FDA 510(k) Premarket Notification

K813510 CARDIOSONIC ACOUSTIC AMPLIFIER

CLEARED

Jan 12, 1982
28 days to decision

K813510 · Product code: **LDE** · Cardiovascular
Source: https://www.510kdatabase.net/k813510/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stethoscope, Manual (LDE)
Date received	Dec 15, 1981
Decision date	Jan 12, 1982
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Srs Co.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1982-1983