FDA 510(k) Premarket Notification

CLEARED

K251666 SYSTEM 1E Liquid Chemical Sterilant Processing System, Model P6700

Apr 8, 2026
313 days to decision

K251666 · 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