

**K823844 HORIZON ECG SYSTEM**Jan 10, 1985  
751 days to decisionK823844 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k823844/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Dec 21, 1982
Decision date	Jan 10, 1985
Days to decision	751 days
Third-party review	No

**APPLICANT**

---

Company	<b>Beckman Instruments, Inc.</b>
Location	Mchenry, IL, US
Contact	JACK SOROKIN
Website	<a href="https://www.beckman.com">https://www.beckman.com</a>
510(k) history	281 submissions · 281 cleared · 1976-1998

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k823844/> 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