

**K101409 SYSTEM IE LIQUID CHEMICAL CTERILANT
PROCESSING SYSTEM**Aug 2, 2010
75 days to decisionK101409 · Product code: **MED** · General Hospital
Source: <https://www.510kdatabase.net/k101409/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Sterilant, Medical Devices (MED)
Date received	May 19, 2010
Decision date	Aug 2, 2010
Days to decision	75 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k101409/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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