complaint issuance, I am willing to support the allegations relating to these two markets.

The proposed relief appears to be necessary and appropriate to address the complaint allegations in the refining, pipeline, and terminal markets. In my view, the Commission's staff and the merging parties have worked diligently and creatively to craft relief to remedy the competitive concerns in these markets. However, given the extraordinary complexity of the divestitures and other relief negotiated, I welcome public comments addressing whether the order would fulfill its remedial purpose without causing unintended adverse effects on competition or consumers. In particular, I would be interested in public comment on whether the merging parties should be required to divest the Exxon refinery in Benicia, California, and the Exxon retail gasoline stations in California to a single buyer. From a purely economic basis, there seems to be little logic in forcing the divestiture of the refinery and the retail stations to a single buyer.

[FR Doc. 00–570 Filed 1–10–00; 8:45 am]

BILLING CODE 6750–01–P

GENERAL SERVICES ADMINISTRATION

Notice of Availability

The General Services Administration (GSA) has prepared a Record of Decision as the final document in the Environmental Impact Statement process for the renovation of the Tecate Port of Entry, Tecate, California. This project is designed to relocate the commercial operations, improve the working conditions for the U.S. Customs Service and U.S. Immigration and Naturalization Service, and improve the water systems on the port. For a copy of the Record of Decision contact: General Services Administration, 450 Golden Gate, Portfolio Division, San Francisco, California 94102, Attn: Rosanne Nieto, Phone: (415) 522–3490.

Arlin M. Timberlake,

Director, Portfolio Division, Public Buildings Service, General Services Administration.

[FR Doc. 00–1003 Filed 1–14–00; 8:45 am]

BILLING CODE 6820–BR–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control And Prevention

[60Day–00–18]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of Section 3506 (c) (2) (A) of the Paperwork Reduction Act of 1995, the Center for Disease Control and Prevention is providing opportunity for public comment on proposed data collection projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call the CDC Reports Clearance Officer on (404) 639–7090.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques for other forms of information technology. Send comments to Seleda Perryman, CDC Assistant Reports Clearance Officer, 1600 Clifton Road, MS–D24, Atlanta, GA 30333. Written comments should be received within 60 days of this notice.

Proposed Projects

1. PHS Supplements to the Application for Federal Assistance—SF–424 (0920–0428)—Extension. The Centers for Disease Control and Prevention (CDC) is requesting a three-year extension for continued use of the Supplements to the Request for Federal Assistance Application (SF–424). The Checklist, Program Narrative, and the Public Health System Impact Statement (third party notification) (PHSIS) are a part of the standard application for State and local governments and for private non-profit and for-profit organizations when applying for financial assistance from PHS grant programs. The Checklist assists applicants to ensure that they have included all required information necessary to process the application. The Checklist data helps to reduce the time required to process and review grant applications, expediting the issuance of grant awards. The PHSIS Third Party Notification Form is used to inform State and local health agencies of community-based proposals submitted by non-governmental applicants for Federal funding.

The total annual cost to the respondents is $1,184,452.

Respondents	No. of respondents	No. of responses/respondent	Avg. burden/response (in hrs.)	Total burden (in hrs.)
State and local health departments; non-profit and for-profit organizations	7,755	1	4.215	32,687
Total				32,687

Dated: January 11, 2000.

Nancy Cheal,

Acting Associate Director for Policy, Planning and Evaluation, Centers for Disease Control and Prevention (CDC).

[FR Doc. 00–1026 Filed 1–14–00; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control And Prevention

[60 Day–00–19]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of Section 3506 (c)(2)(A) of the Paperwork reduction Act of 1995, the Center for Disease Control and Prevention is providing opportunity for public comment on proposed data collection projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call the CDC Reports Clearance Officer on (404) 639–7090.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance

of the functions of the agency, including whether the information shall have practical utility; (b) The accuracy of the agency's estimate of the burden of the proposed collection of information; (c) Ways to enhance the quality, utility, and clarity of the information to be collected; and (d) Ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques for other forms of information technology. Send comments to Seleda Perryman, CDC Assistant Reports Clearance Officer, 1600 Clifton Road, MS–D24, Atlanta, GA 30333. Written comments should be received within 60 days of this notice.

Proposed Projects

1. Preventive Health and Health Services Block Grant, Annual Application and Reports (0920–0106)—Renewal—The National Center for Chronic Disease Prevention and Health Promotion—In 1994, the Office of Management and Budget approved the collection of information provided in the grant applications and annual reports for the Preventive Health and Health Services Block Grant (0920–0106). This approval expires on November 30, 2000. CDC is requesting an extension of OMB clearance for this legislatively mandated information collection until November 30, 2001. The extension is limited to one year to allow for the development and adherence to *Healthy People 2010* to be released the Spring of 2000. The Preventive Health and Health Services Block Grant is mandated according to section 1904 to adhere to the Healthy People framework, therefore, the current application and report format will be restructured to coincide with 2010 and resubmitted for OMB clearance at that time.

This information collected through the applications from the official State health agencies is required from section 1905 of the Public Health Service Act. There is no change in the proposed information collection from previous years. The information collected from the annual reports is required by section 1906, specifically the requirement for uniform data sets matching the uses of funds. The total cost to all respondents is $137,250, estimated at $25/burden hour.

Respondents	No. of respondents	No. of responses/respondent	Average burden per response	Total burden
Application	61	1	30	1830
Report	61	1	60	3660
Total				5490

Dated: January 11, 2000.

Nancy Cheal,

Acting Associate Director for Policy, Planning, and Evaluation, Centers for Disease Control and Prevention (CDC).

[FR Doc. 00–1027 Filed 1–14–00; 8:45 am]

BILLING CODE 4163–18–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 00N–0002]

Agency Information Collection Activities; Proposed Collection; Comment Request; Application for Exemption From Federal Preemption of State and Local Medical Device Requirements

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on FDA's requirements for State and local government applications for exemption from preemption for medical device requirements.

DATES: Submit written comments on the collection of information by March 20, 2000.

ADDRESSES: Submit written comments on the collection of information to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Peggy Schlosburg, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.