

Data Management (CDM), NIEHS, P.O. Box 12233, MD E1-02, Research Triangle Park, NC 27709, T: 919-541-3419, FAX: 919-541-3687, or e-mail: CDM@niehs.nih.gov.

The NTP Board of Scientific Counselors Technical Reports Review Subcommittee meeting is open to the public and public comment on any of the Technical Reports is welcome. Time will be provided at the meeting for public comment on each of the reports under review. In order to facilitate planning for the meeting, persons requesting time for an oral presentation on a particular Technical Report are asked to notify the Executive Secretary, Dr. Mary S. Wolfe, at P.O. Box 12233, MD A3-07, Research Triangle Park, NC 27709, T: 919-541-3971, F: 919-541-0295, e-mail: wolfe@niehs.nih.gov. Persons registering to make brief comments are asked to provide, if possible, a written copy of their statement by April 20, to enable review by the Subcommittee and staff prior to the meeting. Written statements can

supplement and may expand the oral presentation. Each speaker is asked to provide his/her name, affiliation, mailing address, phone, fax, e-mail and supporting organization (if any). At least seven minutes will be allotted to each speaker, and if time permits, may be extended to 10 minutes. Each organization is allowed one time slot for each report being reviewed. Registration for making public comments will also be available on-site. If registering on-site to speak and reading oral comments from printed copy, the speaker is asked to bring 25 copies of the text. These copies will be distributed to the Chair and Subcommittee members and supplement the record.

Written comments, in lieu of making oral comments, are also welcome. The comments should include name, affiliation, mailing address, phone, fax, e-mail and sponsoring organization (if any) and preferably be received by April 20, to enable review by the Subcommittee and staff prior to the meeting.

Request for Additional Information

The NTP would welcome receiving toxicology and carcinogenesis information from completed, ongoing or planned studies as well as current production data, human exposure information, and use patterns for any of the chemicals listed in this announcement. Please forward this information to CDM at the address given above. CDM will forward the information to the appropriate staff scientist.

The agenda and a roster of subcommittee members will be available prior to the meeting on the NTP web homepage at <http://ntp-server.niehs.nih.gov> and upon request from the Executive Secretary. Following the meeting, summary minutes will be available on the NTP web homepage and upon request to Dr. Wolfe.

Dated: March 23, 2001.

Samuel H. Wilson,
Deputy Director, National Institute of Environmental Health Sciences.

TECHNICAL REPORTS TENTATIVELY SCHEDULED FOR REVIEW BY THE NTP BOARD OF SCIENTIFIC COUNSELORS TECHNICAL REPORTS REVIEW SUBCOMMITTEE ON MAY 3, 2001

Chemical CAS number	Report No.	Primary uses	Route & exposure levels	Review order
Acrylonitrile 107-13-1 ..	TR-506	Used in the production of acrylic fibers, elastomers, resins, and a variety of chemical or intermediates; annual production is in the millions of tons.	Gavage (deionized water vehicle) Mice: 2.5, 10, or 20 mg/kg	3
Citral 5392-40-5	TR-505	Used in lemon flavoring in foods and beverages and as a lemon fragrance in detergents, perfumes, and toiletries.	Microencapsulated citral in feed Rats: 0, 1000, 2000, or 4000 ppm. Mice: 0, 500, 1000, or 2000 ppm	5
Methacrylonitrile 126-98-7.	TR-497	Used in the production of polymers, elastomers, and plastics including those used in beverage containers.	Gavage (deionized water vehicle) Rats: 0, 3, 10, or 30 mg/kg	4
o-Nitrotoluene 88-72-2	TR-504	Used in synthesis of agricultural and rubber chemicals and of a variety of dyes.	Mice: 0, 1.5, 3, or 6 mg/kg	1
p-Nitrotoluene 99-99-0	TR-498	Used in synthesis of agricultural and rubber chemicals and of a variety of dyes.	Feed Rats: 0, 625, 1250, or 2000 ppm; Male rats: 2000 or 5000 ppm (stop study) Mice: 0, 1250, 2500, or 5000 ppm	2
			Feed Rats & Mice: 0, 1250, 2500, or 5000 ppm	

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Agency Information Collection Activities: Proposed Collection; Comment Request

In compliance with section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 concerning opportunity for public comment on

proposed collections of information, the Substance Abuse and Mental Health Services Administration will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the information collection plans, call the SAMHSA Reports Clearance Officer on (301) 443-7978.

