

K161683 SYSTEM 1E Liquid Chemical Sterilant Processing System

Aug 5, 2016
49 days to decisionK161683 · Product code: **MED** · General Hospital
Source: <https://www.510kdatabase.net/k161683/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Sterilant, Medical Devices (MED)
Date received	Jun 17, 2016
Decision date	Aug 5, 2016
Days to decision	49 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k161683/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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