

**K893398 CENTRALITE BACKPOINTER**Aug 17, 1989  
108 days to decisionK893398 · Product code: **IWE** · Radiology  
Source: <https://www.510kdatabase.net/k893398/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Patient Position, Light-beam (IWE)
Date received	May 1, 1989
Decision date	Aug 17, 1989
Days to decision	108 days
Third-party review	No

**APPLICANT**

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Company	<b>Diacor, Inc.</b>
Location	Salt Lake City, UT, US
Contact	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/k893398/>; 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