

**K001201 MULTIX COMPACT K AND MULTIX L RADIOGRAPHIC  
X-RAY SYSTEMS**Jun 1, 2000  
49 days to decisionK001201 · Product code: **KPR** · Radiology  
Source: <https://www.510kdatabase.net/k001201/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Apr 13, 2000
Decision date	Jun 1, 2000
Days to decision	49 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Siemens Medical Solutions USA, Inc.</b>
Location	Hoffman Estates, IL, US
Contact	MALGORZATA STANEK
510(k) history	778 submissions · 778 cleared · 1980-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k001201/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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