Therefore, based on the review and observations described in section III of this final notice, we have determined that NCQA's requirements for HMOs and local PPOs continue to meet or exceed our requirements. We renew the MA deeming authority of the NCQA for HMOS and PPOs for a term of 4 years. The new term of approval began October 19, 2010, and ends October 18, 2014.

IV. Results of the Review Process

Using the information listed in section III of this final notice, we determined that NCQA's current accreditation program for HMO and PPO MA plans continues to be at least as stringent as the MA requirements contained in the 6 categories specified in section 1852(e)(4)(C) of the Act and our methods of evaluation for those areas.

V. Collection of Information Requirements

This document does not impose information collection and recordkeeping requirements. Consequently, it need not be reviewed by the Office of Management and Budget under the authority of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35).

VI. Regulatory Impact Statement

In accordance with the provisions of Executive Order 12866, this regulation was not reviewed by the Office of Management and Budget.

(Catalog of Federal Domestic Assistance Program No. 93.773, Medicare—Hospital Insurance; and Program No. 93.774, Medicare—Supplementary Medical Insurance Program)

Dated: March 9, 2011.

Donald M. Berwick,

Administrator, Centers for Medicare & Medicaid Services.

[FR Doc. 2011–6222 Filed 3–24–11; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Voluntary Establishment of Paternity—NPRM.

OMB No.: 0970–0175.

Description: Section 466(a)(5)(C) of the Social Security Act requires States to pass laws ensuring a simple civil process for voluntarily acknowledging paternity under which the State must provide that the mother and putative father must be given notice, orally and in writing, of the benefits and legal responsibilities and consequences of acknowledging paternity. The information is to be used by hospitals, birth record agencies, and other entities participating in the voluntary paternity establishment program that collect information from the parents of children that are born out of wedlock.

Respondents:

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Disclosure	1,167,097	1	0.17	198,406.49

Estimated Total Annual Burden Hours: 198,406.49.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, *Attn:* ACF Reports Clearance Officer. All requests should be identified by the title of the information collection. *E-mail address: infocollection@acf.hhs.gov.*

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following:

Office of Management and Budget, Paperwork Reduction Project, *Fax:* 202–395–7285, *E-mail: OIRA_SUBMISSION@OMB.EOP.GOV,* Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2011–7077 Filed 3–24–11; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0620]

The National Antimicrobial Resistance Monitoring System Strategic Plan 2011–2015; Request for Comments; Extension of Comment Period

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; request for comments; extension of comment period.

SUMMARY: The Food and Drug Administration (FDA) is extending the comment period for the notice that appeared in the **Federal Register** of January 24, 2011 (76 FR 4120). In the notice, FDA requested comments on a document for the National Antimicrobial Resistance Monitoring System (NARMS) entitled "NARMS Strategic Plan 2011–2015." The Agency is taking this action in response to requests for an extension to allow interested persons additional time to submit comments. Based on requests received, additional information is being placed in the docket related to the development of the Strategic Plan. This information can also be viewed at the Web sites listed in section III of this document.

DATES: Submit either electronic or written comments by May 24, 2011.

ADDRESSES: Submit electronic comments to *http://www.regulations.gov.* Submit written comments to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Patrick McDermott, Center for Veterinary Medicine (HFV–530), Food and Drug Administration, 8401 Muirkirk Rd., Laurel, MD 20708, 301–210–4213, *e-mail: patrick.mcdermott@fda.hhs.gov.*