

K800788 SIREGRAPH CMay 14, 1980
36 days to decisionK800788 · Product code: **KPR** · Radiology
Source: <https://www.510kdatabase.net/k800788/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Apr 8, 1980
Decision date	May 14, 1980
Days to decision	36 days
Third-party review	No

APPLICANT

Company	Siemens Medical Solutions USA, Inc.
Location	Hoffman Estates, IL, US
510(k) history	778 submissions · 778 cleared · 1980-2026

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k800788/> Data sourced from FDA 510(k) public records (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. - Generated May 15, 2026