

K813174 BRADYCARDIA ALARMJan 22, 1982
77 days to decisionK813174 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k813174/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Nov 6, 1981
Decision date	Jan 22, 1982
Days to decision	77 days
Third-party review	No

APPLICANT

Company	Douglas Scientific Products
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1980-1982

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Device record: <https://www.510kdatabase.net/k813174/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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