PERSON TO CONTACT FOR INFORMATION: Judith Ingram, Press Officer, Telephone: (202) 694–1220.

Shawn Woodhead Werth,

Secretary and Clerk of the Commission.

[FR Doc. 2015–08906 Filed 4–14–15; 4:15 pm]

BILLING CODE 6715–01–P

FEDERAL RESERVE SYSTEM

Notice of Proposals To Engage in or To Acquire Companies Engaged in Permissible Nonbanking Activities; Correction

This notice corrects a notice (FR Doc. 2015–08045) published on pages 18442 and 18443 of the issue for Wednesday, April 8, 2015.

Under the Federal Reserve Bank of San Francisco heading, the entry for Cathay Financial Holding Co., Ltd., Taipei, Taiwan, is revised to read as follows:

A. Federal Reserve Bank of San Francisco (Gerald C. Tsai, Director, Applications and Enforcement) 101 Market Street, San Francisco, California 94105–1579:

1. *Cathay Financial Holding Co., Ltd., Cathay Life Insurance Co., Ltd., Liang Ting Industrial Co., Ltd., Lin Yuan Investment Co., Ltd., Pai Hsing Investment Co., Ltd., Tung Chi Capital Co., Ltd., and Wan Ta Investment Co., Ltd., all in Taipei, Taiwan, and Wan Bao Development Co., Ltd.,* New Taipei, Taiwan; to acquire Conning Holdings Corp., Hartford, Connecticut, and thereby engage in financial and investment advisory activities, and agency transactional services for customer investments, pursuant to sections 225.28(b)(6) and (b)(7).

Comments on this application must be received by April 23, 2015.

Board of Governors of the Federal Reserve System, April 13, 2015.

Michael J. Lewandowski,

Associate Secretary of the Board.

[FR Doc. 2015–08713 Filed 4–15–15; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[CDC–2014–0013; Docket Number NIOSH–274]

Issuance of Final Guidance Publication

AGENCY: National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice of issuance of final guidance publication.

SUMMARY: The National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC), announces the availability of the following publication: *''NIOSH Current Intelligence Bulletin 67: Promoting Health and Preventing Disease and Injury through Workplace Tobacco Policies''* [2015–113].

ADDRESSES: This document may be obtained at the following link: *http://www.cdc.gov/niosh/docs/2015-113/*.

FOR FURTHER INFORMATION CONTACT: Michelle Martin, NIOSH Division of Respiratory Disease Studies, 1095 Willowdale Road, Mailstop H–2900, Morgantown, WV 26505–2888. (304) 285–5734 (not a toll free number).

Dated: April 9, 2015.

John Howard,

Director, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention.

[FR Doc. 2015–08737 Filed 4–15–15; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects: Immediate Disaster Case Management Intake Assessment (hardcopy and electronic versions).

Title: Immediate Disaster Case Management Intake Assessment.

OMB No.: 0970–NEW.

Description: Section 426 of the Robert T. Stafford Disaster Relief and Emergency Assistance Act (Stafford Act), as amended, 42 U.S.C. 5189d authorizes the Federal Emergency Management Agency (FEMA) and the U.S. Department of Health Services' Administration for Children and Families (ACF) to provide Immediate Disaster Case Management (IDCM) services under the federal Disaster Case Management Program (DCMP).

The use of the Electronic Case Management Record System (ECMRS) is aligned with Executive Order of the President 13589 and the memorandum to the Heads of Executive Departments and Agencies M–12–12 from the Office of Management and Budget to ''Promote Efficient Spending to Support Agency Operations.''

The primary purpose of the information collection pertains to ACF/OHSEPR's initiative to improve the intake process and delivery of case management services to individuals and households impacted by a disaster. Further, the information collection will be used to support ACF/OHSEPR's goal to quickly identify critical gaps, resources, needs, and services to support State, local and non-profit capacity for disaster case management and to augment and build capacity where none exists. All information gathered will be exclusively used to inform the delivery of disaster case management services and programmatic strategies and improvements.

Respondents: Individuals impacted by a disaster.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Burden hours per response	Total burden hours
IDCM Intake Assessment	3,500	1	40	2,333

Estimated Total Annual Burden Hours: 2,333 hours or 140,000 minutes

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 Planning, Research and Evaluation, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. Email 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.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2015–08684 Filed 4–15–15; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–N–0672]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Prominent and Conspicuous Mark of Manufacturers on Single-Use Devices

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by May 18, 2015.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–7285, or emailed to *oira_submission@omb.eop.gov.* All comments should be identified with the OMB control number 0910–0577. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 8455 Colesville Rd., COLE–14526, Silver Spring, MD 20993–0002, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Prominent and Conspicuous Mark of Manufacturers On Single-Use Devices (OMB Control Number 0910–0577)—Extension

Section 502 of the Federal Food, Drug, and Cosmetic Act (the FD&C Act) (21 U.S.C. 352), among other things, establishes requirements that the label or labeling of a medical device must meet so that it is not misbranded and subject to regulatory action. Section 301 of the Medical Device User Fee and Modernization Act of 2002 (Pub. L. 107–250) amended section 502 of the FD&C Act to add section 502(u) to require devices (both new and reprocessed) to bear prominently and conspicuously the name of the manufacturer, a generally recognized abbreviation of such name, or a unique and generally recognized symbol identifying the manufacturer.

Section 2(c) of the Medical Device User Fee Stabilization Act of 2005 (Pub. L. 109–43) amends section 502(u) of the FD&C Act by limiting the provision to reprocessed single-use devices (SUDs) and the manufacturers who reprocess them. Under the amended provision, if the original SUD or an attachment to it prominently and conspicuously bears the name of the manufacturer, then the reprocessor of the SUD is required to identify itself by name, abbreviation, or symbol in a prominent and conspicuous manner on the device or attachment to the device. If the original SUD does not prominently and conspicuously bear the name of the manufacturer, the manufacturer who reprocesses the SUD for reuse may identify itself using a detachable label that is intended to be affixed to the patient record.

The requirements of section 502(u) of the FD&C Act impose a minimal burden on industry. This section of the FD&C Act only requires the manufacturer, packer, or distributor of a device to include their name and address on the labeling of a device. This information is readily available to the establishment and easily supplied. From its registration and premarket submission database, FDA estimates that there are 67 establishments that distribute approximately 427 reprocessed SUDs. Each response is anticipated to take 0.1 hours (6 minutes) resulting in a total burden to industry of 43 hours.

In the **Federal Register** of December 30, 2014 (79 FR 78445), FDA published a 60-day notice requesting public comment on the proposed collection of information. No comments were received.

FDA estimates the burden of this collection of information as follows:

TABLE 1—ESTIMATED ANNUAL THIRD-PARTY DISCLOSURE BURDEN [1] [2]

Type of respondent	Number of respondents	Number of disclosures per respondent	Total annual disclosures	Average burden per disclosure	Total hours
Establishments listing fewer than 10 SUDs	58	2	116	0.1 (6 minutes)	12
Establishments listing 10 or more SUDs	9	34	306	0.1 (6 minutes)	31
Total					43

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.

[2] Numbers have been rounded.