FDA 510(k) Premarket Notification

K911114 INTERSON ENDOCAVITY PROBE/ALOKA ULTRASOUND SYSTEM

CLEARED

Nov 19, 1991
257 days to decision

K911114 · Product code: **ITX** · Radiology
Source: https://www.510kdatabase.net/k911114/

SUBMISSION DETAILS

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

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