FDA 510(k) Premarket Notification

K132934 MULTIX SELECT DR

CLEARED

Apr 10, 2014
204 days to decision

K132934 · Product code: **KPR** · Radiology
Source: https://www.510kdatabase.net/k132934/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Sep 18, 2013
Decision date	Apr 10, 2014
Days to decision	204 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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