FDA 510(k) Premarket Notification

K041029 KINETDX

CLEARED

Jul 8, 2004
78 days to decision

K041029 · 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