

**K973336 IMEX STETHODOP**Feb 5, 1998  
154 days to decisionK973336 · Product code: **JAF** · Radiology  
Source: <https://www.510kdatabase.net/k973336/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Ultrasonic, Nonfetal (JAF)
Date received	Sep 4, 1997
Decision date	Feb 5, 1998
Days to decision	154 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Imex Medical Systems, Inc.</b>
Location	Mchenry, IL, US
Contact	DAVID W WAGNER
510(k) history	19 submissions · 19 cleared · 1983-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k973336/> 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