FDA 510(k) Premarket Notification

K803124 SIDOS U.

CLEARED

Jan 22, 1981
42 days to decision

K803124 · Product code: **KPQ** · Radiology
Source: https://www.510kdatabase.net/k803124/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Simulation, Radiation Therapy (KPQ)
Date received	Dec 11, 1980
Decision date	Jan 22, 1981
Days to decision	42 days
Third-party review	No

APPLICANT

Company	**Siemens Corp.**
Location	Mchenry, IL, US
Website	http://www.siemens.it/
510(k) history	66 submissions · 66 cleared · 1978-2010

Siemens Corp. is a global technology company headquartered in McHenry, US. The company develops medical imaging and diagnostic equipment for healthcare providers worldwide. Siemens has received FDA 510(k) clearances from total submissions. The company's regulatory focus centers on Radiology devices, which represent the dominant category of its cleared portfolio. FDA 510(k) clearances span from 1978 to 2010, establishing a significant historical record in medical device regulation. Recent cleared devices include advanced imaging systems such as CT scanners, MR systems, X-r...