

**K103785 MOBILE MIM**Feb 4, 2011  
39 days to decisionK103785 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k103785/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Dec 27, 2010
Decision date	Feb 4, 2011
Days to decision	39 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Mim Software, Inc.</b>
Location	Cleveland, OH, US
Contact	LYNN HANIGAN
510(k) history	22 submissions · 22 cleared · 2011-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k103785/> 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 17, 2026