FDA 510(k) Premarket Notification

K951358 SIRESKOP SX

CLEARED

May 15, 1995
52 days to decision

K951358 · Product code: **JAA** · Radiology
Source: https://www.510kdatabase.net/k951358/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Fluoroscopic, Image-intensified (JAA)
Date received	Mar 24, 1995
Decision date	May 15, 1995
Days to decision	52 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Siemens Medical Solutions USA, Inc.**
Location	Hoffman Estates, IL, US
Contact	KATHLEEN RUTHERFORD
510(k) history	778 submissions · 778 cleared · 1980-2026