

**K883855 PREDICTOR II SIGNAL AVERAGING ECG**Nov 3, 1988  
51 days to decisionK883855 · Product code: **DXG** · CardiovascularSource: <https://www.510kdatabase.net/k883855/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Diagnostic, Pre-programmed, Single-function (DXG)
Date received	Sep 13, 1988
Decision date	Nov 3, 1988
Days to decision	51 days
Third-party review	No

**APPLICANT**

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Company	<b>Corazonix Corp.</b>
Location	Oklahoma City, OK, US
Contact	JAMES H LUBY
510(k) history	2 submissions · 2 cleared · 1987-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k883855/>; 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 July 7, 2026