FDA 510(k) Premarket Notification

K791338 DUAL HEAD STETHOSCOPE

CLEARED

Aug 10, 1979
21 days to decision

K791338 · Product code: **DQD** · Cardiovascular
Source: 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