

**K812519 2000 SERIES PORTABLE RESUSCITATION SYST.**Sep 25, 1981  
23 days to decisionK812519 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k812519/>**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	<b>Life Science Instrumentation, Inc.</b>
Location	Mchenry, IL, US
510(k) history	20 submissions · 20 cleared · 1981-1985

---

**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k812519/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated July 8, 2026