

K113543 ONDEMAND3DDec 16, 2011
15 days to decisionK113543 · Product code: **LLZ** · Radiology
Source: <https://www.510kdatabase.net/k113543/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Dec 1, 2011
Decision date	Dec 16, 2011
Days to decision	15 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	Cybermed, Inc.
Location	San Leandro, CA, US
Contact	YONG KI IM
510(k) history	6 submissions · 6 cleared · 2001-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k113543/> 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 28, 2026