

**K090136 STERIS CMAX XLT SURGICAL TABLE**Mar 20, 2009  
58 days to decisionK090136 · Product code: **FQO** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k090136/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k090136/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 23, 2026