

**K113787 DDU-120 FULLY AUTOMATIC EXTERNAL
DEFIBRILLATOR AND ACCESSORIES**Jan 4, 2013
379 days to decisionK113787 · Product code: **MKJ** · Cardiovascular
Source: <https://www.510kdatabase.net/k113787/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Dec 22, 2011
Decision date	Jan 4, 2013
Days to decision	379 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Defibtech, LLC
Location	Guilford, CT, US
Contact	Ed Horton
510(k) history	9 submissions · 8 cleared · 2002-2021

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k113787/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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