FDA 510(k) Premarket Notification

K800788 SIREGRAPH C

CLEARED

May 14, 1980
36 days to decision

K800788 · 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