinvestment, and competition.

Some of the Agencies will collect limited supplemental information in certain cases. For example, the OCC and OTS will collect information regarding CRA commitments, the Federal Reserve will collect information on debt servicing from certain institutions, and all Agencies will require additional information on the competitive impact of proposed mergers.

Current actions: On January 5, 1998, the Board granted initial approval of the proposal. A joint notice of the proposed action was published in the *Federal Register* on January 21, 1998 (63 FR 3182), and the comment period expired on March 23, 1998. The Agencies received five public comments from the Texas Department of Banking, the Independent Bankers Association of America, the National Community Investment Coalition, the Center for Community Change, and the Conference of State Bank Supervisors, as well as comments from staff at each agency. Most of the commenters suggested modifications to the forms and instructions. As a result of the comments, the application was further revised to include an ''Other'' category under the ''Filed Pursuant To'' section and information on Tier 3 capital (if any), the addresses of directors and senior executive officers, how the proposal will meet the convenience and needs of the community (including needs of the community under the applicable criteria of the Community Reinvestment Act, and debt servicing (if applicable). In addition, certain branch information requested in the initial proposal was eliminated. The additional changes proposed in response to the comments would not affect most applicants; on average for all applicants, the estimated burden would be unchanged. The other agencies submitted the same revised information collection to OMB for approval.

Board of Governors of the Federal Reserve System, December 23, 1998.

Jennifer J. Johnson,

Secretary of the Board.

[FR Doc. 98–34549 Filed 12–29–98; 8:45AM]

Billing Code 6210–01–F

DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000–0031]

Proposed Collection; Comment Request Entitled Contractor Use of Government Supply Sources

AGENCIES: Department of Defense (DOD), General Services Administration (GSA), and National Aeronautics and Space Administration (NASA).

ACTION: Notice of request for public comments regarding an extension to an existing OMB clearance.

SUMMARY: Under the provisions of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35), the Federal Acquisition Regulation (FAR) Secretariat will be submitting to the Office of Management and Budget (OMB) a request to review and approve an extension of a currently approved information collection requirement concerning Contractor Use of Government Supply Sources. The clearance currently expires on April 30, 1999.

DATES: Comments may be submitted on or before March 1, 1999.

ADDRESSES: Comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, should be submitted to: FAR Desk Officer, OMB, Room 10102, NEOB, Washington, DC 20503, and a copy to the General Services Administration, FAR Secretariat (MVRS), 1800 F Street, NW, Room 4035, Washington, DC 20405. Please cite OMB Control No. 9000–0031, Contractor Use of Government Supply Sources, in all correspondence.

FOR FURTHER INFORMATION CONTACT: Linda Klein, Federal Acquisition Policy Division, GSA (202) 501–3775.

SUPPLEMENTARY INFORMATION:

A. Purpose

When it is in the best interest of the Government and when supplies and services are required by a Government contract, contracting officers may authorize contractors to use Government supply sources in performing certain contracts. Contractors placing orders under Federal Supply Schedules or Personal Property Rehabilitation Price Schedules must follow the terms of the applicable schedule. To place orders, firms will submit the initial FEDSTRIP or MILSTRIP requisitions or the Optional Form 347, a copy of the authorization to order, and a statement regarding authorization to the firm holding the schedule contract.

The information informs the schedule contractor that the ordering contractor is authorized to use this Government supply source and fills the ordering contractor's order under the terms of the Government contract.

B. Annual Reporting Burden

Public reporting burden for this collection of information is estimated to average *15* minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information.

The annual reporting burden is estimated as follows: Respondents, *300*; responses per respondent, *7*; total annual responses, *2,100*; preparation hours per response, *.25*; and total response burden hours, *525*.

Obtaining copies of proposals: Requester may obtain a copy of the justification from the General Services Administration, FAR Secretariat (MVRS), Room 4035, 1800 F Street, NW, Washington, DC 20405, telephone (202) 208–7312. Please cite OMB Control No. 9000–0031, Contractor Use of Government Supply Sources, in all correspondence.

Dated: December 23, 1998.

Victoria E. Moss,

Acting Director, Federal Acquisition Policy Division.

[FR Doc. 98–34402 Filed 12–29–98; 8:45 am]

BILLING CODE 6820–34–U

DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000–0032]

Proposed Collection; Comment Request Entitled Contractor Use of Interagency Motor Pool Vehicles

AGENCIES: Department of Defense (DoD), General Services Administration (GSA),

and National Aeronautics and Space Administration (NASA).

ACTION: Notice of request for public comments regarding an extension to an existing OMB clearance.

SUMMARY: Under the provisions of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35), the Federal Acquisition Regulation (FAR) Secretariat will be submitting to the Office of Management and Budget (OMB) a request to review and approve an extension of a currently approved information collection requirement concerning Contractor Use of Interagency Motor Pool Vehicles. The clearance currently expires on April 30, 1999.

