Respondents	No. of respondents	No. of responses/ respondent	Average burden/ response (in hrs)
D. Followback for evaluations without onsite evaluations:			
Year 1	75	1	10/60
Year 2	75	1	15/60

Dated: August 10, 2004.

Alvin Hall,

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

[FR Doc. 04–18677 Filed 8–13–04; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Notice of Public Consultation

AGENCY: Administration for Native Americans (ANA).

ACTION: Notice of Public Consultation.

SUMMARY: The Administration for Children and Families (ACF) will be holding a half-day Tribal Consultation Session on September 20, 2004 at the Rayburn House Office Building in Washington, DC.

DATES: September 20, 2004.

FOR FURTHER INFORMATION CONTACT: Kim Vigue, Administration for Native Americans, toll free at 1–877–922–9262 or *www.masterkeyconsulting.com/acfconference.*

SUBMISSION INFORMATION: Tribal leaders and representatives interested in submitting written testimony or topics to be discussed on the Consultation Session agenda should contact Kim Vigue toll free at 1–877–922–9262.

If you are proposing a topic to be addressed in the Consultation Session, please be sure to include a brief description of the topic area along with the name and contact information of a suggested presenter.

The public record will remain open for 60 days following the September 20, 2004 consultation. Written comment and testimony can be submitted until November 19, 2004.

SUPPLEMENTARY INFORMATION:

The Administration for Children and Families would like to invite Tribal leaders to participate in a formal consultation Session with ACF senior officials and program directors. The Consultation Session will take place Monday, September 20, 2004 from 8:30 a.m. to 12:30 p.m. in Rayburn House Office Building Room B–339.

The intent of this Consultation Session is to allow ACF officials to hear first hand from Tribal leaders and representatives of Tribal organizations and Native Americans non-profit organizations about the implementation of ACF programs in Native Americans communities. Of particular interest are the challenges that Tribes and Tribal organizations face in accessing ACF program funding and using program funding to support social and economic development activities in Native American communities. ACF offices such as the Administration for Native Americans, Office of Child Support Enforcement, Office of Community Services, Office of Family Assistance, Child Care Bureau, Children's Bureau, Head Start Bureau, and the Family and Youth Services Bureau will be represented.

Because of the limited time, ACF has collaborated with Master Key Consulting to plan and facilitate the session. Master Key Consulting will be responsible for coordinating the stakeholders who wish to participate in the Consultation Session and will work with a planning committee to develop a structured agenda, identifying key issues to be raised and spokespersons to present testimony on the issues.

Dated: August 6, 2004.

Quanah Crossland Stamps,

Commissioner, Administration for Native Americans.

[FR Doc. 04–18588 Filed 8–13–04; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2004N–0355]

Scientific Considerations Related to Developing Follow-On Protein Products

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshop.

SUMMARY: The Food and Drug Administration (FDA) is announcing a public workshop on scientific and technical considerations related to the development of follow-on protein pharmaceutical products. The agency is planning to develop draft guidance on this topic during the coming year. The purpose of this workshop is to obtain input from interested persons on the topics outlined in this document related to developing and approving follow-on protein pharmaceutical products. The agency will consider presentations made at the workshop and comments submitted to the docket before and after the workshop when developing the draft guidance.

DATES: The public workshop will be held on Tuesday, September 14, 2004, from 8:30 a.m. to 5 p.m. and Wednesday, September 15, 2004 from 8 a.m. to 12 noon. Submit requests to make a presentation by September 7, 2004.

ADDRESSES: The public workshop will be held at the University of Maryland—Shady Grove Conference Center, 9630 Gudelsky Dr., Rockville, MD 20850.

Submit written comments on scientific topics related to follow-on protein products to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Two copies of any comments are to be submitted, except that individuals may submit one copy. Comments are to be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT:

To register to present: Marilyn Welschenbach, Center for Drug Evaluation and Research (HFD–121), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20852, 301–443–5089, FAX: 301–443–5245, e-mail: *Marilyn.Welschenbach@fda.gov.*

With regard to the scientific topics outlined in this notice: Keith Webber, Center for Drug Evaluation and Research, Food and Drug Administration (HFD–121), 5600 Fishers Lane, Rockville, MD 20852, 301–443–5089, FAX: 301–443–5234, e-mail: *Keith.Webber@fda.gov*, or Chris Joneckis, Center for Biologics Evaluation and Research (HFM–1),