

**K132934 MULTIX SELECT DR**Apr 10, 2014  
204 days to decisionK132934 · Product code: **KPR** · Radiology  
Source: <https://www.510kdatabase.net/k132934/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k132934/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 16, 2026