

K233532 MULTIX Impact ENov 29, 2023
27 days to decisionK233532 · Product code: **KPR** · Radiology
Source: <https://www.510kdatabase.net/k233532/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, X-ray, Stationary (KPR)
Date received	Nov 2, 2023
Decision date	Nov 29, 2023
Days to decision	27 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Siemens Medical Solutions
Location	Danvers, MA, US
Contact	Camila Rodriguez Valentin
510(k) history	15 submissions · 15 cleared · 2003-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k233532/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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