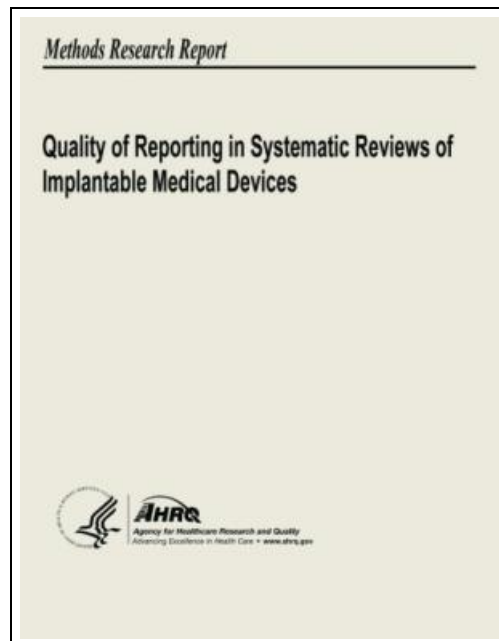


## Quality of Reporting in Systematic Reviews of Implantable Medical Devices



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***(Mr. Noah Cummerata IV)***

## QUALITY OF REPORTING IN SYSTEMATIC REVIEWS OF IMPLANTABLE MEDICAL DEVICES



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CreateSpace Independent Publishing Platform. Paperback. Condition: New. This item is printed on demand. 106 pages. Dimensions: 11.0in. x 8.5in. x 0.2in. Over the past decade, advances in the technologies used in medical devices have profoundly transformed clinical practice and patient management. According to a recent position paper, studies of medical devices typically do not address in sufficient detail study design features and factors that affect safety and effectiveness, including standardization of outcomes and endpoints, evaluation of deviceoperator interactions, and evaluation of characteristics of the clinical practice setting. The evaluation of medical devices poses a number of methodological challenges, and empirical research evaluating such devices has not been performed. The challenges in conducting assessments of medical devices are secondary to the rapid evolution of technology, practitioner input into design, operator learning curve, and variation in the skill with which they are used. These features can potentially impact study outcomes. Systematic reviews have an established role in medical practice and research. Clinicians use systematic reviews to keep abreast of current research and to provide information on the effectiveness of competing interventions. Agencies including the Centers for Medicare and Medicaid in the United States and the National Institute of Clinical Excellence in the United Kingdom use systematic reviews to prioritize funding or coverage decisions. Health care organizations use systematic reviews to shape policy and inform clinical practice guidelines. Despite there being many published systematic reviews of medical devices, no empirical evaluation of the reviews has been performed. A thorough empirical appraisal of systematic reviews is needed to critically assess current practices and identify issues and gaps in reporting. The information generated from an empirical appraisal of systematic reviews of medical devices could be used to develop new items in reporting guidelines and improve the conduct and the quality of reporting of systematic reviews of...



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