FDA 510(k) Premarket Notification

K831763 MAX I

CLEARED

Jan 27, 1984
240 days to decision

K831763 · Product code: **DQD** · Cardiovascular
Source: 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