

**K891122 DISP. ACCESSORY TO EXTERNAL TRANS. CARDIAC PACE.**Sep 13, 1989  
195 days to decisionK891122 · Product code: **LDD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k891122/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Mar 2, 1989
Decision date	Sep 13, 1989
Days to decision	195 days
Third-party review	No

**APPLICANT**

---

Company	<b>Big Valley Medical</b>
Location	Orangevale, CA, US
Contact	ROBERT S JONES
510(k) history	1 submissions · 1 cleared · 1989-1989

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k891122/> 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 7, 2026