

ENVIRONMENTAL PROTECTION AGENCY**[FRL-9640-9]****Request for Public Comments on the List of Candidates for EPA's Science Advisory Board (SAB), Chemical Assessment Advisory Committee****AGENCY:** Environmental Protection Agency.**ACTION:** Notice.

SUMMARY: The U.S. Environmental Protection Agency (EPA) invites public comments on the list of candidates being considered for appointment to the EPA's Science Advisory Board (SAB) Chemical Assessment Advisory Committee (CAAC).

DATES: Comments should be submitted in time to arrive no later than March 21, 2012.

FOR FURTHER INFORMATION CONTACT:

Members of the public wishing to obtain further information may contact the Dr. Suhair Shallal, the Designated Federal Officer (DFO) for the committee, by email at shallal.suhair@epa.gov or by telephone at 202-564-2057.

Background: The U.S. Environmental Protection Agency (EPA) Science Advisory Board (SAB) Staff Office announced in a **Federal Register** Notice (Vol. 76, No. 223, pp. 71561-62) published on November 18, 2011 that it was forming a new committee under the auspices of the SAB to provide advice to EPA through the chartered SAB regarding the development of IRIS Toxicological Reviews available on EPA's Integrated Risk Assessment System (IRIS). The SAB Staff Office sought public nominations of nationally and internationally recognized experts with knowledge in human health risk assessment and expertise in a range of disciplines including, but not limited to: *public health; epidemiology; toxicology; modeling; biostatistics; and risk assessment*. The SAB Staff Office hereby invites public comments on the list of candidates under consideration for the SAB Chemical Assessment Advisory Committee.

How To Submit Comments: Any interested person or organization may submit comments to Dr. Suhair Shallal, Designated Federal Officer, no later than March 21, 2012. Emailing comments (shallal.suhair@epa.gov) is the preferred mode of receipt. Please be advised that public comments are subject to release under the Freedom of Information Act.

Dated: February 22, 2012.

Vanessa T. Vu,

Director, EPA Science Advisory Board Staff Office.

[FR Doc. 2012-4841 Filed 2-28-12; 8:45 am]

BILLING CODE 6560-50-P**FEDERAL COMMUNICATIONS COMMISSION****Information Collection Being Reviewed by the Federal Communications Commission****AGENCY:** Federal Communications Commission.**ACTION:** Notice and request for comments.

SUMMARY: As part of its continuing effort to reduce paperwork burden and as required by the Paperwork Reduction Act (PRA) of 1995 (44 U.S.C. 3501-3520), the Federal Communications Commission invites the general public and other Federal agencies to take this opportunity to comment on the following information collection(s). Comments are requested concerning: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the Commission, including whether the information shall have practical utility; (b) the accuracy of the Commission's burden estimate; (c) ways to enhance the quality, utility, and clarity of the information collected; (d) ways to minimize the burden of the collection of information on the respondents, including the use of automated collection techniques or other forms of information technology; and (e) ways to further reduce the information burden for small business concerns with fewer than 25 employees.

The FCC may not conduct or sponsor a collection of information unless it displays a currently valid OMB control number. No person shall be subject to any penalty for failing to comply with a collection of information subject to the Paperwork Reduction Act (PRA) that does not display a valid OMB control number.

DATES: Written Paperwork Reduction Act (PRA) comments should be submitted on or before April 30, 2012. If you anticipate that you will be submitting PRA comments, but find it difficult to do so within the period of time allowed by this notice, you should advise the FCC contact listed below as soon as possible.

ADDRESSES: Submit your PRA comments to Judith B. Herman, Federal Communications Commission, via the

Internet at Judith-b.herman@fcc.gov. To submit your PRA comments by email send them to: PRA@fcc.gov.

FOR FURTHER INFORMATION CONTACT:

Judith B. Herman, Office of Managing Director, (202) 418-0214.

SUPPLEMENTARY INFORMATION:

OMB Control Number: 3060-0936.

Title: Section 95.1215, Medical Device Radiocommunication Service (MedRadio); Disclosure Policies and Section 95.1217, Labeling Requirements.

Form Number: N/A.

Type of Review: Revision of a currently approved collection.

Respondents: Business or other for-profit entities and Not-for-profit institutions.

Number of Respondents: 100 respondents; 100 responses.

Estimated Time per Response: 1 hour for each manufacturer (20 manufacturers).

Frequency of Response: On occasion reporting requirement and third party disclosure requirement.

Obligation to Respond: Required to obtain or retain benefits. Statutory authority for this information collection is contained in 47 U.S.C. Sections 151, and 303 of the Communications Act of 1934, as amended.

Total Annual Burden: 100 hours.

Total Annual Cost: N/A.

Privacy Impact Assessment: N/A.

Needs and Uses: The Commission will submit this information collection to the Office of Management and Budget (OMB) for approval of a revision. The Commission adopted and released a Report and Order, FCC 11-176, Amendment of Parts 2 and 95 of the Commission's rules to provide additional spectrum for the Medical Device Radiocommunication Service which requires manufacturers of MedRadio programmer/control transmitters shall include the following statement on the device(s) in conspicuous location, or if it is not feasible to place the statement on the device, in the instruction manual.

"This device may not interfere with stations operating in the 400.150-406.000 MHz band in the Meteorological Aids, Meteorological Satellite, and Earth Exploration Satellite Services and must accept any interference received, including interference that may cause undesired operation."

Also section 95.1215(b) was revised as follows for which the Commission also seeks OMB approval:

Manufacturers of MedRadio transmitters operating in the 413-419 MHz, 426-432 MHz, 438-444 MHz, and 451-457 MHz bands must include with