FDA 510(k) Premarket Notification

K810797 LOW DOSE SERVOCHEST

CLEARED

Apr 14, 1981
22 days to decision

K810797 · 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