FDA 510(k) Premarket Notification

K102462 SYSTEM 1E LIQUID CHEMICAL STERILANT PROCESSING SYSTEM

CLEARED

Sep 21, 2010
25 days to decision

K102462 · Product code: **MED** · General Hospital
Source: https://www.510kdatabase.net/k102462/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Sterilant, Medical Devices (MED)
Date received	Aug 27, 2010
Decision date	Sep 21, 2010
Days to decision	25 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.net
Device record: https://www.510kdatabase.net/k102462/ Data sourced from FDA 510(k) public records (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. - Generated June 20, 2026