

**K911117 INTERSON ENDOCAVITY PROBE FOR ATL
ULTRASOUND SYST**Dec 17, 1991
285 days to decisionK911117 · Product code: **ITX** · Radiology
Source: <https://www.510kdatabase.net/k911117/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Ultrasonic, Diagnostic (ITX)
Date received	Mar 7, 1991
Decision date	Dec 17, 1991
Days to decision	285 days
Third-party review	No

APPLICANT

Company	Interson Corp.
Location	Pleasanton, CA, US
Contact	MICHAEL CHRISTY
510(k) history	9 submissions · 9 cleared · 1989-1991

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k911117/>; 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 21, 2026