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 “Register Now” link on the conference Web site at http://www.XavierGOC.com. 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, email, and payment information for the fee to Xavier University, Attention: Sue Bensman, 3800 Victory Pkwy., Cincinnati, OH 45207. An email will be sent confirming your registration. Attendees are responsible for their own accommodations. The conference headquarters hotel is the Downtown Cincinnati Hilton Netherlands Plaza, 35 West Fifth St., Cincinnati, OH 45202, 513–421–9100. To make reservations online, please visit the “Venue & Logistics” link at http://www.XavierGOC.com to make reservations. The hotel is expected to sell out during this timeframe, so early reservation in the conference room-block is encouraged.

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 International Initiatives
- European Union Regulator Perspective

### Table 1—Registration Fees 1

<table>
<thead>
<tr>
<th>Attendee</th>
<th>Fee on or before August 5th</th>
<th>Fee August 6th–September 2nd</th>
<th>Fee September 3rd–September 23rd</th>
</tr>
</thead>
<tbody>
<tr>
<td>Industry</td>
<td>$995</td>
<td>$1,295</td>
<td>$1,495</td>
</tr>
<tr>
<td>Small Business (&lt;100 employees)</td>
<td>800</td>
<td>900</td>
<td>1,000</td>
</tr>
<tr>
<td>Consultants</td>
<td>500</td>
<td>600</td>
<td>700</td>
</tr>
<tr>
<td>Startup Manufacturers/Academic</td>
<td>200</td>
<td>250</td>
<td>300</td>
</tr>
<tr>
<td>Media/Government</td>
<td>Free</td>
<td>Free</td>
<td>Free</td>
</tr>
</tbody>
</table>

1 The fourth registration from the same company is free.
The Food and Drug Administration (FDA) is announcing the following public meeting entitled “Regulatory Science Considerations for Medical Countermeasure (MCM) Radiation Biodosimetry Devices.” The purpose of the public meeting is to obtain input from academia, Government, industry, and other stakeholders on the clinical application and scientific and technological challenges for performance validation of radiation biodosimetry devices.

**Date and Time:** The public meeting will be held on September 27 and 28, 2012, from 8:30 a.m. to 5 p.m.

**Location:** The public meeting will be held at FDA’s White Oak Campus, 10903 New Hampshire Ave., Bldg. 31 Conference Center, the Great Room (rm. 1503), Silver Spring, MD 20993–0002.

Entrance for the public meeting participants (non-FDA employees) is through Bldg. 1 where routine security check procedures will be performed. For parking and security information, please visit the following Web site: http://www.fda.gov/AboutFDA/WorkingatFDA/BuildingsandFacilities/WhiteOakCampus/Information/ucm241740.htm. The public meeting will also be webcast.

**Contact:** Jennifer S. Dickey, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, rm. 4254, Silver Spring, MD 20993–0002, 301–796–5028, Fax: 301–847–8512, email: Jennifer.Dickey@fda.hhs.gov.

**Registration:** Registration is free and will be on a first-come, first-served basis. Persons interested in attending this public meeting must register online by 4 p.m., September 13, 2012. Early registration is recommended because facilities are limited and, therefore, FDA may limit the number of participants from each organization. If time and space permits, onsite registration on the day of the public meeting will be provided beginning at 7 a.m.

If you need special accommodations due to a disability, please contact Susan Monahan, Center for Devices and Radiological Health, Food and Drug Administration, 10003 New Hampshire Ave., Bldg. 66, rm. 4321, Silver Spring, MD 20993–0002, 301–796–5661, email: Susan.Monahan@fda.hhs.gov at least 7 days in advance of the meeting.

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To register for the public meeting, please visit FDA’s Medical Devices News & Events—Workshops & Conferences calendar at http://www.fda.gov/MedicalDevices/NewsEvents/WorkshopsConferences/default.htm. (Select this public meeting from the posted events list.) Please provide complete contact information for each participant, including name, title, affiliation, email, and telephone number. Those without Internet access should contact Susan Monahan to register (see previous paragraph). Registrants will receive confirmation after they have been accepted. You will be notified if you are on a waiting list.

**Streaming Webcast of the Public Meeting:** This public meeting will also be webcast. Persons interested in viewing the webcast must register online by 4 p.m., September 13, 2012. Early registration is recommended because webcast connections are limited. Organizations are requested to register all participants, but to view using one connection per location. Webcast participants will be sent technical system requirements after registration and will be sent connection access information after September 20, 2012. If you have never attended a Connect Pro meeting before, test your connection at: https://collaboration.fda.gov/common/help/en/support/meeting_test.htm. To get a quick overview of the Connect Pro program, visit: http://www.adobe.com/go/connectpro_overview. FDA has verified the Web site addresses in this document, but FDA is not responsible for any subsequent changes to the Web sites after this document publishes in the Federal Register.

**Requests for Oral Presentations:** This public meeting includes public comment sessions. During online registration you may indicate if you wish to present during a public comment session or participate in a specific session, and which topics you wish to address. FDA has included general topics in this document. FDA will do its best to accommodate requests to make public comment. Individuals and organizations with common interests are urged to consolidate or coordinate their presentations, and request time for a joint presentation. Following the close of registration, FDA will determine the amount of time allotted to each presenter and the approximate time that each oral presentation is to begin, and will select and notify participants by September 18, 2012. All requests to make oral presentations must be received by the close of registration on September 13, 2012 by 4 p.m. If selected for presentation, any presentation materials must be emailed to Jennifer Dickey (see Contact) no later than September 24, 2012. No commercial or promotional material will be permitted to be presented or distributed at the meeting.

**Comments:** FDA is holding this public meeting to obtain information on the clinical application and scientific and technological challenges for performance validation of radiation biodosimetry devices. In order to permit