Workshop on Adverse Outcome Pathways: From Research to Regulation

Abstract Submission Guidelines

The steering committee of "Adverse Outcome Pathways: From Research to Regulation" invites the submission of abstracts for scientific posters. Posters should address how adverse outcome pathways may be developed or used (i) to predict toxic effects in humans or ecological systems or (ii) to improve regulatory assessment of chemical toxicity. Posters that address related issues or topics may be considered as space allows.

Abstracts should be submitted by email to Michael Paris (paris@niehs.nih.gov) by close of business on **July 25, 2014**. The workshop steering committee will review the submitted abstracts. The corresponding author will be notified of an abstract's acceptance and provided with guidelines for poster presentation no later than August 4, 2014.

Requirements for Submission:

- The abstract should be submitted as a Microsoft Word document with one-inch margins and text in 12 point font or larger.
- Heading information should include the names of all authors and their affiliations.
- Abstracts must contain a statement of the rationale and scope of the study presented, a brief description of procedures used, data collected, and principal conclusion(s) based on interpretation of the results.
- Abstract body may not exceed 300 words.
- Key references may be included after the abstract body.
- Complete abstract including heading, body, and references may not exceed one page.
- Name and contact information (physical address, phone number, fax number, and e-mail address) for the corresponding or senior author should be provided at the end of the abstract.

Additional Requirements:

- All abstracts describing studies using animals or animal tissues should include a statement that all animal use was carried out in accordance with all applicable animal care and use laws, regulations, and guidelines and that the study was approved by the appropriate Institutional Animal Care and Use Committee.
- All abstracts describing human studies should include a statement that such studies were conducted in accordance with all applicable laws, regulations, and guidelines and that the appropriate Institutional Review Board approved the study.