

K082135 SHOWCASEOct 21, 2008
84 days to decisionK082135 · Product code: **LLZ** · Radiology
Source: <https://www.510kdatabase.net/k082135/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Jul 29, 2008
Decision date	Oct 21, 2008
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Trillium Technology, Inc.
Location	San Leandro, CA, US
Contact	GARY J ALLSEBROOK
510(k) history	1 submissions · 1 cleared · 2008-2008

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k082135/> 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 17, 2026