Secretary of the Department of Health and Human Services (HHS) (the Secretary) the authority to establish an advisory panel if the Secretary finds the panel necessary and in the public interest. The Secretary signed the charter establishing the Advisory Panel on Medicare Education (the Panel) on January 21, 1999 and the renewed the charter on January 14, 2005. The Panel advises HHS and the Centers for Medicare & Medicaid Services (CMS) on opportunities to enhance the effectiveness of consumer education materials serving the Medicare program.

The goals of the Panel are to provide advice on the following:

- Developing and implementing a national Medicare education program that describes the options for selecting health plans and prescription drug benefits under Medicare.
- Enhancing the Federal government's effectiveness in informing the Medicare consumer, including the appropriate use of public-private partnerships.
- Expanding outreach to vulnerable and underserved communities, including racial and ethnic minorities, in the context of a national Medicare education program.
- Assembling an information base of best practices for helping consumers evaluate health plan options and building a community infrastructure for information, counseling, and assistance.

The Panel shall consist of a maximum of 20 members. The charter requires that meetings be held approximately four times per year. Members are expected to attend all meetings.

This notice is an invitation to interested organizations or individuals to submit their nominations for membership on the Panel. The Secretary, or his designee, will appoint the new members to the Panel from among those candidates determined to have the expertise required to meet specific agency needs, and in a manner to ensure an appropriate balance of membership.

Each nomination must state that the nominee has expressed a willingness to serve as a Panel member and must be accompanied by a resume and a brief summary of the nominee's experience. In order to permit an evaluation of possible sources of conflict of interest, potential candidates will be asked to provide detailed information concerning such matters as financial holdings, consultancies, and research grants or contracts. Self-nominations will also be accepted.

Authority: (Section 222 of the Public Health Service Act (42 U.S.C. 217(a)) and section 10(a) of Pub. L. 92–463 (5 U.S.C. App. 2, section 10(a) and 41 CFR 102–3)

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

Dated: April 6, 2005.

Mark B. McClellan,

Administrator, Centers for Medicare & Medicaid Services.

[FR Doc. 05–7954 Filed 4–21–05; 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: Supporting Healthy Marriage Project Focus Group.

OMB No.: New Collection.

Description: The Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS), is conducting a demonstration and evaluation called the Supporting Health Marriage (SHM) Project. The project is a large-scale, multi-site, multi-year, rigorous test of marriage education programs for interested low-income married couples and is based on a substantial body of research that has shown a relationship between healthy marriages and a variety of positive child and family outcomes. The SHM Project is designed to inform program operators and policymakers of the most effective ways to help couples who voluntarily choose to participate in demonstrations designed to strengthen and maintain healthy marriages. The focus groups will provide key information about the perspectives of low-to-moderate-income couples regarding marriage, relationships, and marriage education programs; assist ACF and program managers in designing responsive healthy marriage programs; and will help to ensure that the project is testing the strongest possible program models for its target populations.

Respondents: The respondents will be selected to represent low-to-moderate income couples in each of the following categories, whose views can help us to design effective SHM programs: Married couples and those planning to marry, couples with and without children, and couples who have had experience with marriage education programs as well as those who have not. There will also be an effort to include African American and Hispanic couples. Focus groups may be divided into separate discussions for those who are married and for those who are planning to marry. They may also be further separated into discussions for couples, for men only, and for women only.

Each focus group will have approximately 10 respondents for a total of 180 respondents over 2 years.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Mixed Gender Focus Group Protocol	30	1	3	90
Men's Focus Group Protocol	30	1	3	90
Women's Focus Group Protocol	30	1	3	90
Estimated Total Annual Burden Hours	90	1	3	270

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: *grjohnson@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, Attn: Desk Officer in ACF, E-mail address: *Katherine_T._Astrich@omb.eop.gov*.

Dated: April 15, 2005.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 05–8051 Filed 4–21–05; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Required Elements for Voluntary Establishment of Paternity Affidavits.

OMB No.: 0970–0171.

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.

Respondents: State and Tribal IV–D birth record agencies.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
None	862,043	Variable	.166	143,099

Estimated Total Annual Burden Hours: 143,099

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: *grjohnson@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, Attn: Desk Officer for ACF, E-mail address: *Katherine_T._Astrich@omb.eop.gov.*

Dated: April 15, 2005.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 05–8052 Filed 4–21–05; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2005N–0058]

Hospira, Inc. et al.; Withdrawal of Approval of 76 New Drug Applications and 60 Abbreviated New Drug Applications; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** of March 4, 2005 (70 FR 10651). The document announced the withdrawal of approval of 76 new drug applications (NDAs) and 60 abbreviated new drug applications (ANDAs). The document inadvertently withdrew approval of ANDA 76–214 for Sotalol Hydrochloride Tablets, 80 milligrams (mg), 120 mg, and 160 mg, held by TorPharm, c/o Apotex Corp., 616 Heathrow Dr., Lincolnshire, IL 60069. FDA confirms that approval of ANDA 76–214 is still in effect.

EFFECTIVE DATE: April 4, 2005.

FOR FURTHER INFORMATION CONTACT: Florine P. Purdie, Center for Drug Evaluation and Research (HFD–7), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–594–2041.

SUPPLEMENTARY INFORMATION: In FR Doc. 05–4158, appearing on page 10651 in the **Federal Register** of Friday, March 4, 2005, the following correction is made:

1. On page 10656, in the table, the entry for ANDA 76–214 is removed.

Dated: April 14, 2005.

Steven Galson, Acting Director.

Center for Drug Evaluation and Research.

[FR Doc. 05–8049 Filed 4–21–05; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

The Eighth Annual FDA–Orange County Regulatory Affairs Educational Conference; ''Reality of Regulatory Affairs''

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of meeting.

The Food and Drug Administration (FDA) is announcing its eighth annual educational conference cosponsored with the Orange County Regulatory Affairs Discussion Group (OCRA). The conference is intended to provide the drug, device, and biologics industries with an opportunity to interact with FDA reviewers and compliance officers from the centers and district offices, as well as other industry experts. The main focus of this interactive conference will be product approval, compliance, and risk management in the three medical product areas. Industry speakers, interactive question and answer and workshop sessions will also be included to assure open exchange and dialogue on the relevant regulatory issues.