children and families and repeating the data collection procedures used in the original study. Selection of a new cohort will allow the comparison of characteristics of children who are entering the child welfare system today with those who entered prior to the implementation of the Adoption and Safe Families Act and prior to the advent of the Child and Family Services Review process. The data collection will follow the same format as that used in previous rounds of data collection, and will employ, with only modest revisions, the same instruments that have been used in previous rounds. Currently, HHS intends to collect baseline data and one follow-up 18 months later, with future follow-up rounds contingent on funding availability. Data from NSCAW are made available to the research community though licensing arrangements from the National Data Archive on Child Abuse and Neglect at Cornell University.

Respondents: 5,700 children and their associated permanent or foster caregivers, caseworkers, and teachers; in addition, an administrator will be interviewed in each location from which children are sampled.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Child Interview	5,700	1	1.2	6,840
Permanent Caregiver Interview	3,800	1	2.0	7,600
Foster Caregiver Interview	1,990	1	1.5	2,985
Caseworker Interview	5,700	1	1.0	5,700
Teacher Questionnaire	3,000	1	.75	2,250
Agency Questionnaire	97	1	1.0	97
Estimated Total Annual Burden Hours:				25,472

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: 202–395–6974, Attn: Desk Officer for ACF.

Dated: December 17, 2007.

Brendan Kelly,

Reports Clearance Officer.

[FR Doc. 07–6143 Filed 12–21–07; 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: Compassion Capital Fund Impact Evaluation Process Study.

OMB No.: New Collection.

Description: The information collection activity proposed under this notice will obtain information about intermediary grantee agencies providing capacity building assistance to faith-based and community organizations under the Compassion Capital Fund (CCF) Demonstration program. The information gathered under this data collection activity will be used to describe the approach and methods used by intermediaries to provide the services that are being evaluated in the CCF impact evaluation. Information collection will be through informal discussions and observations on-site at the organizations, using uniform protocols.

Respondents: Directors and staff providing technical assistance and related services to faith-based and community organizations and directors and staff in faith-based and community organizations that have received capacity building assistance.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Intermediary Protocol for Executive Director	10	1	3	30
Intermediary Protocol for Key Staff	30	1	1	30
Faith-based or Community Organization Protocol for Executive Director	30	1	2	60
Faith-based or Community Organization Protocol for Key Staff	60	1	1	60

Estimated Total Annual Burden Hours: 180.

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: 202–395–6974, Attn: Desk Officer for ACF.

Dated: December 18, 2007.

Brendan Kelly,

Reports Clearance Officer.

[FR Doc. 07–6158 Filed 12–21–07; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2007N–0472]

Agency Emergency Processing Under the Office of Management and Budget Review; Certification to Accompany Drug, Biological Product, and Device Applications or Submissions; 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 December 12, 2007 (72 FR 70599). The document announced that a proposed collection of information had been submitted to the Office of Management and Budget (OMB) for emergency processing under the Paperwork Reduction Act of 1995. The document was published with an error. This document corrects that error.

FOR FURTHER INFORMATION CONTACT: Joyce Strong, Office of Policy, Planning, and Preparedness (HF–27), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–7010.

SUPPLEMENTARY INFORMATION:IN FR DOC. 07–6023, APPEARING ON PAGE 70599 IN THE Federal Register OF WEDNESDAY, DECEMBER 12, 2007, THE FOLLOWING CORRECTION IS MADE: 1. On page 70599, in the third column, in the second full paragraph, the second sentence is corrected to read ''Specifically, at the time of submission of an application under sections 505, 515, or 520(m) of the FD&C Act (21 U.S.C. 355, 360e, or 360j(m)), or under section 351 of the PHS Act (42 U.S.C. 262), or submission of a report under section 510(k) of the FD&C Act (21 U.S.C. 360(k)), such application or submission must be accompanied by a certification that all applicable requirements of section 402(j) of the PHS Act have been met.''

Dated: December 17, 2007.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E7–24914 Filed 12–21–07; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2007C–0474]

DSM Nutritional Products, Inc.; Filing of Color Additive Petition; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration is correcting a notice that appeared in the **Federal Register** of December 4, 2007 (72 FR 68166). The document announced that DSM Nutritional Products, Inc., has filed a petition proposing that the color additive regulations be amended to provide for the safe use of astaxanthin dimethyldisuccinate as a color additive in the feed of salmonid fish to enhance the color of their flesh.

FOR FURTHER INFORMATION CONTACT: Joyce Strong, Office of Policy, Planning, and Preparedness, Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–7010.

SUPPLEMENTARY INFORMATION: In FR Doc. E7–23473, appearing on page 68166 in the **Federal Register** of Tuesday, December 4, 2007, the following correction is made:

1. On page 68166, in the third column, in the heading of the document, ''[Docket No. 2007N–0453]'' is corrected to read ''[Docket No. 2007C–0474]''.

Dated: December 17, 2007.

Laura M. Tarantino,

Director, Office of Food Additive Safety, Center for Food Safety and Applied Nutrition.

[FR Doc. E7–24911 Filed 12–21–07; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Clinical Trial Design for Community-Acquired Pneumonia; Public Workshop

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshop.

SUMMARY: The Food and Drug Administration (FDA) is announcing a public workshop, cosponsored with the Infectious Diseases Society of America (IDSA), regarding scientific issues in clinical trial design for community-acquired pneumonia. This public workshop is intended to provide information for and to gain perspective from health care providers, academia, and industry on various aspects of antimicrobial drug development for community-acquired pneumonia, including diagnosis of community-acquired pneumonia, effect of antimicrobial treatment for community-acquired pneumonia, endpoints for trials of community-acquired pneumonia, and statistical issues in analysis of results of trials in community-acquired pneumonia. The input from this public workshop will help in developing topics for further discussion.

Date and Time: The public workshop will be held on January 17, 2008, from 8 a.m. to 6 p.m. and on January 18, 2008, from 8 a.m. to 5 p.m.

Location: The public workshop will be held at the Crowne Plaza Hotel, Kennedy Room, 8777 Georgia Ave., Silver Spring, MD 20910, 301–589–0800. Seating is limited and available only on a first-come, first-served basis.

Contact Person: Chris Moser or Lori Benner, Center for Drug Evaluation and Research, Office of Antimicrobial Products, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 22, rm. 6413, Silver Spring, MD 20993–0002, 301–796–0767, or 301–796–0849.

Registration: There is no registration fee for the public workshop. Space is limited; therefore, interested parties are encouraged to register early. Seating will be available on a first-come, first-served basis. To register electronically, e-mail registration information (including name, title, firm name, address, telephone, and fax number) to *CAPwkshp@fda.hhs.gov* by January 9, 2008. Persons without access to the Internet can call 301–796–1300 to register. Persons needing a sign language interpreter or other special