

K812520 280/4 MINI-DEFIBRILLATORSep 25, 1981
23 days to decisionK812520 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k812520/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Sep 2, 1981
Decision date	Sep 25, 1981
Days to decision	23 days
Third-party review	No

APPLICANT

Company	Life Science Instrumentation, Inc.
Location	Mchenry, IL, US
510(k) history	20 submissions · 20 cleared · 1981-1985

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Device record: <https://www.510kdatabase.net/k812520/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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