

K813002 MAVIS CMar 1, 1982
126 days to decisionK813002 · 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

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Device record: <https://www.510kdatabase.net/k813002/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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