FDA 510(k) Premarket Notification

K801723 UR 150-UR 150T

CLEARED

Aug 13, 1980
20 days to decision

K801723 · Product code: **KQS** · Radiology
Source: https://www.510kdatabase.net/k801723/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Table, Cystometric, Non-electric And Accessories (KQS)
Date received	Jul 24, 1980
Decision date	Aug 13, 1980
Days to decision	20 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...