

K201670 YSIO X.preeOct 21, 2020
124 days to decisionK201670 · Product code: **KPR** · Radiology
Source: <https://www.510kdatabase.net/k201670/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Jun 19, 2020
Decision date	Oct 21, 2020
Days to decision	124 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Siemens Medical Solutions USA, Inc.
Location	Hoffman Estates, IL, US
Contact	Andrew Turner
510(k) history	779 submissions · 779 cleared · 1980-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k201670/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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