and NIOSH made the final determination regarding proposed additions to the 2016 hazardous drug list.

NIOSH reviewed the recommendations of the peer reviewers and stakeholders and determined that 33 drugs in addition to the 3 drugs with manufacturer's warnings, were determined to have one or more characteristics of a hazardous drug and this list of 36 drugs is being published for comment in CDC–2015–0034 and NIOSH Docket Number 233–A. The list of proposed additions can be found at *www.regulations.gov.*

Dated: May 20, 2015.

John Howard,

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

[FR Doc. 2015–12857 Filed 5–27–15; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–15–15KZ]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The notice for the proposed information collection is published to obtain comments from the public and affected agencies.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address any of the following: (a) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; (b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (c) Enhance the quality, utility, and clarity of the information to be collected; (d) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submission of responses; and (e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570 or send an email to *omb@cdc.gov.* Written comments and/or suggestions regarding the items contained in this notice should be directed to the Attention: CDC Desk Officer, Office of Management and Budget, Washington, DC 20503 or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

Research on the Efficacy and Feasibility of Essentials for Parenting Toddlers and Preschoolers—New—National Center for Injury Prevention and Control (NCIPC), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Centers for Disease Control and Prevention (CDC) seeks a two-year OMB approval to conduct a new information collection for a study entitled, ''Research on the Efficacy and Feasibility of Essentials for Parenting Toddlers and Preschoolers''.

Child maltreatment is both widespread and impactful. It is estimated that 1 in 58 U.S. children had been maltreated in a 1-year period (*i.e.,* victims of physical, sexual, and emotional abuse or neglect). Millions of other American children are exposed to maltreatment that does not meet thresholds for clinical significance, but is nonetheless detrimental to child health.

Parent training is arguably the single most effective prevention initiative developed to date. Although there are potentially far-reaching impacts of parent training to improve public health, empirically-supported parent training is not widely available. The public health challenge is how to make the content of these empirically-supported parent training programs—which largely focus on the same parenting skills and approaches—accessible to the majority of American parents.

To leverage the strength of empirically supported parent training as a broadly disseminated prevention tool, the CDC has developed a resource tool called ''Essentials for Parenting Toddlers and Preschoolers (EFP)''. This web-based resource includes the typical content of empirically supported parent training programs and uses a psychoeducational approach including modeling (through its videos) and practice (through its activities).

This study is an empirical evaluation using an intensive repeated measures design to test the efficacy, feasibility, and use of EFP as administered in guided and unguided formats. The proposed data collection fits into NCIPC's research agenda's priorities in preventing child maltreatment.

There are no costs to respondents other than their time. The total estimated annual burden hours are 2,050.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Parents (both Natural Navigation [NN] and Guided Navigation [GN] groups).	Form 1—Screening and Demographics Questionnaires—Attachment I1.	400	1	15/60
	Form 2—Detailed Assessment Measures—Attachment I2.	200	2	45/60
	Form 3—Core Assessment Measures (Rotating)—Attachment I3.	200	18	15/60
	Form 4—Parental EFP Skills Knowledge Scale—Attachment I4.	200	8	15/60
	Form 5—Parental EFP Skills Usefulness Scale—Attachment I5.	200	6	15/60

ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
	Form 6—Therapy Attitude Inventory and System Usability Scale—Attachment I6.	200	1	15/60

Leroy A. Richardson,

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science,Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2015–12809 Filed 5–27–15; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: Head Start Facilities Construction, Purchase and Major Renovation.

OMB No.: 0970–0193.

Description: The Office of Head Start within the Administration for Children and Families, United States Department of Health and Human Services, is proposing to renew authority to collect information on funding for the purchase, construction or renovation of facilities. All information is collected electronically through the Head Start Enterprise System (HSES). The information required is in conformance with Section 644(f) and (g) of the Act. Federal funding officials use the information to determine that the proposed purchase has resulted in savings when compared to the costs that would be incurred to acquire the use of an alternative facility, or that the lack of alternative facilities will prevent, or would have prevented, the operation of the program. The rule further describes the assurances which are necessary to protect the Federal interest in real property and the conditions under which federal interest may be subordinated and protected when grantees make use of debt instruments when purchasing facilities. The information is used by funding officials to determine if grantee's arrangements adequately conform to other applicable statutes which apply to the expenditure of public funds for the purchase of real property.

Respondents: Head Start and Early Head Start program grant recipients.

ANNUAL BURDEN ESTIMATES

Instruments	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Administrative Requirements	225	1	41	9225

Estimated Total Annual Burden Hours: 9225.

Cost per respondent is $40 estimated at 2 hours × $20.00 per hour.

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. 2015–12924 Filed 5–27–15; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–D–0369]

M7 Assessment and Control of DNA Reactive (Mutagenic) Impurities in Pharmaceuticals to Limit Potential Carcinogenic Risk; International Conference on Harmonisation; Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of a guidance entitled ''M7 Assessment and Control of DNA Reactive (Mutagenic) Impurities in Pharmaceuticals to Limit Potential Carcinogenic Risk.'' The guidance was prepared under the auspices of the International Conference on Harmonisation of Technical