

K791334 MULTISCOPEAug 10, 1979
21 days to decisionK791334 · Product code: **DQD** · CardiovascularSource: <https://www.510kdatabase.net/k791334/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stethoscope, Electronic (DQD)
Date received	Jul 20, 1979
Decision date	Aug 10, 1979
Days to decision	21 days
Third-party review	No

APPLICANT

Company	Amdia Medical Products
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1979-1979

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Device record: <https://www.510kdatabase.net/k791334/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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