

K813510 CARDIOSONIC ACOUSTIC AMPLIFIERJan 12, 1982
28 days to decisionK813510 · Product code: **LDE** · CardiovascularSource: <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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k813510/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 2, 2026