

**K911604 MDPHONE CELLULAR + RESCUE (DEFIB SYSTEM)**Sep 23, 1991  
166 days to decisionK911604 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k911604/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Apr 10, 1991
Decision date	Sep 23, 1991
Days to decision	166 days
Third-party review	No

**APPLICANT**

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Company	<b>Medphone Corp.</b>
Location	Washington, DC, US
Contact	GLENN KRAPF
510(k) history	2 submissions · 2 cleared · 1987-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k911604/>; 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 6, 2026