FDA 510(k) Premarket Notification

K822316 HEART MONITOR BRACELET ALARM

CLEARED

Sep 21, 1982
49 days to decision

K822316 · Product code: **DRI** · Cardiovascular
Source: https://www.510kdatabase.net/k822316/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Line Isolation (DRI)
Date received	Aug 3, 1982
Decision date	Sep 21, 1982
Days to decision	49 days
Third-party review	No

APPLICANT

Company	**Hmba**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1982-1982