

K811934 DSS 500Aug 13, 1981
38 days to decisionK811934 · Product code: **KPR** · Radiology
Source: <https://www.510kdatabase.net/k811934/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Jul 6, 1981
Decision date	Aug 13, 1981
Days to decision	38 days
Third-party review	No

APPLICANT

Company	Philips Medical Systems (Cleveland), Inc.
Location	Mchenry, IL, US
510(k) history	190 submissions · 190 cleared · 1977-2017

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Device record: <https://www.510kdatabase.net/k811934/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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