

**K002079 MULTI-FUNCTION DEFIBRILLATION ELECTRODE**Feb 22, 2001  
227 days to decisionK002079 · Product code: **MKJ** · CardiovascularSource: <https://www.510kdatabase.net/k002079/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Jul 10, 2000
Decision date	Feb 22, 2001
Days to decision	227 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Medicotest A/S</b>
Location	Washington, DC, US
Contact	EMALEE G MURPHY
510(k) history	7 submissions · 7 cleared · 1997-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k002079/> 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 June 30, 2026