

K052379 MIM 3.5 (CIRCA)Oct 31, 2005
62 days to decisionK052379 · Product code: **LLZ** · Radiology
Source: <https://www.510kdatabase.net/k052379/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Aug 30, 2005
Decision date	Oct 31, 2005
Days to decision	62 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Mimvista Corp.
Location	Cleveland, OH, US
Contact	PETER SIMMELINK
510(k) history	4 submissions · 4 cleared · 2005-2007

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k052379/> Data sourced from FDA 510(k) public records (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. - Generated May 18, 2026