Dated: November 6, 2006.

Joan F. Karr,

Acting Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. E6–19146 Filed 11–13–06; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–07–0604]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639–5960 or send an e-mail to *omb@cdc.gov*. Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395–6974. Written comments should be received within 30 days of this notice.

Proposed Project

School Associated Violent Death Surveillance System—Extension—National Center for Injury Prevention and Control (NCIPC), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Division of Violence Prevention (DVP), National Center for Injury Prevention and Control (NCIPC) proposes to maintain a system for the surveillance of school-associated homicides and suicides. The system will rely on existing public records and interviews with law enforcement officials and school officials. The purpose of the system is to (1) estimate the rate of school-associated violent death in the United States and (2) identify common features of school-associated violent deaths. The system will contribute to the understanding of fatal violence associated with schools, guide further research in the area, and help direct ongoing and future prevention programs.

Violence is the leading cause of death among young people, and increasingly recognized as an important public health and social issue. In 1998, over 3,500 school aged children (5 to 18 years old) in the United States died violent deaths due to suicide, homicide, and unintentional firearm injuries. The vast majority of these fatal injuries were not school associated. However, whenever a homicide or suicide occurs in or around school, it becomes a matter of particularly intense public interest and concern. NCIPC conducted the first scientific study of school-associated violent deaths during the 1992–99 academic years to establish the true extent of this highly visible problem. Despite the important role of schools as a setting for violence research and prevention interventions, relatively little scientific or systematic work has been done to describe the nature and level of fatal violence associated with schools. Until NCIPC conducted the first nationwide investigation of violent deaths associated with schools, public health and education officials had to rely on limited local studies and estimated numbers to describe the extent of school-associated violent death.

The system will draw cases from the entire United States in attempting to capture all cases of school-associated violent deaths that have occurred. Investigators will review public records and published press reports concerning each school-associated violent death. For each identified case, investigators will also interview an investigating law enforcement official (defined as a police officer, police chief, or district attorney), and a school official (defined as a school principal, school superintendent, school counselor, school teacher, or school support staff) who are knowledgeable about the case in question. Researchers will request information on both the victim and alleged offender(s)—including demographic data, their academic and criminal records, and their relationship to one another. They will also collect data on the time and location of the death; the circumstances, motive, and method of the fatal injury; and the security and violence prevention activities in the school and community where the death occurred, before and after the fatal injury event. There are no costs to the respondents other than their time. The total estimated annualized burden hours are 70.

ESTIMATED ANNUALIZED BURDEN HOURS

Respondents	Number of respondents	Number of responses per respondent	Average burden/response (in hours)
School Officials	35	1	1
Police Officials	35	1	1

Dated: November 7, 2006.

Joan F. Karr,

Acting Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. E6–19147 Filed 11–13–06; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Committee for Injury Prevention and Control: Notice of Charter Renewal

This gives notice under the Federal Advisory Committee Act (Pub. L. 92–463) of October 6, 1972, that the Advisory Committee for Injury Prevention and Control, Centers for Disease Control and Prevention, Department of Health and Human Services, has been renewed for a 2-year period through October 28, 2008.

For information, contact Amy Harris, Executive Secretary, Advisory Committee for Injury Prevention and Control, Centers for Disease Control and Prevention, Department of Health and Human Services, 1600 Clifton Road, NE., Mailstop K61, Atlanta, Georgia 30333, telephone 770/488–1484 or fax 770/488–4222.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of

meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dated: November 6, 2006.

Alvin Hall,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. E6–19151 Filed 11–13–06; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2006N–0329]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Medicated Feed Mill License Application

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 December 14, 2006

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: 202–395–6974.

FOR FURTHER INFORMATION CONTACT: Denver Presley, Jr., Office of the Chief Information Officer (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1472.

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.

Medicated Feed Mill Licensing Application—21 CFR Part 515 (OMB Control No. 0910–0337)—Extension

The Animal Drug Availability Act (ADAA) of October 9, 1996, amended section 512 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b) to replace the system for the approval of specific medicated feed with a general licensing system for feed mills. Before passage of the ADAA, medicated feed manufacturers were required to obtain approval of Medicated Feed Applications (MFAs) in order to manufacture certain types of medicated feeds. An individual approved MFA was required for each and every applicable medicated feed. The ADAA streamlined the paperwork process for gaining approval to manufacture medicated feeds by replacing the MFA system with a facility license for each medicated feed manufacturing facility. Implementing regulations are at part 515 (21 CFR part 515).

In the **Federal Register** of August 25, 2006 (71 FR 50433), FDA solicited comments on the information collection provisions of this proposed collection. In response to that request, FDA received no comments.

Description of Respondents: Medicated feed manufacturers.

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

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

21 CFR Section	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
515.10(b)	7	1	7	0.25	1.75
515.11(b)	100	1	100	0.25	25
515.23	25	1	25	0.25	6.25
515.30(c)	0.15	1	0.15	24	3.6
Total					36.6

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

TABLE 2.—ESTIMATED ANNUAL RECORDKEEPING BURDEN[1]

21 CFR Section	No. of Recordkeepers	Annual Frequency per Recordkeeping	Total Annual Records	Hours per Recordkeeper	Total Hours
510.305	1,070	1	1,070	0.03	32.10

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

The estimated annual reporting burden on industry is 36.6 hours as shown in table 1 of this document. Industry estimates it takes about 1/4 hour to submit the application. We estimate 132 original and supplemental applications, and voluntary revocations for a total of 33 hours (132 submissions x 1/4 hour). An additional 3.6 hours is added for the rare notice of opportunity for a hearing to not approve or revoke an application. Finally, we estimate 36 hours for maintaining and retrieving labels as required by 21 CFR 510.305. We estimated .03 hours for each of approximately 1,070 licensees. Thus, the total burden for recordkeeping requirements is 32.10 hours (1,070 x 0.03).

Dated: November 7, 2006.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E6–19152 Filed 11–13–06; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2006D–0441]

Draft Guidance for Industry: Protocols for the Conduct of Method Transfer Studies for Type C Medicated Feed Assay Methods; Availability

AGENCY: Food and Drug Administration, HHS.