

**K033896 DEFIBTECH AED WITH ATTENUATED  
DEFIBRILLATION/MONITORING PADS, MODELS DDU-100 WITH  
DDP-200P**Jun 16, 2004  
183 days to decisionK033896 · Product code: **MKJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k033896/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Dec 16, 2003
Decision date	Jun 16, 2004
Days to decision	183 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Defibtech, LLC</b>
Location	Guilford, CT, US
Contact	JOHN L ROGERS
510(k) history	9 submissions · 8 cleared · 2002-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k033896/> 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 May 25, 2026