FDA 510(k) Premarket Notification

K780023 BLACK BAG, ELECTRONIC

CLEARED

Feb 21, 1978
48 days to decision

K780023 · Product code: **LDE** · Cardiovascular
Source: https://www.510kdatabase.net/k780023/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stethoscope, Manual (LDE)
Date received	Jan 4, 1978
Decision date	Feb 21, 1978
Days to decision	48 days
Third-party review	No

APPLICANT

Company	**Marshall Electronics, Inc.**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1978-1981