

**K833552 TREND PROCESSOR OPTION FOR #515 NEONAT**Jan 27, 1984  
105 days to decisionK833552 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k833552/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Oct 14, 1983
Decision date	Jan 27, 1984
Days to decision	105 days
Third-party review	No

**APPLICANT**

---

Company	<b>Ge Medical Systems Information Technologies</b>
Location	Mchenry, IL, US
510(k) history	136 submissions · 132 cleared · 1978-2012

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k833552/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 26, 2026