FDA 510(k) Premarket Notification

K810104 ECG ARRHYTHMIA STIMULATOR

CLEARED

Jan 29, 1981
13 days to decision

K810104 · Product code: **DPS** · Cardiovascular
Source: https://www.510kdatabase.net/k810104/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Jan 16, 1981
Decision date	Jan 29, 1981
Days to decision	13 days
Third-party review	No

APPLICANT

Company	**Belmont Instrument Corp.**
Location	Mchenry, IL, US
510(k) history	20 submissions · 19 cleared · 1981-2013

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k810104/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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