

**K970908 HOWTEK 960**May 20, 1997  
70 days to decisionK970908 · Product code: **LMA** · Radiology  
Source: <https://www.510kdatabase.net/k970908/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Digitizer, Image, Radiological (LMA)
Date received	Mar 11, 1997
Decision date	May 20, 1997
Days to decision	70 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Howtek, Inc.</b>
Location	Hudson, NH, US
Contact	M. RUSSELL LEONARD
510(k) history	5 submissions · 5 cleared · 1996-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k970908/> 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