FDA 510(k) Premarket Notification

K813002 MAVIS C

CLEARED

Mar 1, 1982
126 days to decision

K813002 · Product code: **DXK** · Radiology
Source: https://www.510kdatabase.net/k813002/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Echocardiograph (DXK)
Date received	Oct 26, 1981
Decision date	Mar 1, 1982
Days to decision	126 days
Third-party review	No

APPLICANT

Company	**Philips Medical Systems (Cleveland), Inc.**
Location	Mchenry, IL, US
510(k) history	190 submissions · 190 cleared · 1977-2017