FDA 510(k) Premarket Notification

K832354 HOLTA-MED INFORMER PLAYBACK

CLEARED

Jan 27, 1984
193 days to decision

K832354 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k832354/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jul 18, 1983
Decision date	Jan 27, 1984
Days to decision	193 days
Third-party review	No

APPLICANT

Company	**Holta-Med**
Location	Walker, MI, US
510(k) history	2 submissions · 2 cleared · 1983-1984

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k832354/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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