

K810797 LOW DOSE SERVOCHESTApr 14, 1981
22 days to decisionK810797 · Product code: **KPR** · Radiology
Source: <https://www.510kdatabase.net/k810797/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Mar 23, 1981
Decision date	Apr 14, 1981
Days to decision	22 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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k810797/> 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 16, 2026