

K251666 SYSTEM 1E Liquid Chemical Sterilant Processing System, Model P6700Apr 8, 2026
313 days to decisionK251666 · Product code: **MED** · General Hospital
Source: <https://www.510kdatabase.net/k251666/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sterilant, Medical Devices (MED)
Date received	May 30, 2025
Decision date	Apr 8, 2026
Days to decision	313 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	SYSTEM 1 endo Liquid Chemical Sterilant Processing System, Model P6900

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k251666/> 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 May 13, 2026