

**K880533 VITEL III CM**May 13, 1988  
94 days to decisionK880533 · Product code: **DXH** · Cardiovascular  
Source: <https://www.510kdatabase.net/k880533/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Electrocardiograph, Telephone (DXH)
Date received	Feb 9, 1988
Decision date	May 13, 1988
Days to decision	94 days
Third-party review	No

**APPLICANT**

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Company	<b>Equimed Corp.</b>
Location	Plymouth, MN, US
Contact	DAVID C GUST
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/k880533/>; 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 28, 2026