FDA 510(k) Premarket Notification

K881273 HORIZON 2000 ARRHYTHMIA MONITORING OPTION

CLEARED

Jul 12, 1988
109 days to decision

K881273 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k881273/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Mar 25, 1988
Decision date	Jul 12, 1988
Days to decision	109 days
Third-party review	No

APPLICANT

Company	**Mennen Medical, Inc.**
Location	Clarence, NY, US
Contact	THOMAS CONNELLY
510(k) history	34 submissions · 34 cleared · 1985-2004