DATES: Comments may be submitted on or before March 1, 1999.

ADDRESSES: Comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, should be submitted to: FAR Desk Officer, OMB, Room 10102, NEOB, Washington, DC 20503, and a copy to the General Services Administration, FAR Secretariat (MVRS), 1800 F Street, NW, Room 4035, Washington, DC 20405. Please cite OMB Control No. 9000–0032, Contractor Use of Interagency Motor Pool Vehicles, in all correspondence.

FOR FURTHER INFORMATION CONTACT: Linda Klein, Federal Acquisition Policy Division, GSA (202) 501–3775.

SUPPLEMENTARY INFORMATION:

A. Purpose

If it is in the best interest of the Government, the contracting officer may authorize cost-reimbursement contractors to obtain, for official purposes only, interagency motor pool vehicles and related services. Contractors' requests for vehicles must obtain two copies of the agency authorization, the number of vehicles and related services required and period of use, a list of employees who are authorized to request the vehicles, a listing of equipment authorized to be serviced, and billing instructions and address.

A written statement that the contractor will assume, without the right of reimbursement from the Government, the cost or expense of any use of the motor pool vehicles and services not related to the performance of the contract is necessary before the contracting officer may authorize cost-reimbursement contractors to obtain interagency motor pool vehicles and related services.

The information is used by the Government to determine that it is in the Government's best interest to authorize a cost-reimbursement contractor to obtain, for official purposes only, interageny motor pool vehicles and related services, and to provide those vehicles.

B. Annual Reporting Burden

Public reporting burden for this collection of information is estimated to average *30* minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information.

The annual reporting burden is estimated as follows: Respondents, *70*; responses per respondent, *2*; total annual responses, *140*; preparation hours per response, *.5*; and total response burden hours, *70*.

Obtaining copies of proposals: Requester may obtain a copy of the justification from the General Services Administration, FAR Secretariat (MVRS), Room 4035, 1800 F Street, NW, Washington, DC 20405, telephone (202) 208–7312. Please cite OMB Control No. 9000–0032, Contractor Use of Interagency Motor Pool Vehicles, in all correspondence.

Dated: December 23, 1998.

Victoria E. Moss,

Acting Director, Federal Acquisition Policy Division.

[FR Doc. 98–34403 Filed 12–29–98; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket Nos. 98M–0136, 98M–0217, 98M–0138, 98M–0327, 98M–0328, 98M–0219, 98M–0137, 98M–0404, 98M–0200, 98M–0140, 98M–0231, 98M–0187, 98M–0139, 98M–0201, 98M–0403, 98M–0162, 97M–0084, 98M–0329, 98M–0450, 98M–0451, 98M–0251, 98M–0507, 98M–0618, 98M–0604, 98M–0619, 96M–0678, 98M–0679, 98M–0715, 98M–0711, and 98M–0725]

Medical Devices; List of Premarket Approval Actions

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is publishing a list of premarket approval application (PMA) approvals. This list is intended to inform the public of the existence and the availability of summaries of safety and effectiveness of approved PMA's through the Internet and the agency's Dockets Management Branch.

ADDRESSES: Summaries of safety and effectiveness are available on the World Wide Web (WWW) at http://www.fda.gov/cdrh/pma page.html. Copies of summaries of safety and effectiveness are also available by submitting a written request to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Please cite the appropriate docket number as listed in Table 1 in the **SUPPLEMENTARY INFORMATION** section of this document, when submitting a written request.

FOR FURTHER INFORMATION CONTACT: Kathy M. Poneleit, Center for Devices and Radiological Health (HFZ–402), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301–594–2186.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of January 30, 1998 (63 FR 4571), FDA published a final rule to revise §§ 814.44(d) and 814.45(d) (21 CFR 814.44(d) and 814.45(d)) to discontinue publication of individual PMA approvals and denials in the **Federal Register**. Revised §§ 814.44(d) and 814.45(d) state that FDA will notify the public of PMA approvals and denials by posting them on FDA's home page on the Internet (http://www.fda.gov), by placing the summaries of safety and effectiveness on the Internet and in FDA's Dockets Management Branch, and by publishing in the **Federal Register** after each quarter a list of the PMA approvals and denials announced in that quarter.

FDA believes that this procedure expedites public notification of these actions because announcements can be placed on the Internet more quickly than they can be published in the **Federal Register**, and FDA believes that the Internet is accessible to more people than the **Federal Register**.

In accordance with section 515(d)(3) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 360e(d)(3)), notification of an order approving, denying, or withdrawing approval of a PMA will continue to include a notice of opportunity to request review of the order under section 515(g) of the act. The 30-day period for requesting reconsideration of an FDA action under § 10.33(b) (21 CFR 10.33(b)) for notices announcing approval of a PMA begins on the day the notice is placed on the Internet. Section 10.33(b) provides that FDA may, for good cause, extend this 30-day period. Reconsideration of a denial or withdrawal of approval of a