Technological and administrative changes necessitate that a new toll-free telephone number and a new, more sophisticated routing system be operational by January 1, 2005. This entails collecting administrative information from each of the 144 crisis

centers currently participating in the network (e.g., location, hours of operation, call capacity, non-English language capability) and programming the data into a new routing system to ensure that each caller is linked to the nearest and most appropriate crisis

center. The form developed to secure this information requests only factual information that is essential to the design of the routing system.

The following table is the estimated hour burden:

Number of respondents	Responses/ respondent	Burden/ response (hrs.)	Total burden hours
144	1	.17	24.50

Emergency approval is being requested because SAMHSA does not want to risk the possibility of disrupting the functioning of the national suicide prevention hotline network, and the subsequent possibility that even one individual who tries calling a non-working hotline might ultimately take his/her own life.

Written comments and recommendations concerning the proposed information collection should be sent within two weeks of this notice to: John Kraemer, Human Resources and Housing Branch, Office of Management and Budget, New Executive Office Building, Room 10235, Washington, DC 20503; due to potential delays in OMB's receipt and processing of mail sent through the U.S. Postal Service, respondents are encouraged to submit comments by fax to: 202–395–6974.

Dated: November 4, 2004.

### Patricia S. Bransford,

Acting Executive Officer, SAMHSA.
[FR Doc. 04–25028 Filed 11–9–04; 8:45 am]
BILLING CODE 4162–20–P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

[Request for Application 05017]

### Intervention and Evaluation Trials To Prevent Intimate Partner Violence; Notice of Availability of Funds-Amendment

A notice announcing the availability of fiscal year (FY) 2005 funds for a research cooperative agreement program to conduct efficacy and effectiveness trials of intervention strategies to prevent intimate partner violence and/or its negative consequences, particularly studies of strategies that have not been well studied, for at-risk or underserved populations was published in the **Federal Register** on October 27, 2004, Vol. 69, No. 207, pages 62694–62701. The notice is

amended as follows: On page 62696, Column 3, Line 2, delete \$1,800,000 and replace with the new amount of \$2,250,000.

Dated: November 3, 2004.

#### William Nichols,

Acting Director, Procurement and Grants Office, Centers for Disease Control and Prevention.

[FR Doc. 04–25027 Filed 11–9–04; 8:45 am] BILLING CODE 4163–18–P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

## Centers for Disease Control and Prevention

# Public Health Service Act; Delegation of Authority

Notice is hereby given that I have delegated to the Director, Centers for Disease Control and Prevention (CDC), the authority vested in the Secretary of Health and Human Services under Section 319F–2 of Title III of the Public Health Service Act, as amended, to enter into agreements with recipients of the Stockpile material will be deployed. The authority to deploy the Stockpile referred to herein is limited to the CHEMPACK Program.

This authority cannot be redelegated. Further, CDC must notify the Office of Public Health Emergency Preparedness before entering into an agreement.

This delegation became effective upon date of signature. In addition, I have affirmed and ratified any actions taken by the Director, CDC, or his/her subordinates which involved the exercise of the authorities delegated therein prior to the effective date of the delegation.

Dated: October 27, 2004.

### Tommy G. Thompson,

Secretary.

[FR Doc. 04–25002 Filed 11–9–04; 8:45 am] BILLING CODE 4160–18–M

### DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. 2004N-0470]

Agency Information Collection Activities: Proposed Collection; Comment Request; New Animal Drugs For Investigational Use

**AGENCY:** Food and Drug Administration,

HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the Federal Register concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the reporting and recordkeeping requirements for "New Animal Drugs for Investigational Use."

**DATES:** Submit written or electronic comments on the collection of information by January 10, 2005.

ADDRESSES: Submit electronic comments on the collection of information to: http://www.fda.gov/dockets/ecomments. Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

#### FOR FURTHER INFORMATION CONTACT:

Denver Presley, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1472.

**SUPPLEMENTARY INFORMATION:** Under the PRA (44 U.S.C. 3501–3520), Federal