FDA 510(k) Premarket Notification

K900882 MULTI DOPPLEX, MD-1

CLEARED

Aug 22, 1990
177 days to decision

K900882 · Product code: **JAF** · Radiology
Source: https://www.510kdatabase.net/k900882/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Ultrasonic, Nonfetal (JAF)
Date received	Feb 26, 1990
Decision date	Aug 22, 1990
Days to decision	177 days
Third-party review	No

APPLICANT

Company	**Huntleigh Technology, Inc.**
Location	Walker, MI, US
Contact	JAMES BRITTON
510(k) history	23 submissions · 23 cleared · 1981-1999