

K830282 ELECTRONIC EXERCISE #313-PULSE RATEApr 14, 1983
78 days to decisionK830282 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k830282/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Jan 26, 1983
Decision date	Apr 14, 1983
Days to decision	78 days
Third-party review	No

APPLICANT

Company	Walton Manufacturer Co.
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1983-1983

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Device record: <https://www.510kdatabase.net/k830282/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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