FDA 510(k) Premarket Notification

K811934 DSS 500

CLEARED

Aug 13, 1981
38 days to decision

K811934 · 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