amended (5 U.S.C. App.), notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable materials, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Institute of Mental Health Initial Review Group; Mental Health Services in Non-Specialty Settings.

Date: June 8–9, 2010.

Time: 8 a.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: Melrose Hotel, 2430 Pennsylvania Avenue, NW., Washington, DC 20037.

Contact Person: Aileen Schulte, PhD, Scientific Review Officer, Division of Extramural Activities, National Institute of Mental Health, NIH, Neuroscience Center, 6001 Executive Blvd., Room 6154, MSC 9606, Bethesda, MD 20892–9606, 301–443–1225, aschulte@mail.nih.gov.

Name of Committee: National Institute of Mental Health Initial Review Group; Interventions Committee for Adult Disorders.

Date: June 8–9, 2010.

Time: 8:30 a.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: Hyatt Regency Bethesda, One Bethesda Metro Center, 7400 Wisconsin Avenue, Bethesda, MD 20814.

Contact Person: David Sommers, PhD, Scientific Review Officer, Division of Extramural Activities, National Institute of Mental Health, NIH, Neuroscience Center, 6001 Executive Blvd., Room 6154, MSC 9606, Bethesda, MD 20892–9606, 301–443–7861, dsommers@mail.nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.242, Mental Health Research Grants; 93.281, Scientist Development Award, Scientific Development Award for Clinicians, and Research Scientist Award; 93.282, Mental Health National Research Service Awards for Research Training, National Institutes of Health, HHS)


Jennifer Spaeth,
Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2010–2846 Filed 4–27–10; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meetings. The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Bioengineering Sciences & Technologies Integrated Review Group; Biomaterials and Biointerfaces Study Section.

Date: May 25–26, 2010.

Time: 8 a.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: Hilton Crystal City, 2399 Jefferson Davis Highway, Arlington, VA 22202.

Contact Person: Steven J Zullo, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5146, MSC 7849, Bethesda, MD 20892, 301–435–2810, zullost@csr.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict: Gastrointestinal Physiology and Pathophysiology.

Date: May 28, 2010.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0001]

Food and Drug Administration/Xavier University Global Outsourcing Conference

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public conference.

SUMMARY: The Food and Drug Administration (FDA), Cincinnati District, in co-sponsorship with Xavier University, is announcing a public conference entitled “FDA/Xavier University Global Outsourcing Conference.” This 3-day public conference for the pharmaceutical industry includes presentations from key FDA officials, global regulators, and industry experts. The conference will focus on global compliance challenges associated with pharmaceutical outsourcing relationships and supply chain control, as well as expectations from global regulators. Pharmaceutical companies and contract organizations are invited to this conference to address the issues that reside on both sides. In addition to expert presentations, participants will be engaged through live polling and a small group discussion session on sharing best practices with each other.

Dates and Times: The public conference will be held on June 14 and 15, 2010, from 8 a.m. to 5 p.m. and June 16, 2010, from 8 a.m. to 1 p.m.

Location: The public conference will be held on the campus of Xavier University.
University, 3800 Victory Pkwy.,
Cincinnati, OH 45207, 513–745–3073 or
513–745–3396.

Contact Persons:
For information regarding this notice:
Steven Eastham, Food and Drug
Administration, 6751 Steger Dr.,
Cincinnati, OH 45237, 513–679–2700,
ext. 123, e-mail:
steven.eastham@fda.hhs.gov.

For information regarding the
conference and registration: Marla
Phillips, Xavier University, 3800
Victory Pkwy., Cincinnati, OH 45207,
513–745–3073, e-mail:
phillipsm@xavier.edu.

