

This form may be submitted via E-mail to [mweldon@ansi.org](mailto:mweldon@ansi.org)

**PINS: PROJECT INITIATION NOTIFICATION SYSTEM FORM** *(Effective 01.01.08)*

\*NOTE: Adoptions of an ISO or IEC standards require compliance with the *ANSI Policy Regarding Rights to Nationally Adopt IEC and ISO Standards or Otherwise Use IEC and ISO Material* and with the *ANSI Procedures for the Adoption of ISO and IEC Standards as American National Standards*.

<b>1. Designation of Proposed Standard:</b>	ANSI C63.32	
<b>2. Title of Standard:</b>	American National Standard for Evaluating Immunity of Portable Electronic Medical Devices to Electronic Article Surveillance (EAS) Systems and Metal Detectors	
<b>3. Project Intent:</b> (Check the applicable box below)		
Create new American National Standard (ANS)	X	
*Adopt identical ISO or IEC standard		
*Adopt modified ISO or IEC standard		
<b>*AND</b> this adoption revises this current ANS		
Revise current ANS		
Revise and Redesignate current ANS		
Revise, Redesignate and Consolidate current ANS		
Revise and Partition current ANS		
Reaffirm current ANS		
Reaffirm and Redesignate current ANS		
Addenda to a current ANS under Continuous Maintenance: (this document relates to/updates the following base document that is registered under Continuous Maintenance)		
Supplement to current ANS		
Withdraw current ANS		
Maintain ANS under stabilized maintenance		
<b>4. This standard contains excerpted text from an ISO or IEC standard, but is not an ISO or IEC adoption.</b>		Check here if this standard includes excerpted text from an ISO or IEC standard but is not an identical or modified adoption of an ISO or IEC standard.
<b>5. Provide a brief explanation of the need for the project</b> (see 2.5 of the <i>ANSI Essential Requirements</i> ):	The proliferation of portable electronic medical devices and their potential proximity to electronic article surveillance systems in retail environments and metal detectors in security applications creates the need to evaluate the performance of medical devices when exposed to these electromagnetic fields	
<b>6. Identify the stakeholders</b> (e.g., telecom, consumer, medical, environmental, etc.) <b>likely to be directly impacted by the standard</b> (see 2.5 of the <i>ANSI Essential Requirements</i> ):	EAS system manufacturers and users Metal detector manufacturers and users Medical device manufacturers Test labs Regulators	
<b>7. Unit of Measure: Non Applicable, US, Metric, or Both</b>	M	
<b>8. This PINS revises a previous PINS submittal</b> (see 2.5 of the <i>ANSI Essential Requirements</i> ):		Note: A revised PINS is only required if the previously identified stakeholders have changed substantively (see item 6 on this form.).

<b>9. Description of Contents of Standard:</b> (Provide a one paragraph description, not to exceed 500 characters. Please note in the scope if this standard is intended to be submitted for consideration as an ISO or ISO/IEC JTC-1 standard.)	Test methods and test levels for testing immunity of portable electronic medical devices to EAS systems and metal detectors, using actual EAS systems and metal detectors or simulated signals. Information on incorporation of results into risk management.	
<b>10. Request an Announcement in Standards Action to Solicit New Consensus Body Members</b> (Note that participants from diverse interest categories shall be sought with the objective of achieving balance. See 1.3 and 2.3 of the <i>ANSI Essential Requirements</i> .)		Check here to request the publication in Standards Action of a call for membership on the relevant consensus body.
<b>11. Consumer Product or Service:</b>		Check here if standard covers Consumer Product or Service
<b>12. Accredited Standards Developer Acronym:</b>		
<b>13. Submitter:</b> (Specify Accredited Standards Developer submitter's name and complete contact information, address, phone, email, etc.)	Name:	David Schaefer
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