

**K811638 DATASCOPE INTERNAL DEFIBRILLATOR PADDLES**Jul 10, 1981  
30 days to decisionK811638 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k811638/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Jun 10, 1981
Decision date	Jul 10, 1981
Days to decision	30 days
Third-party review	No

**APPLICANT**

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Company	<b>Datascope Corp.</b>
Location	Mchenry, IL, US
510(k) history	136 submissions · 135 cleared · 1976-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k811638/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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