

K222093 V-PRO maX 2 Low Temperature Sterilization System, V-PRO maX Low Temperature Sterilization SystemSep 9, 2022
56 days to decisionK222093 · Product code: MLR · General Hospital
Source: <https://www.510kdatabase.net/k222093/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Sterilizer, Chemical (MLR)
Date received	Jul 15, 2022
Decision date	Sep 9, 2022
Days to decision	56 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Steris
Location	Mentor, OH, US
Contact	Anthony Piotrkowski
Website	https://www.steris.com
510(k) history	21 submissions · 19 cleared · 2021-2026

Steris is a leading global provider of products and services supporting patient care with emphasis on infection prevention. The company operates with a manufacturing facility in Mentor, Ohio, and serves hospitals, surgery centers, pharmaceutical manufacturers, and research laboratories worldwide. Steris has received FDA 510(k) clearances from total submissions since 2021. The company specializes in General Hospital devices, which represent 81% of its regulatory submissions. Recent clearances include sterilization systems, chemical indicators, biological indicators, and wa...