

K041029 KINETDXJul 8, 2004
78 days to decisionK041029 · Product code: **LLZ** · Radiology
Source: <https://www.510kdatabase.net/k041029/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Apr 21, 2004
Decision date	Jul 8, 2004
Days to decision	78 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Siemens Medi Cal Solutions, Inc.
Location	Ann Arbor, MI, US
Contact	ANA LADINO
510(k) history	32 submissions · 32 cleared · 2004-2020

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k041029/> 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 June 19, 2026