

K831763 MAX IJan 27, 1984
240 days to decisionK831763 · Product code: **DQD** · CardiovascularSource: <https://www.510kdatabase.net/k831763/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stethoscope, Electronic (DQD)
Date received	Jun 1, 1983
Decision date	Jan 27, 1984
Days to decision	240 days
Third-party review	No

APPLICANT

Company	Trade Max, Inc.
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1984-1984

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Device record: <https://www.510kdatabase.net/k831763/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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