

**K821755 TELCOR 2**Jul 2, 1982  
17 days to decisionK821755 · Product code: **DRT** · Cardiovascular  
Source: <https://www.510kdatabase.net/k821755/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Jun 15, 1982
Decision date	Jul 2, 1982
Days to decision	17 days
Third-party review	No

**APPLICANT**

---

Company	<b>Burdick Corp.</b>
Location	Mchenry, IL, US
Website	<a href="https://www.cardinalhealth.com">https://www.cardinalhealth.com</a>
510(k) history	38 submissions · 38 cleared · 1976-1994

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k821755/>; 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 22, 2026