

**K102723 MODEL 950-12L HOLTER RECORDER WITHOUT ANALYSIS**Dec 9, 2010  
79 days to decisionK102723 · Product code: **MWJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k102723/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph, Ambulatory (without Analysis) (MWJ)
Date received	Sep 21, 2010
Decision date	Dec 9, 2010
Days to decision	79 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Memtec Corp.</b>
Location	Salem, NH, US
Contact	DENNIS GARBOSKI
510(k) history	3 submissions · 3 cleared · 1997-2011

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k102723/> 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 June 19, 2026