

K822316 HEART MONITOR BRACELET ALARMSep 21, 1982
49 days to decisionK822316 · 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

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Device record: <https://www.510kdatabase.net/k822316/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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