assesses the effectiveness of CMS' quality reporting activities.

• Effectively communicates program policies related to the Prescription Drug and Medicare Advantage (MA) programs to heath plans and drug plan contractors, employer group sponsors, beneficiary advocates and other stakeholders in the health care field.

• Develops new policies (e.g. health plan access, benefits, special needs plans) and programs to reflect changes in program objectives, the health care delivery system, beneficiary health care needs, and new plan types to support an appropriate range of choices for beneficiaries.

• Collaborates with our partners, such as industry, other government entities and advocacy groups, to understand their perspectives on Prescription Drug and Medicare Managed Care policies and procedures and to drive best practices in the health care industry.

• Develops and implements a comprehensive strategic plan, objectives and measures for overseeing an effective compliance and oversight program for all Part C (Medicare Advantage) and Part D (Medicare Prescription Drug) contractors in close collaboration with CMHPO, the Medicare Drug Benefit and C & D Data Group, the Medicare Drug and Health Plan Contract Administration Group and other Center for Drug and Health Plan Choice components.

• Develops and implements a comprehensive and effective audit program for all Part C (Medicare Advantage) and Part D (Medicare Prescription Drug) contractors.

Dated: June 21, 2008.

James W. Weber,

Acting Director, Office of Operations Management, Centers for Medicare & Medicaid Services.

[FR Doc. E8–14896 Filed 6–30–08; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2004–N–0188] (formerly Docket No. 2004N–0081)

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Request for Designation as Country Not Subject to the Restrictions Applicable to Human Food and Cosmetics Manufactured From, Processed With, or Otherwise Containing, Material From Cattle

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Request for Designation as Country not Subject to the Restrictions Applicable to Human Food and Cosmetics Manufactured from, Processed With, or Otherwise Containing, Material from Cattle'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of the Chief Information Officer (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of April 17, 2008 (73 FR 20785), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0623. The approval expires on June 30, 2011. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain*.

Dated: June 24, 2008.

Jeffrey Shuren,

Associate Commissioner for Policy and Planning.

[FR Doc. E8–14882 Filed 6–30–08; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–N–0354]

Agency Information Collection Activities; Proposed Collection; Comment Request; Mental Models Study of Farmers' Understanding and Implementation of Good Agricultural Practices

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 and to allow 60 days for public comment in response to the notice. This notice solicits comments on the proposed study entitled ''Mental Models Study of Farmers' Understanding and Implementation of Good Agricultural Practices.''

DATES: Submit written or electronic comments on the collection of information by September 2, 2008.

ADDRESSES: Submit electronic comments on the collection of information to *http://www.regulations.gov*. Submit written comments on the collection of information to the Division of Dockets Management (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: Jonna Capezzuto, Office of the Chief Information Officer (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

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