

K120484 VISIA ONCOLOGYMar 27, 2012
39 days to decisionK120484 · Product code: **LLZ** · Radiology
Source: <https://www.510kdatabase.net/k120484/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Feb 17, 2012
Decision date	Mar 27, 2012
Days to decision	39 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Mevis Medical Solutions AG
Location	Pewaukee, WI, US
Contact	THOMAS E TYNES
510(k) history	6 submissions · 6 cleared · 2011-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k120484/> 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 26, 2026