

**K960329 DISPOSABLE HYDROGEL DEFIBRILLATION  
ELECTRODE/PAD**Jul 2, 1996  
160 days to decisionK960329 · Product code: **MKJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k960329/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Jan 24, 1996
Decision date	Jul 2, 1996
Days to decision	160 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Contour Medical Technology, Inc.</b>
Location	Lavergne, TN, US
Contact	JOSH TRANTUM
510(k) history	9 submissions · 9 cleared · 1996-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960329/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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