

**K102330 AMSCO V-PRO MAX LOW TEMPERATURE
STERILIZATION SYSTEM**Aug 12, 2011
360 days to decisionK102330 · Product code: MLR · General Hospital
Source: <https://www.510kdatabase.net/k102330/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sterilizer, Chemical (MLR)
Date received	Aug 17, 2010
Decision date	Aug 12, 2011
Days to decision	360 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k102330/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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