utilized for authorizing the marketing of FCS's, except where FDA determines that the submission and premarket review of an FAP under section 409(b) of the act is necessary to provide adequate assurance of safety. Section 409(h)(1) of the act requires that a notification include information on the identity and the intended use of the FCS and the basis for the notifier's determination that the FCS is safe under the intended conditions of use. Because section 409(h)(1) of the act references the general safety standard for food additives, the data in a PMN should be comparable to the data in an FAP. FDA is announcing the availability for comment of two draft guidance documents that are part of the agency's implementation of the PMN program, which will largely replace the FAP process for those food additives that are FCS's. The information to be collected is information on the manufacture and intended use of the FCS, studies relating to the safety of the FCS, and other information necessary to demonstrate that the FCS is safe under the intended conditions of use.

FDA is also making available for comment FDA Form No. 3480 entitled ''Notification for New Use of a Food Contact Substance'' for a notification for a new use of a FCS. FDA believes that this form will facilitate both preparation and review of notifications since the form will serve to organize information necessary to support the safety of the use of the FCS. The burden of filling out the appropriate form has been included in the burden estimate for the notification.

Description of Respondents: Manufacturers of food-contact substances.

FDA estimates the burden of this collection of information as follows:

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN [1]

Form	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
FDA 3480 [2]	200	1	200	25	5,000
FDA 3480 [3]	125	2	250	120	30,000
FDA 3480 [4]	45	2	90	150	13,500
FDA 3480 [5]	16	1	16	150	2,400
Total					50,900

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.
[2] Duplicate notifications for uses of FCS's.
[3] Notifications for uses that would currently be the subject of exemptions under 21 CFR 170.39 or very simple FAP's.
[4] Notifications for uses that would currently be the subject of moderately complex FAP's.
[5] Notifications for uses that would currently be the subject of more complex FAP's.

The above estimate is based on the types of submissions that FDA currently receives for FCS's in the TOR and the FAP processes and the following assumptions and information:

1. FDA estimates that the likely increase in PMN's over the number of FAP's and TOR requests will be approximately four times the highest recent influx of these submissions (50 and 54, respectively). This factor is based on an analysis of the number of companies producing various types of FCS's and the types of FCS's for which FAP's and TOR's are most commonly submitted to FDA.

2. FDA also has included 200 expected duplicate submissions in the second lowest tier. FDA expects that the burden for preparing these notifications will primarily consist of the notifier filling out FDA Form No. 3480, verifying that a previous notification is effective, and preparing necessary documentation.

3. Based on the amount of data typically submitted in FAP's and TOR requests, FDA identified three other tiers of PMN's that represent escalating levels of burden required to collect information.

4. FDA estimated the median number of hours necessary for collecting information for each type of notification within each of the three tiers, and the cost of developing necessary data based on input from industry sources.

V. Comments

Interested persons may, on or before February 14, 2000, submit to the Dockets Management Branch (address above) written comments regarding the two draft guidance documents. Two copies of any comments are to be submitted, except that individuals may submit one copy. Comments are to be identified with the docket numbers found in brackets in the heading of this document. Submit written comments concerning this collection of information to the Dockets Management Branch by January 11, 2000. The draft guidance documents and received comments may be seen in the office above between 9 a.m. and 4 p.m., Monday through Friday. Received comments will be considered when determining whether to amend the guidance.

Dated: November 1, 1999.

Margaret M. Dotzel,

Acting Associate Commissioner for Policy.

[FR Doc. 99–29493 Filed 11–10–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Advisory Council; Notice of Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Public Law 92–463), announcement is made of the following National Advisory body scheduled to meet during the month of December 1999.

Name: Advisory Commission on Childhood Vaccines (ACCV).

Date and Time: December 1, 1999; 9:00 a.m.–5:00 p.m.

Place: Parklawn Building, Conference Rooms G & H, 5600 Fishers Lane, Rockville, Maryland 20857.

The meeting is open to the public.

The full Commission will meet on Wednesday, December 1, 1999, from 9:00 a.m. to 5:00 p.m. Agenda items will include, but not be limited to: A discussion of the Government Accounting Office Report; a discussion of the six-month severity criteria to allow compensation for injuries, such as Intussusception; updates from the Department of Justice and the National Vaccine Program Office; and routine program reports.

Public comment will be permitted before lunch and at the end of the Commission meeting on December 1, 1999. Oral presentations will be limited to 5 minutes per public speaker. Persons interested in

providing an oral presentation should submit a written request, along with a copy of their presentation to: Ms. Shelia Tibbs, Principal Staff Liaison, Division of Vaccine Injury Compensation, Bureau of Health Professions, Health Resources and Services Administration, Room 8A–46, 5600 Fishers Lane, Rockville, MD 20857, Telephone (301) 443–6593. Requests should contain the name, address, telephone number, and any business or professional affiliation of the person desiring to make an oral presentation. Groups having similar interests are requested to combine their comments and present them through a single representative. The allocation of time may be adjusted to accommodate the level of expressed interest. The Division of Vaccine Injury Compensation will notify each presenter by mail or telephone of their assigned presentation time.

Persons who do not file an advance request for a presentation, but desire to make an oral statement, may sign-up in Conference Rooms G and H on December 1, 1999. These persons will be allocated time as time permits.

Anyone requiring information regarding the Commission should contact Ms. Tibbs at (301) 443–6593.

