products subject to this order are as follows.

A. Registrations Listed in Table 1 of Unit II

The registrants may continue to sell and distribute existing stocks of products listed in Table 1 of Unit II. until December 28, 2012, which is 1 year after the publication of the Cancellation Order in the Federal Register. Thereafter, the registrants are prohibited from selling or distributing products listed in Table 1, except for export in accordance with FIFRA section 17, or proper disposal. Persons other than the registrants may sell, distribute, or use existing stocks of products listed in Table 1 of Unit II. until existing stocks are exhausted, provided that such sale, distribution, or use is consistent with the terms of the previously approved labeling on, or that accompanied, the cancelled products.

B. Registrations Listed in Table 2 of Unit II Except CA970032 and FL970008

The registrants may sell and distribute existing stocks of these products until January 15, 2012, 1 year after the date on which the maintenance fee was due. Thereafter, the registrants are prohibited from selling or distributing the pesticides identified in Table 2 of Unit II, except for export consistent with FIFRA section 17 or for proper disposal. Persons other than registrants are allowed to sell, distribute, or use existing stocks until such stocks are exhausted, provided that such sale, distribution, or use is consistent with the terms of the previously approved labeling on, or that accompanied, the cancelled products.

C. Registration Numbers CA970032 and FL970008

Mitigation measures included as part of the Reregistration Eligibility Decision (RED) for methyl bromide are being implemented in two phases. The first phase of mitigation is required to be placed on all product labels in 2011. To ensure that all methyl bromide products in the marketplace have the same protections at the same time, the following are the existing stocks provisions for the cancellation of registration numbers CA970032 and FL970008. The registrants may sell and distribute existing stocks of these products until December 31, 2011. Thereafter, the registrants are prohibited from selling and distributing the pesticides identified in Table 2 of Unit II., except for export consistent with FIFRA section 17 or for proper disposal. Persons other than registrants are allowed to sell and distribute existing stocks through April 30, 2012. After December 28, 2011, remaining stocks may be used until such stocks are exhausted, provided that such use is consistent with the terms of the previously approved labeling on, or that accompanied, the cancelled products.

List of Subjects

Environmental protection, Pesticides and pests.

Dated: November 17, 2011.

Richard P. Keigwin, Jr.
Director, Pesticide Re-evaluation Division, Office of Pesticide Programs.

GOVERNMENT ACCOUNTABILITY OFFICE

Medicare Payment Advisory Commission Nomination Letters

AGENCY: Government Accountability Office (GAO).

ACTION: Notice on letters of nomination.

SUMMARY: The Balanced Budget Act of 1997 established the Medicare Payment Advisory Commission (MedPAC) and gave the Comptroller General responsibility for appointing its members. For appointments to MedPAC that will be effective May 1, 2012, I am announcing the following: Letters of nomination should be submitted between January 1 and March 8, 2011, to ensure adequate opportunity for review and consideration of nominees prior to the appointment of new members.


Gene L. Dodaro,
Comptroller of the United States.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier CMS–10249 and CMS–10409]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Centers for Medicare & Medicaid Services (CMS), Department of Health and Human Services, is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the Agency’s function; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

1. Type of Information Collection Request: Revision of a currently approved collection; Title of Information Collection: Administrative Requirements for Section 6071 of the Deficit Reduction Act; Use: Under section 6071 of the Deficit Reduction Act of 2005 (Pub. L. 109–171) subsection (c), the Secretary may require States to meet requirements and provide additional information, provisions, and assurances. Through the Operational Protocol, States provide the requirements, information, provisions and assurances which, following CMS approval, States may enroll individuals in the State’s demonstration program or begin to claim for service dollars. The Act also requires the Money Follows the Person Rebalancing Demonstration (MFP) program be evaluated to determine program effectiveness. One aspect of the evaluation is determining participant quality of life and how the program affects quality of life. Medicaid enrollees who participate in the MFP program are expected to have need for long-term care services for the rest of their lives and are a particularly vulnerable population if the community setting cannot adequately meet their...