Registration: There is a registration
fee. The conference registration fees
cover the cost of the presentations,
training materials, receptions,
breakfasts, lunches, dinners, and dinner
speakers for the 3 days of the
conference. Early registration ends May
14, 2010. Standard registration ends
June 13, 2010. There will be onsite
registration. The cost of registration is as
follows:

<table>
<thead>
<tr>
<th>Attendees</th>
<th>Fees by May 14th</th>
<th>Fees by June 13th</th>
</tr>
</thead>
<tbody>
<tr>
<td>Industry</td>
<td>$995</td>
<td>$1,200</td>
</tr>
<tr>
<td>Small Business (&lt; 100 employees)</td>
<td>$800</td>
<td>$1,000</td>
</tr>
<tr>
<td>Academic/Government</td>
<td>$600</td>
<td>$700</td>
</tr>
<tr>
<td>Student</td>
<td>$200</td>
<td>$250</td>
</tr>
<tr>
<td>FDA Employee</td>
<td>Fee waived</td>
<td>Fee waived</td>
</tr>
</tbody>
</table>

† The fourth registration from the same company is free.

The following forms of payment will
be accepted: American Express, Visa,
Mastercard, and company checks.

To register online for the public
conference, please visit the
“Registration” link on the conference
(FDA has verified the Web site address,
but is not responsible for subsequent
changes to the Web site after this
document publishes in the Federal
Register.)

To register by mail, please send your
name, title, firm name, address,
telephone and fax numbers, e-mail, and
payment information to the fee to
Xavier University, Attn: Sue Bensman,
3800 Victory Pkwy., Cincinnati, OH
45207. An e-mail will be sent
confirming your registration.

Attendees are responsible for their
own accommodations. The conference
headquarter hotel is the Downtown
Cincinnati Hilton Netherlands Plaza, 35
West 5th Street, Cincinnati, OH 45202,
513–421–9100. To make reservations
online, please visit the "Venue/

If you need special accommodations
due to a disability, please contact Marla
Phillips (see Contact Persons) at least 7
days in advance of the conference.

SUPPLEMENTARY INFORMATION: The
public conference helps fulfill the
Department of Health and Human
Services and FDA’s important mission
to protect the public health. The
conference will provide those engaged
in FDA-regulated outsourcing with
information on the following topics:

- FDA Center to present on initiatives
  from Congress and FDA, and resulting
  impact;
- Global regulator perspective on
  global compliance initiatives,
  challenges, and expectations;
- FDA Field perspective on the most
  common and significant deficiencies
  specific to outsourcing relationships;
- Global compliance of
  manufacturing in Asia;
- Pharmaceutical companies—how to
  manage varying global regulatory
  expectations while working with
  contractors in various states of
  compliance;
- Contract organizations—
  compliance strategy for managing
  global regulatory requirements while
  managing multiple client expectations;
- Contract Organization Selection
  Process;
- The Client Selection Process—the
  criteria a contract organization should
  use to consider saying no to a contract
  relationship;
- Regulatory challenges—Drug Master
  File Fitness;
- Due diligence audit—how to audit
  in 1 day;
- Quality Agreement Development
  throughout the product and process
  lifecycle;
- Supply Chain Transparency and
  Pedigree;
- How to Audit the Supply Chain;
- Rx-360 and International
  Pharmaceutical Excipients Council
  initiatives—impact to industry;
- Risk-based Performance
  Management best practices;
- International Conference on
  Harmonisation Triple Q’s (Q8, Q9, and
  Q10)—how quality can drive down the
cost of business, and how innovation
can increase business opportunities;
- Rebuilding the Trust case studies;
and
- Small group discussion on sharing
  best practices.

FDA has made education of the drug
and device manufacturing community a
high priority to help ensure the quality
of FDA-regulated drugs and devices.
The conference helps to achieve
objectives set forth in section 406 of the
Food and Drug Administration
393), which includes working closely
with stakeholders and maximizing the
availability and clarity of information to
stakeholders and the public. The
conference also is consistent with the
Small Business Regulatory Enforcement
Fairness Act of 1996 (Public Law 104–
121) by providing outreach activities by
Government agencies to small
businesses.


Leslie Kux,
Acting Assistant Commissioner for Policy.
[FR Doc. 2010–9795 Filed 4–27–10; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND
HUMAN SERVICES

Centers for Disease Control and
Prevention

Statement of Organization, Functions,
and Delegations of Authority

Part C (Centers for Disease Control
and Prevention) of the Statement of