recommendations in VICH GL9 when differences among the guidances occur.

This Level 1 final guidance is being issued consistent with FDA's good guidance practices (21 CFR 10.115; 65 FR 56468, September 19, 2000). This guidance document represents FDA's current thinking on design and conduct of all clinical studies of veterinary products in the target species. It does not create or confer any rights for or on any person, and does not operate to bind FDA or the public. An alternative method may be used as long as it satisfies the requirements of applicable statutes and regulations.

Information collected is covered under OMB control number 0910–0032.

III. Electronic Access

Copies of the final guidance documents entitled ''Good Clinical Practice'' (VICH GL9) may be obtained on the Internet from the CVM home page at http://www.fda.gov/cvm. Comments may also be submitted electronically on the Internet at http://www.fda.gov/dockets/ecomments. Once on this Internet site, select ''99D–2406 Good Clinical Practice'' and follow the directions.

IV. Comments

As with all of FDA's guidances, the public is encouraged to submit written comments with new data or other new information pertinent to this final guidance. FDA will periodically review the comments in the docket and, where appropriate, will amend this guidance. The agency will notify the public of any such amendments through a notice in the **Federal Register**.

Interested persons may submit to the Dockets Management Branch (address above) written or electronic comments regarding this final guidance document at any time. Two copies of any comments are to be submitted, except that individuals may submit one copy. Comments should be identified with the docket number found in brackets in the heading of this document. A copy of this final guidance document and received comments are available in the Dockets Management Branch between 9 a.m. and 4 p.m., Monday through Friday.

Dated: May 7, 2001.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 01–12092 Filed 5–14–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

[HCFA–R–235]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Health Care Financing Administration, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Health Care Financing Administration (HCFA), Department of Health and Human Services, is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Type of Information Collection Request: Revision of a currently approved collection;

Title of Information Collection: Data Use Agreement and Information Collection Requirements, model language, and Supporting Regulations in 45 CFR, part 5b;

Form No.: HCFA–R–235 (OMB# 0938–0734);

Use: This agreement is used as a binding agreement stating conditions under which HCFA will disclose and user will maintain HCFA data that are protected by the Privacy Act.;

Frequency: On occasion;

Affected Public: Not-for-profit institutions;

Number of Respondents: 1,500;

Total Annual Responses: 1,500;

Total Annual Hours: 750.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access HCFA's Web Site address at *http://www.hcfa.gov/regs/prdact95.htm,* or E-mail your request, including your address, phone number, OMB number, and HCFA document identifier, to *Paperwork@hcfa.gov,* or call the Reports Clearance Office on (410) 786–1326. Written comments and recommendations for the proposed information collections must be mailed within 60 days of this notice directly to the HCFA Paperwork Clearance Officer designated at the following address: HCFA, Office of Information Services, Security and Standards Group, Division of HCFA Enterprise Standards, Attention: Dawn Willinghan, HCFA–R–235, Room N2–14–26, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

Dated: April 24, 2001.

John P. Burke, III,

HCFA Reports Clearance Officer, HCFA Office of Information Services, Security and Standards Group, Division of HCFA Enterprise Standards.

[FR Doc. 01–12203 Filed 5–14–01; 8:45 am]

BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Special Projects of National Significance; Targeted HIV Outreach and Intervention Model Development; Evaluation and Program Support Center

AGENCY: Health Resources and Services Administration, HHS.

ACTION: Correction.

In the **Federal Register** of April 13, 2001, appearing on page 19180, second column, line 21 is an incorrect website to locate the guidance. The correct HRSA web site for the guidance should read, *www.hab.hrsa.gov/grants.html.*

Dated: May 9, 2001.

Jane M. Harrison,

Director, Division of Policy Review and Coordination.

[FR Doc. 01–12180 Filed 5–14–01; 8:45 am]

BILLING CODE 4160–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Advisory Committee; Notice of Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Public Law 92–463), announcement is made of the following National Advisory body scheduled to meet during the month of June 2001.

Name: Maternal and Child Health Research Grants Review Committee.

Date and Time: June 20–22, 2001; 8 am–5 pm.