

K791338 DUAL HEAD STETHOSCOPEAug 10, 1979
21 days to decisionK791338 · Product code: **DQD** · CardiovascularSource: <https://www.510kdatabase.net/k791338/>**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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k791338/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 2, 2026