FDA 510(k) Premarket Notification

K911115 INTERSON TRANSRECTAL PROBE/ATL ULTRASOUND SYSTEM

CLEARED

Dec 17, 1991
285 days to decision

K911115 · Product code: **ITX** · Radiology
Source: https://www.510kdatabase.net/k911115/

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	GARY J ALLSEBROOK
510(k) history	9 submissions · 9 cleared · 1989-1991