FDA 510(k) Premarket Notification

K231551 Stethophone

CLEARED

Oct 12, 2023
135 days to decision

K231551 · Product code: **DQD** · Cardiovascular
Source: https://www.510kdatabase.net/k231551/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stethoscope, Electronic (DQD)
Date received	May 30, 2023
Decision date	Oct 12, 2023
Days to decision	135 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Sparrow Acoustics, Inc.**
Location	Lucasville, CA
Contact	Nadia Ivanova
510(k) history	4 submissions · 4 cleared · 2023-2025