Comments are invited on: (a) Whether the proposed collections of information are necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection

of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology.

Proposed Project

Participant Feedback on Training Under the Cooperative Agreement for Mental Health Care Provider Education in HIV/AIDS Program II (OMB No. 0930-0195, Extension)—The Substance Abuse and Mental Health Services Administration's (SAMHSA) Center for

Mental Health Services (CMHS) intends to continue to conduct a multi-site assessment of its Cooperative Agreement for Mental Health Care Provider Education in HIV/AIDS Program II until the end of the sites' expenditure of Program II funds (anticipated end date of September 2002). The education programs funded under this cooperative agreement are designed to disseminate knowledge of the psychological and neuropsychiatric sequelae of HIV/AIDS to both traditional (e.g., psychiatrists, psychologists, nurses, primary care physicians, medical students, and social workers) and non-traditional (e.g., clergy, and alternative health care workers) first-line providers of mental health services.

The multi-site assessment is designed to assess the effectiveness of particular training curricula, document the

integrity of training delivery formats, and assess the effectiveness of the various training delivery formats. Analyses will assist CMHS in documenting the numbers and types of traditional and non-traditional mental health providers accessing training; the content, nature and types of training participants receive; and the extent to which trainees experience knowledge, skill and attitude gains/changes as a result of training attendance. The multi-site data collection design uses a two-tiered data collection and analytic strategy to collect information on (1) the organization and delivery of training, and (2) the impact of training on participants' knowledge, skills and abilities.

Information about the organization and delivery of training will be collected from trainers and staff who are

funded by these cooperative agreements hence there is no respondent burden. All training participants attending sessions lasting less than 6 hours will be asked to complete a brief feedback form at the end of the training session. Trainees attending sessions lasting 6 hours or longer will be asked to complete brief pre- and post-session feedback questionnaires. A sample of trainees attending sessions lasting 6 hours or longer will also be asked to complete a brief follow-up telephone interview three months after the training session. CMHS has funded seven education sites under the Cooperative Agreement for Mental Health Care Provider Education in HIV/AIDS Program II. The annual burden estimates for this activity are shown below:

Form	Responses per respondent	Estimated number of respondents (× 7 sites)	Hours per response	Total hours
All Sessions				
Session Report Form	1	60 × 7 = 420	0.080	34
Sessions Less than 6 Hours				
Participant Feedback Form	1	600 × 7 = 4200	0.167	701
Neuropsychiatric Participant Feedback Form	1	75 × 7 = 525	0.167	88
Ethics Participant Feedback Form	1	75 × 7 = 525	0.167	88
Sessions 6 Hours or Longer				
Pre-Training Participant Inventory	1	200 × 7 = 1400	0.167	234
Post-Training Participant Inventory	1	200 × 7 = 1400	0.250	350
Neuropsychiatric Pre-Training Participant Inventory	1	50 × 7 = 350	0.167	58
Neuropsychiatric Post-Training Participant Inventory	1	50 × 7 = 350	0.250	88
Participant Follow-up Form	1	45 × 7 = 315	0.250	79
Monthly Form Submission				
Monthly Form Mailing	¹ 12	84	0.167	14
Total		7,504		1,733

¹ Per site.

Send comments to Nancy Pearce, SAMHSA Reports Clearance Officer, Room 16-105, Parklawn Building, 5600 Fishers Lane, Rockville, MD 20857. Written comments should be received within 60 days of this notice.

Dated: March 27, 2001.

Richard Kopanda,

Executive Officer, SAMHSA.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Current List of Laboratories Which Meet Minimum Standards To Engage in Urine Drug Testing for Federal Agencies

AGENCY: Substance Abuse and Mental Health Services Administration, HHS.

ACTION: Notice.

SUMMARY: The Department of Health and Human Services notifies Federal agencies of the laboratories currently certified to meet standards of Subpart C

of Mandatory Guidelines for Federal Workplace Drug Testing Programs (59 FR 29916, 29925). A similar notice listing all currently certified laboratories will be published during the first week of each month, and updated to include laboratories which subsequently apply for and complete the certification process. If any listed laboratory's certification is totally suspended or revoked, the laboratory will be omitted from updated lists until such time as it is restored to full certification under the Guidelines.

If any laboratory has withdrawn from the National Laboratory Certification Program during the past month, it will