

K173256 System 1 endo Liquid Chemical Sterilant Processing System

Mar 2, 2018
143 days to decisionK173256 · Product code: **MED** · General Hospital
Source: <https://www.510kdatabase.net/k173256/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sterilant, Medical Devices (MED)
Date received	Oct 10, 2017
Decision date	Mar 2, 2018
Days to decision	143 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k173256/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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