with a specified biological, chemical, radiological, or nuclear agent or agents—in this case, Bacillus anthracis, although there is no current domestic emergency involving anthrax, no current heightened risk of an anthrax attack, and no credible information indicating an imminent threat of an attack involving Bacillus anthracis. Pursuant to section 564(b) of the Federal Food, Drug and Cosmetic Act, 21 U.S.C. 360bbb–3(b), and on the basis of such determination, on October 1, 2008, former Secretary of Health and Human Services, Michael O. Leavitt, declared an emergency justifying the authorization of the emergency use of doxycycline hyclate tablets accompanied by emergency use information subject to the terms of any authorization issued under 21 U.S.C. 360bbb–3(a).[1] Pursuant to section 564(b)(2)(B) of the Federal Food, Drug and Cosmetic Act, 21 U.S.C. 360bbb–3(b), and on the basis of Secretary Chertoff's September 23, 2008 determination, I hereby renew former Secretary Leavitt's October 1, 2008 declaration of an emergency, which I previously renewed on October 1, 2009, justifying the authorization of the emergency use of doxycycline hyclate tablets accompanied by emergency use information subject to the terms of any authorization issued under 21 U.S.C. 360bbb–3(a). I am issuing this notice in accordance with section 564(b)(4) of the Federal Food, Drug, and Cosmetic Act, 21 U.S.C. 360bbb–3(b)(4).

Dated: September 24, 2010.

Kathleen Sebelius,

Secretary.

[FR Doc. 2010–24840 Filed 9–30–10; 11:15 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

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: The parents of children that are born out of wedlock.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
None	1,167,097	1	0.17	198,406.49
Estimated Total Annual Burden Hours:				198,406.49

In compliance with the requirements of Section 506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded 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. E-mail address: *infocollection@acf.hhs.gov.* All requests should be identified by the title of the information collection.

The Department specifically requests comments 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) 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 or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: September 30, 2010.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2010–24893 Filed 10–4–10; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0502]

Agency Information Collection Activities; Proposed Collection; Comment Request; National Consumer Surveys on Understanding the Risks and Benefits of FDA-Regulated Medical Products

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

[1] Pursuant to section 564(b)(4) of the FFDCA, notice of the determination by the Secretary, DHS, and the declaration by the Secretary, HHS, was provided at 73 FR 58242 (October 6, 2008).