FDA 510(k) Premarket Notification

K013896 SENTRY SEMIAUTOMATIC EXTERNAL DEFIBRILLATOR

CLEARED

Jun 19, 2002
208 days to decision

K013896 · 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