

**K031894 SCANTEC PAD**Jul 18, 2003  
29 days to decisionK031894 · Product code: **MUI** · Radiology  
Source: <https://www.510kdatabase.net/k031894/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Media, Coupling, Ultrasound (MUI)
Date received	Jun 19, 2003
Decision date	Jul 18, 2003
Days to decision	29 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Sonotech, Inc.</b>
Location	Bellingham, WA, US
Contact	MARGARET J LARSON
510(k) history	14 submissions · 14 cleared · 1995-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k031894/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 18, 2026