

**K911114 INTERSON ENDOCAVITY PROBE/ALOKA  
ULTRASOUND SYSTEM**Nov 19, 1991  
257 days to decisionK911114 · Product code: **ITX** · Radiology  
Source: <https://www.510kdatabase.net/k911114/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Ultrasonic, Diagnostic (ITX)
Date received	Mar 7, 1991
Decision date	Nov 19, 1991
Days to decision	257 days
Third-party review	No

**APPLICANT**

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Company	<b>Interson Corp.</b>
Location	Pleasanton, CA, US
Contact	GARY J ALLSEBROOK
510(k) history	9 submissions · 9 cleared · 1989-1991

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k911114/>; 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 21, 2026