

ANNUAL BURDEN ESTIMATES

Instrument	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Total annual burden hours
Discussion Guide for Use with Tribal TANF administrators .....	13	1	2	26
Discussion Guide for Use with Tribal TANF staff .....	12	1	1	12
Discussion Guide for Focus Groups with Tribal TANF clients .....	20	1	2	40
Discussion Guide for Use with staff of related programs .....	20	1	1	20
All Instruments .....	65	.....	.....	98

*Estimated Total Annual Burden Hours:* 98.

In compliance with the requirements of Section 3506(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: OPRE Reports Clearance Officer. Email address: [OPREinfocollection@acf.hhs.gov](mailto:OPREinfocollection@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.

Dated: October 26, 2011.

**Steven M. Hanmer,**

*Reports Clearance, Officer.*

[FR Doc. 2011-28273 Filed 11-1-11; 8:45 am]

**BILLING CODE 4184-09-M**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Administration for Children and Families**

**President's Committee for People With Intellectual Disabilities Meeting, Via Conference Call, Cancellation**

**AGENCY:** President's Committee for People with Intellectual Disabilities (PCPID).

**ACTION:** Notice of PCPID Conference Call Cancellation.

**DATES:** The conference call was scheduled for October 28, 2011, 1 p.m. to 2:30 p.m.

**FOR FURTHER INFORMATION CONTACT:** Laverdia Taylor Roach, Senior Advisor, President's Committee for People with Intellectual Disabilities, The Aerospace Center, Second Floor West, 370 L'Enfant Promenade SW., Washington, DC 20447. Telephone: (202) 619-0634. Fax: (202) 205-9519. Email: [LRoach@acf.hhs.gov](mailto:LRoach@acf.hhs.gov).

Further meetings will be announced through a separate **Federal Register** notice.

Dated: October 26, 2011.

**Jamie Kendall,**

*Deputy Commissioner, Administration on Developmental Disabilities.*

[FR Doc. 2011-28292 Filed 11-1-11; 8:45 am]

**BILLING CODE 4184-01-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

[Docket No. FDA-1999-D-2955]

**Revised Guidance for Industry on Impurities: Residual Solvents in New Veterinary Medicinal Products, Active Substances and Excipients (Revision), VICH GL18(R); Availability**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing the

availability of a revised guidance for industry (#100) entitled "Impurities: Residual Solvents in New Veterinary Medicinal Products, Active Substances and Excipients (Revision)" VICH GL18(R). This revised guidance has been developed for veterinary use by the International Cooperation on Harmonisation of Technical Requirements for Registration of Veterinary Medicinal Products (VICH). The guidance is intended to recommend acceptable amounts of residual solvents in new animal drugs (referred to as pharmaceuticals or veterinary medicinal products in this guidance) for the safety of the target animal as well as for the safety of human consumers of products derived from treated food producing animals. It is intended to assist in developing new animal drug applications (referred to as marketing applications in this guidance) submitted to the European Union, Japan, and the United States.

**DATES:** Submit either electronic or written comments on Agency guidances at any time.

**ADDRESSES:** Submit written requests for single copies of the guidance to the Communications Staff (HFV-12), Center for Veterinary Medicine, Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855. Send one self-addressed adhesive label to assist that office in processing your requests. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the guidance document.

Submit electronic comments on the guidance to <http://www.regulations.gov>. Submit written comments to the Division of Dockets Management (HFA-305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

**FOR FURTHER INFORMATION CONTACT:** Mai Huynh, Center for Veterinary Medicine (HFV-142), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, (240) 276-8273, [mai.huynh@fda.hhs.gov](mailto:mai.huynh@fda.hhs.gov).

**SUPPLEMENTARY INFORMATION:**

## I. Background

In recent years, many important initiatives have been undertaken by regulatory authorities and industry associations to promote the international harmonization of regulatory requirements. FDA has participated in efforts to enhance harmonization and has expressed its commitment to seek scientifically based harmonized technical procedures for the development of pharmaceutical products. One of the goals of harmonization is to identify and then reduce differences in technical requirements for drug development among regulatory agencies in different countries.

FDA has actively participated in the International Conference on Harmonization of Technical Requirements for Approval of Pharmaceuticals for Human Use for several years to develop harmonized technical requirements for the approval of human pharmaceutical and biological products among the European Union, Japan, and the United States. The VICH is a parallel initiative for veterinary medicinal products. The VICH is concerned with developing harmonized technical requirements for the approval of veterinary medicinal products in the European Union, Japan, and the United States, and includes input from both regulatory and industry representatives.

