

**K923054 WITT BIOMEDICAL SERIES II-EP**Feb 25, 1993  
246 days to decisionK923054 · Product code: **MHX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k923054/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Jun 24, 1992
Decision date	Feb 25, 1993
Days to decision	246 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Witt Biomedical Corporation</b>
Location	Indian Harbour Bch, FL, US
Contact	TERENCE R WITT
510(k) history	6 submissions · 6 cleared · 1988-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k923054/> 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 19, 2026