

**K950112 CENTRALITE(R)-DLL SERIES**Aug 16, 1995  
217 days to decisionK950112 · Product code: **IWE** · Radiology  
Source: <https://www.510kdatabase.net/k950112/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Patient Position, Light-beam (IWE)
Date received	Jan 11, 1995
Decision date	Aug 16, 1995
Days to decision	217 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Diacor, Inc.</b>
Location	Salt Lake City, UT, US
Contact	GLENN N WATERMAN
510(k) history	10 submissions · 10 cleared · 1986-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k950112/> 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 17, 2026