FDA 510(k) Premarket Notification

K800091 COMBISON 100-MODEL 100 & 110

CLEARED

Feb 11, 1980
28 days to decision

K800091 · Product code: **IYO** · Radiology
Source: https://www.510kdatabase.net/k800091/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Jan 14, 1980
Decision date	Feb 11, 1980
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Kt Medical, Inc.**
Location	Mchenry, IL, US
510(k) history	13 submissions · 13 cleared · 1980-1980