

K961692 SONOSCANAug 22, 1996
113 days to decisionK961692 · Product code: **ITX** · Radiology
Source: <https://www.510kdatabase.net/k961692/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Ultrasonic, Diagnostic (ITX)
Date received	May 1, 1996
Decision date	Aug 22, 1996
Days to decision	113 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k961692/> Data sourced from FDA 510(k) public records (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. - Generated May 18, 2026