

**K003819 HEARTSTREAM ATTENUATED DEFIBRILLATION  
PADS, MODEL M3870A**May 2, 2001  
142 days to decisionK003819 · Product code: **MKJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k003819/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Dec 11, 2000
Decision date	May 2, 2001
Days to decision	142 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Agilent Technologies, Inc.</b>
Location	Pittsburgh, PA, US
Contact	TERESA SKARR
Website	<a href="http://www.agilent.com">http://www.agilent.com</a>
510(k) history	30 submissions · 30 cleared · 1985-2017

Agilent Technologies, Inc. is an American global company that provides instruments, software, services, and consumables for laboratories. Headquartered in Santa Clara, California, Agilent was established in 1999 as a spin-off from Hewlett-Packard. The company has received FDA 510(k) clearances from total submissions. Cardiovascular devices represent the dominant focus, accounting for approximately 80% of regulatory submissions. Agilent's FDA 510(k) clearance history spans from 1985 to 2017, establishing a long track record in medical device regulation. Notable cleared dev...

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

Device record: <https://www.510kdatabase.net/k003819/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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