The VICH Steering Committee is composed of member representatives from the European Commission, European Medicines Evaluation Agency, European Federation of Animal Health, Committee on Veterinary Medicinal Products, the U.S. Food and Drug Administration, the U.S. Department of Agriculture, the Animal Health Institute, the Japanese Veterinary Pharmaceutical Association, the Japanese Association of Veterinary Biologics, and the Japanese Ministry of Agriculture, Forestry, and Fisheries.

Four observers are eligible to participate in the VICH Steering Committee: One representative from the government of Australia/New Zealand, one representative from the industry in Australia/New Zealand, one representative from the government of Canada, and one representative from the industry of Canada. The VICH Secretariat, which coordinates the preparation of documentation, is provided by the International Federation for Animal Health (IFAH). An IFAH representative also participates in the VICH Steering Committee meetings.

## II. Revised Guidance on Residual Solvents in New Veterinary Medicinal Products, Active Substances and Excipients

In the **Federal Register** of August 17, 2010 (75 FR 50771), FDA published a notice of availability for a draft revised guidance entitled "Residual Solvents in New Veterinary Medicinal Products, Active Substances and Excipients (Revision) VICH GL18(R)" giving interested persons until October 18, 2010, to comment on the draft revised guidance. This draft incorporated a lower permissible daily exposure limit for N-Methylpyrrolidone, which is still being kept in Class 2, and placed tetrahydrofuran into Class 2 from Class 3. Based on comments received from the draft revised guidance, additional information was added in section 3.2 of this guidance to include reference to the International Conference on Harmonisation of Technical Requirements for Registration of Pharmaceuticals for Human Use guideline entitled "Impurities: Guideline for Residual Solvents (Q3C(R4))." The revised guidance announced in this notice finalizes the draft revised guidance announced on August 17, 2010. The revised guidance is a product of the Quality Expert Working Group of the VICH.

## III. Paperwork Reduction Act of 1995

This revised guidance refers to previously approved collections of information found in FDA regulations. These collections of information are subject to review by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (44 U.S.C. 3501–3520). The collections of information in this document have been approved under OMB control number 0910–0032.

## IV. Significance of Guidance

This revised document, developed under the VICH process, has been revised to conform to FDA's good guidance practices regulation (21 CFR 10.115). For example, the document has been designated "guidance" rather than "guideline". In addition, guidance documents must not include mandatory language such as "shall", "must", "require", or "requirement", unless FDA is using these words to describe a statutory or regulatory requirement.

The revised VICH guidance (GFI #100) is consistent with the Agency's current thinking on this topic. This guidance does not create or confer any rights for or on any person and will 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.

## V. Comments

Interested persons may, at any time, submit either electronic or written comments regarding this revised guidance document to the Division of Dockets Management (see **ADDRESSES**). It is only necessary to send one set of comments. It is no longer necessary to send two copies of mailed comments. Identify comments with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

## VI. Electronic Access

Persons with access to the Internet may obtain the guidance at either <http://www.fda.gov/AnimalVeterinary/GuidanceComplianceEnforcement/GuidanceforIndustry/default.htm> or <http://www.regulations.gov>.

Dated: October 27, 2011.

**Leslie Kux,**

*Acting Assistant Commissioner for Policy.*

[FR Doc. 2011–28371 Filed 11–1–11; 8:45 am]

**BILLING CODE 4160–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

### Interagency Autism Coordinating Committee; Call for Nominations

In accordance with Public Law 112–32, The Combating Autism Reauthorization Act the Department of Health and Human Services has been authorized to continue to support the Interagency Autism Coordinating Committee (IACC) until September 30, 2014 and is seeking nominations for public membership on this committee. The Secretary of the Department of Health and Human Services, who will make the final selections and appointments of public members, has directed the Office of Autism Research Coordination (OARC) to assist the Department in conducting an open and transparent nomination process. Nominations of new public members are encouraged, but current members may also be re-nominated to continue to serve. Self-nominations and nominations of other individuals are both permitted. Only one nomination per individual is required. Multiple nominations for the same individual will not increase likelihood of selection. The Secretary may select public members from the pool of submitted