

**K013896 SENTRY SEMIAUTOMATIC EXTERNAL
DEFIBRILLATOR**Jun 19, 2002
208 days to decisionK013896 · Product code: **MKJ** · Cardiovascular
Source: <https://www.510kdatabase.net/k013896/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - ST
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Nov 23, 2001
Decision date	Jun 19, 2002
Days to decision	208 days
Third-party review	No
Other names	EXTERNAL DEFIBRILLATION PADS (ELECTRODES) DDP-100; BATTERY PACK 1200MAH; BA

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k013896/>; Data sourced from FDA 510(k) public records (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. - Generated May 25, 2026