vitae, and a brief summary of relevant experience and qualifications.

#### Request for Data

NICEATM invites the submission of data for substances tested in standardized in vivo acute dermal systemic toxicity tests. Corresponding acute oral LD50 data for the same compounds tested dermally would be particularly useful. Oral data from rat tests and dermal data from rat and/or rabbit tests are preferred. Although data can be accepted at any time, please submit data by September 6, 2012 to ensure consideration during the ICCVAM evaluation process. Relevant data received after this date will be considered where feasible. All information submitted in response to this notice will be made publicly available and may be incorporated into future NICEATM and ICCVAM reports and publications, as appropriate.

When submitting data, please reference this Federal Register notice and provide appropriate contact information (name, affiliation, mailing address, phone, fax, email, and sponsoring organization, as applicable). NICEATM prefers that data be submitted electronically as copies of pages from study notebooks, spreadsheets, and/or study reports. Each submission for a substance should preferably include the following information, as appropriate: common and trade name, Chemical Abstracts Service Registry Number (CASRN), commercial source, in vivo test protocols used, extent to which the data were collected in accordance with national or international Good Laboratory Practice guidelines, date and testing organization, physical and chemical properties (e.g., molecular weight, pH, water solubility, log Kow, etc.), estimated LD50, and incidence of death and other adverse effects.

### Background Information on ICCVAM and NICEATM

ICCVAM is an interagency committee composed of representatives from 15 Federal regulatory and research agencies that require, use, generate, or disseminate toxicological and safety testing information. ICCVAM conducts technical evaluations of new, revised, and alternative safety testing methods and integrated testing strategies with regulatory applicability and promotes the scientific validation and regulatory acceptance of testing methods that more accurately assess the safety and hazards of chemicals and products and that reduce, refine (enhance animal wellbeing and lessen or avoid pain and distress), or replace animal use. The

ICCVAM Authorization Act of 2000 (42 U.S.C. 2851-3) established ICCVAM as a permanent interagency committee of the NIEHS under NICEATM. NICEATM administers ICCVAM, provides scientific and operational support for ICCVAM-related activities, and conducts independent validation studies to assess the usefulness and limitations of new, revised, and alternative test methods and strategies. NICEATM and ICCVAM welcome the public nomination of new, revised, and alternative test methods and strategies for validation studies and technical evaluations. Additional information about NICEATM and ICCVAM can be found on the NICEATM-ICCVAM Web site (http://iccvam.niehs.nih.gov).

#### References

Bronstein AC, Spyker DA, Cantilena LR, Jr, Green JL, Rumack BH, Giffin SL. 2010. 2009 Annual Report of the American Association of Poison Control Centers' National Poison Data System (NPDS): 27th Annual Report. Clinical Toxicology 48: 979–1178.

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ICCVAM. 2008. NICEATM—ICCVAM Five-Year Plan (2008—2012): A Plan to Advance Alternative Test Methods of High Scientific Quality to Protect and Advance the Health of People, Animals and the Environment. Research Triangle Park, NC: National Institute of Environmental Health Sciences. Available: http://iccvam.niehs.nih.gov/ docs/5yearplan.htm.

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OECD. 2001. Test No. 425. Acute Oral Toxicity—Up-and-Down Procedure [adopted 17 December 2001]. In: OECD Guidelines for the Testing of Chemicals, Section 4: Health Effects. Paris:OECD Publishing. Available: www.oecd.org/ dataoecd/17/51/1948378.pdf.

Dated: July 12, 2012.

#### John R. Bucher,

Associate Director, National Toxicology Program.

[FR Doc. 2012–17787 Filed 7–20–12; 8:45 am]

### DEPARTMENT OF HEALTH AND HUMAN SERVICES

# **Centers for Disease Control and Prevention**

# Safety and Occupational Health Study Section: Notice of Charter Renewal

This gives notice under the Federal Advisory Committee Act (Pub. L. 92–463) of October 6, 1972, that the Safety and Occupational Health Study Section, Centers for Disease Control and Prevention, Department of Health and Human Services, has been renewed for a 2-year period through June 30, 2014.

For more information contact: Price Connor, Ph.D., Executive Secretary, Safety and Occupational Health Study Section, Department of Health and Human Services, 1600 Clifton Road NE., Mailstop E74, Atlanta, Georgia 30333, telephone 404/498–2511 or fax 404/498–2571.

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 Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dated: July 16, 2012.

#### Elaine L. Baker,

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

[FR Doc. 2012–17879 Filed 7–20–12; 8:45 am] BILLING CODE 4163–18–P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

#### Centers for Disease Control and Prevention

#### Advisory Board on Radiation and Worker Health (ABRWH or Advisory Board), National Institute for Occupational Safety and Health (NIOSH); Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following meeting of the aforementioned committee:

**TIME AND DATE:** 11:00 a.m.-3:00 p.m., August 15, 2012.

**PLACE:** Audio Conference Call via FTS Conferencing. The USA toll-free, dial-in number is 1–866–659–0537 and the pass code is 9933701.

**STATUS:** Open to the public, but without a verbal public comment period.