6712-01

FEDERAL COMMUNICATIONS COMMISSION

Information Collection Being Reviewed by the Federal Communications Commission

AGENCY: Federal Communications Commission.

ACTION: Notice; 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: 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; the accuracy of the Commission's burden estimate; ways to enhance the quality, utility, and clarity of the information collected; 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 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 **[INSERT DATE 60 DAYS AFTER DATE OF PUBLICATION IN THE FEDERAL**

REGISTER]. 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 Benish Shah, Federal Communications Commission, via the Internet at Benish.shah@fcc.gov. To submit your PRA comments by email send them to: PRA@fcc.gov.

FOR FURTHER INFORMATION CONTACT: Benish Shah, Office of Managing Director, (202) 418-7866.

SUPPLEMENTARY INFORMATION:

OMB Control No.: 3060-0936.

Title: Sections 95.1215, 95.1217, 95.1223 and 95.1225 - Medical Device Radiocommunications Service (MedRadio).

Form No.: N/A.

Type of Review: Revision of a currently approved collection.

Respondents: Business or other for-profit and not-for-profit institutions.

Number of Respondents: 3,120 respondents; 3,120 responses.

Estimated Time Per Response: 1-3 hours.

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

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

Total Annual Burden: 9,120 hours.

Total Annual Cost: $462,600.

Privacy Act Impact Assessment: N/A.

Nature and Extent of Confidentiality: No information is requested that would require assurance of confidentiality.

Needs and Uses: The Commission will submit this information collection to the Office of Management and Budget (OMB) after this 60 day comment period in order to obtain the full three year clearance from them. The Commission is requesting a revision (there has been a program change in the reporting, recordkeeping requirements and/or third party disclosure requirements, the number of respondents/operators increased from 100 to 2,620, therefore, the annual burden and cost has also increased).

The Commission now seeks OMB approval for a revision. On May 24, 2012, the Commission released a Report and Order, ET Docket No. 08-59, FCC 12-54, Amendment of Parts 2 and 95 of the Commission's rules which revised the requirements for manufacturers of transmitters for the "Medical Device Radiocommunication Service"

to include with each transmitting device a statement regarding harmful interference and to label the device in a conspicuous location on the device. The Report and Order also adopted rules for "Medical Body Area Network" (MBAN), which requires the Commission to establish a process by which MBAN users will register and coordinate the use of certain medical devices. The frequency coordinator will make the database available to equipment manufacturers and the public. The coordinator will also notify users of potential frequency conflicts.

Federal Communications Commission.

Marlene H. Dortch,

Secretary,

Office of the Secretary,

Office of Managing Director.

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