Agenda items are subject to change as priorities dictate.

Dated: November 4, 1999.

Jane M. Harrison,

Director, Division of Policy Review and Coordination.

[FR Doc. 99–29492 Filed 11–10–99; 8:45 am]

BILLING CODE 4160–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration (SAMHSA)

Notice of Meetings

Pursuant to Public Law 92–463, notice is hereby given of the following meeting of the SAMHSA Special Emphasis Panel I in November and December 1999.

A summary of the meetings and a roster of the members may be obtained from: Ms. Coral Sweeney, Review Specialist, SAMHSA, Office of Policy and Program Coordination, Division of Extramural Activities, Policy, and Review, 5600 Fishers Lane, Room 17–89, Rockville, Maryland 20857. Telephone: 301–443–2998.

Substantive program information may be obtained from the individual named as Contact for the meeting listed below.

The meetings will include the review, discussion and evaluation of individual grant applications. These discussions could reveal personal information concerning individuals associated with the applications. Accordingly, these meetings are concerned with matters exempt from mandatory disclosure in Title 5 U.S.C. 552b© (6) and 5 U.S.C. App.2, § 10(d).

Committee Name: SAMHSA Special Emphasis Panel I (SEP I).

Meeting Dates: November 15–19, 1999.

Place: Bethesda Marriott 5151 Pooks Hill Road, Bethesda, MD 20814.

Closed: November 15–19, 1999, 8:30 a.m.–5:00 p.m./adjournment.

Panel: Community Treatment Program, PA 99–050.

Contact: Michael Koscinski, Room 17–89, Parklawn Building, Telephone: 301–443–6094 and FAX: 301–443–3437.

Committee Name: SAMHSA Special Emphasis Panel I (SEP I).

Meeting Dates: December 13–15, 1999.

Place: Bethesda Marriott 5151 Pooks Hill Road Bethesda, MD 20814.

Closed: December 13–15, 1999 8:30 a.m.–5:00 p.m./adjournment.

Panel: Substance Abuse and Mental Health Services Administration Conference Grant PA 98–090(a).

Contact: Boris Aponte, Room 17–89, Parklawn Building, Telephone: 301–443–9912 and FAX: 301–443–3437.

Dated: November 1, 1999.

Coral Sweeney,

Review Specialist, Substance Abuse and Mental Health Services Administration.

[FR Doc. 99–29491 Filed 11–10–99; 8:45 am]

BILLING CODE 4162–20–P

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

[Docket No. FR–4432–N–45]

Federal Property Suitable as Facilities to Assist the Homeless

AGENCY: Office of the Assistant Secretary for Community Planning and Development, HUD.

ACTION: Notice.

SUMMARY: This Notice identifies unutilized, underutilized, excess, and surplus Federal property reviewed by HUD for suitability for possible use to assist the homeless.

FOR FURTHER INFORMATION CONTACT: Clifford Taffet, room 7266, Department of Housing and Urban Development, 451 Seventh Street SW., Washington, DC 20410; telephone (202) 708–1234; TTY number for the hearing- and speech-impaired (202) 708–2565 (these telephone numbers are not toll-free), or call the toll-free Title V information line at 1–800–927–7588.

SUPPLEMENTARY INFORMATION: In accordance with 24 CFR part 581 and section 501 of the Stewart B. McKinney Homeless Assistance Act (42 U.S.C. 11411), as amended, HUD is publishing this Notice to identify Federal buildings and other real property that HUD has reviewed for suitability for use to assist the homeless. The properties were reviewed using information provided to HUD by Federal landholding agencies regarding unutilized and underutilized buildings and real property controlled by such agencies or by GSA regarding its inventory of excess or surplus Federal property. This Notice is also published in order to comply with the December 12, 1988 Court Order in *National Coalition for the Homeless* v. *Veterans Administration*, No. 88–2503–OG (D.D.C.).

Properties reviewed are listed in this Notice according to the following categories: Suitable/available, suitable/unavailable, suitable/to be excess, and unsuitable. The properties listed in the three suitable categories have been reviewed by the landholding agencies, and each agency has transmitted to HUD: (1) Its intention to make the property available for use to assist the homeless, (2) Its intention to declare the property excess to the agency's needs, or (3) A statement of the reasons that the property cannot be declared excess or made available for use as facilities to assist the homeless.

Properties listed as suitable/available will be available exclusively for homeless use for a period of 60 days from the date of this Notice. Homeless assistance providers interested in any such property should send a written expression of interest to HHS, addressed to Brian Rooney, Division of Property Management, Program Support Center, HHS, room 5B–41, 5600 Fishers Lane, Rockville, MD 20857; (301) 443–2265. (This is not a toll-free number.) HHS will mail to the interested provider an application packet, which will include instructions for completing the application. In order to maximize the opportunity to utilize a suitable property, providers should submit their written expressions of interest as soon as possible. For complete details concerning the processing of applications, the reader is encouraged to refer to the interim rule governing this program, 24 CFR part 581.

For properties listed as suitable/to be excess, the property may, if subsequently accepted as excess by GSA, be made available for use by the homeless in accordance with applicable law, subject to screening for other Federal use. At the appropriate time, HUD will publish the property in a Notice showing it as either suitable/available or suitable/unavailable.

For properties listed as suitable/unavailable, the landholding agency has decided that the property cannot be declared excess or made available for