COST OF REGISTRATION—Continued

Affiliation	Fee
Non-Government (SoCRA Member)	$575.00
Non-Government (Non SoCRA Member)	$650.00

If you need special accommodations due to a disability, please contact Marie Falcone (see *Contact*) at least 7 days in advance of the workshop.

Registration Instructions: To register, please submit a registration form with your name, affiliation, mailing address, telephone, fax number, and e-mail address, along with a check or money order payable to ''Socra.'' Mail to: SoCRA, 530 West Butler Ave., suite 109, Chalfont, PA 18914. To register via the Internet, go to *http://www.socra.org/html/FDA_Conference.htm*. 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**).The registrar will also accept payment by major credit cards (VISA/MasterCard/AMEX only). For more information on the public workshop, or for questions on registration, contact the Society of Clinical Research Associates at 800–762–7292 or 215–822–8644, FAX: 215–822–8633, or e-mail: *SoCRAmail@aol.com*.

SUPPLEMENTARY INFORMATION: The public workshop helps fulfill the Department of Health and Human Services' and FDA's important mission to protect the public health. The workshop will provide those engaged in FDA-regulated (human) clinical trials with information on a number of topics concerning FDA requirements related to informed consent, clinical investigation requirements, institutional review board (IRB) inspections, electronic record requirements, and investigator initiated research. Topics for discussion include the following:

- What FDA Expects in a Pharmaceutical Clinical Trial;
- Adverse Event Reporting—Science, Regulation, Error, and Safety;
- Part 11 Compliance—Electronic Signatures;
- Informed Consent Regulations;
- IRB Regulations and FDA Inspections;
- Keeping Informed and Working Together;
- FDA Conduct of Clinical Investigator Inspections;
- Meetings With FDA: Why, When, and How;
- Investigator Initiated Research;
- Medical Device Aspects of Clinical Research;
- Working With FDA's Center for Biologics Evaluation and Research; and
- The Inspection is Over—What Happens Next? Possible FDA Compliance Actions.

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 public workshop helps to achieve objectives set forth in section 406 of the FDA Modernization Act of 1997 (21 U.S.C. 393) which includes working closely with stakeholders and maximizing the availability and clarity of information to stakeholders and the public. The public workshop also is consistent with the Small Business Regulatory Enforcement Fairness Act of 1996 (Public Law 104–121), as outreach activities by Government agencies to small businesses.

Dated: August 18, 2009.

David Horowitz,

Assistant Commissioner for Policy.

[FR Doc. E9–20340 Filed 8–25–09; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, National Center for Public Health Informatics (BSC, NCPHI)

Correction: The notice was published in the **Federal Register** on August 18, 2009 [Volume 74, Number 158] [page 41712]. The ''Matters To Be Discussed'' has been revised: The board will discuss public health informatics issues related to the H1N1 virus; CDC public health informatics strategies and goals, including future program activities; and how the board can provide informatics scientific input to CDC.

Contact Person for More Information: Dr. Scott McNabb, National Center for Public Health Informatics, CDC, 1600 Clifton Road, NE., Mailstop E–78, Atlanta, Georgia 30333, Telephone (404) 498–6427, Fax (404) 498–6235.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities for both the CDC and the Agency for Toxic Substance and Disease Registry.

Dated: August 18, 2009.

Elaine L. Baker,

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

[FR Doc. E9–20575 Filed 8–25–09; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Center for Complementary and Alternative Medicine Announcement of Wellness Workshop

ACTION: Notice.

SUMMARY: The National Center for Complementary and Alternative Medicine (NCCAM) invites the research community to participate in a workshop focused on wellness.

The purpose of this workshop is to review several measures of wellness, identify their strengths and weaknesses, and make recommendations on how best to capture the construct. This information will help NCCAM guide development of questions for the 2012 National Health Interview Survey.

The Workshop will take place on September 25, 2009. Those interested in CAM research are particularly encouraged to attend.

Background: The National Center for Complementary and Alternative Medicine (NCCAM) was established in 1999 with the mission of exploring complementary and alternative healing practices in the context of rigorous science, training CAM researchers, and disseminating authoritative information to the public and professionals. NCCAM funds research grants that explore the science of CAM. For more information, see *http://nccam.nih.gov/grants/whatnccamfunds/*.

Participating: The public is invited to attend and observe this workshop. Those interested in attending are required to RSVP via e-mail to Edward Culhane Jr. at *culhane@mail.nih.gov* with their name, affiliation, e-mail and phone number. Space constraints limit the number of attendees at this workshop and participation will be on a first come, first served basis. For more information about what will be covered at the workshop, *see http://nccam.nih.gov/news/events/*.

FOR FURTHER INFORMATION CONTACT: To request more information, visit the NCCAM Web site at *http://nccam.nih.gov/news/events/*, call 301–594–3391 (Edward Culhane Jr.) or e-mail at *culhanee@mail.nih.gov*.