DEPARTMENT OF HEALTH AND HUMAN SERVICES
National Institutes of Health

National Institute of Mental Health; Notice of Meeting

Pursuant to section 10(a) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of an Interagency Autism Coordinating Committee (IACC or Committee) meeting. The purpose of the IACC meeting is to discuss committee business, updates and issues related to autism spectrum disorder (ASD) research and services activities. The meeting will be open to the public and will be accessible by webcast and conference call.

Name of Committee: Interagency Autism Coordinating Committee (IACC).
Type of Meeting: Open Meeting.
Place: The National Institutes of Health, 31 Center Drive/Building 31, 6th Floor, C Wing, Conference Room 10, Bethesda, MD 20892.
Cost: The meeting is free and open to the public.
Registration: Pre-registration is recommended to expedite check-in. Seating in the meeting room is limited to room capacity and on a first come, first served basis. To register, please visit: https://pointpass.com/events/Interagency_Autism_Coordinating_Committee_July_9_2013/

Deadlines: Notification of intent to present oral comments: Friday, June 28, 2013 by 5:00 p.m. ET. Submission of written/electronic statement for oral comments: Tuesday, July 2, 2013 by 5:00 p.m. ET. Final Deadline for Submission of written comments: Tuesday, July 2, 2013 by 5:00 p.m. ET. Please note: The NIMH Office of Autism Research Coordination (OARC) anticipates that written public comments received by 5:00 p.m. ET, Friday, June 14, 2013 will be provided to the Committee prior to the meeting. Written comments may be sent to the Contact Person listed on this notice by 5:00 p.m. ET on Tuesday, July 2, 2013. The comments should include the name, address, telephone number and when applicable, the business or professional affiliation of the participant. Participants must provide for oral comment statements received between June 14, 2013 and July 2, 2013 will be provided to the Committee either before or after the meeting, depending on the volume of comments received and the staff time required to process them in accordance with privacy regulations and other applicable Federal policies. Written public comments will be provided to the IACC for their consideration and will become part of the public record.

Public Comments: Any member of the public interested in presenting oral comments to the Committee must notify the Contact Person listed on this notice by 5:00 p.m. ET on Friday, June 28, 2013, with their request to present oral comments at the meeting. Interested individuals and representatives of organizations must submit a written/electronic copy of the oral presentation/statement including a brief description of the organization represented by 5:00 p.m. ET on Tuesday, July 2, 2013. Statements submitted will become a part of the public record. Only one representative of an organization will be allowed to present oral comments and presentations will be limited to three to five minutes per speaker, depending on number of speakers to be accommodated within the allotted time. Speakers will be assigned a time to speak in the order of the date and time when their request is received, along with the required submission of the written/electronic statement by the specified deadline.

In addition, any interested person may submit written comments to the IACC prior to the meeting by sending the comments to the Contact Person listed on this notice by 5:00 p.m. ET on Tuesday, July 2, 2013. The comments should include the name, address, telephone number and when applicable, the business or professional affiliation of the participant. Participants must provide for presentation and oral comment statements received by the deadlines for both oral and written public comments will be provided to the IACC for their consideration and will become part of the public record.

In the 2009 IACC Strategic Plan, the IACC listed the “Spirit of Collaboration” as one of its core values, stating that, “We will treat others with respect, listen to diverse views with open minds, discuss submitted public comments, and frame discussions where participants can comfortably offer opposing opinions.” In keeping with this core value, the IACC and the NIMH Office of Autism Research Coordination (OARC) ask that members of the public who provide public comments or participate in meetings for the IACC also seek to treat others with respect and consideration in their communications and actions, even when discussing issues of genuine concern or disagreement.

Remote Access: The meeting will be open to the public through a conference call phone number and access: 512-248-7574. Contact Person: Carolyn A. Baum, Program Analyst, Office of Federal Advisory Committee Policy.
Committee Policy.

The Substance Abuse and Mental Health Services Administration (SAMHSA) is requesting OMB approval for a revision to the Behavioral Health Web site and Resources data collection. SAMHSA is authorized under section 501(d)(16) of the Public Health Service Act (42 USC 290aa(d)(16)) to develop and distribute materials for the prevention, treatment, and recovery from substance abuse and mental health disorders. To improve customer service and lessen the burden on the public to locate and obtain these materials, SAMHSA has developed a Web site that includes more than 1,400 free publications from SAMHSA and its component Agencies: The Center for Substance Abuse Treatment, the Center for Substance Abuse Prevention, the Center for Mental Health Services, the Center for Behavioral Health Statistics and Quality, and other SAMHSA partners, such as the Office of National Drug Control Policy. These products are available to the public for ordering and download. When a member of the public chooses to order hard-copy publications, it is necessary for SAMHSA to collect certain customer information in order to fulfill the request. To further lessen the burden on the public and provide the level of customer service that the public has come to expect from product Web sites, SAMHSA has developed a voluntary registration process for its publication Web site that allows customers to create accounts. Through these accounts, SAMHSA customers are able to access their order histories and save their shipping addresses. This reduces the burden on customers of having to re-identify materials they ordered in the past and to re-enter their shipping information each time they place an order with SAMHSA. During the Web site registration process, SAMHSA also asks customers to provide optional demographic information that helps SAMHSA evaluate the use and distribution of its publications and improve services to the public.

SAMHSA is employing a Web-based form for information collection to avoid duplication and unnecessary burden on customers who register both for an account on the product Web site and for email updates. The Web technology allows SAMHSA to integrate the email update subscription process into the Web site account registration process. Customers who register for an account on the product Web site are given the option of being enrolled automatically to receive SAMHSA email updates. Any optional questions answered by the customer during the Web site registration process automatically are mapped to the profile generated for the email update system, thereby reducing the collection of duplicate information.

SAMHSA collects all customer information submitted for Web site registration and email update subscriptions electronically via a series of Web forms on the samhsa.gov domain. Customers can submit the Web forms at their leisure, or call SAMHSA’s toll-free Call Center and an information specialist will submit the forms on their behalf. The electronic collection of information reduces the burden on the respondent and streamlines the data-capturing process. SAMHSA places Web site registration information into a Knowledge Management database and places email subscription information into a database maintained by a third-party vendor that serves multiple Federal agencies and the White House. Customers can change, add, or delete their information from either system at any time.

The respondents are behavioral health professionals, researchers, parents, caregivers, and the general public.

SAMHSA proposes two changes to the information collection. The first change is increasing the number of responses based on the average annual number of actual responses in 2011 and 2012. The second change is modifying the response options for “Organization Type” in the following ways: “Treatment Facility” will be changed to “Behavioral Health Treatment Facility”, “Individual/Group Practice” will be changed to “Other Health Care Facility”, and adding four new categories including “Military/Veterans Organization,” “Criminal Justice/Courts,” “Health Insurer,” and “Human Resources/Employer Assistance Program.”

SAMHSA estimates the burden of this information collection as follows:

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<tr>
<th>Table 1—Estimated Annual Reporting Burden</th>
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<tr>
<td>Number of respondents</td>
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<td>Web Site Registration</td>
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