FDA 510(k) Premarket Notification

K900881 OR-340 INTRAOPERATIVE ULTRASOUND SYSTEM

CLEARED

Sep 25, 1990
211 days to decision

K900881 · Product code: **IYO** · Radiology
Source: https://www.510kdatabase.net/k900881/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Feb 26, 1990
Decision date	Sep 25, 1990
Days to decision	211 days
Third-party review	No

APPLICANT

Company	**Codman & Shurtleff, Inc.**
Location	Mchenry, IL, US
Contact	JAMES R VEALE
510(k) history	152 submissions · 151 cleared · 1976-2020