FDA 510(k) Premarket Notification

K090136 STERIS CMAX XLT SURGICAL TABLE

CLEARED

Mar 20, 2009
58 days to decision

K090136 · Product code: **FQO** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k090136/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Table, Operating-room, Ac-powered (FQO)
Date received	Jan 21, 2009
Decision date	Mar 20, 2009
Days to decision	58 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**STERIS Corporation**
Location	Mentor, OH, US
Contact	JOHN ROBERT SCOVILLE
510(k) history	204 submissions · 202 cleared · 1997-2026