

**K131120 AMSCO V-PRO 1 LOW, V-PRO 1 PLUS, V-PRO MAX
LOW TEMPERATURE STERILIZATION SYSTEM**

Jul 16, 2013
85 days to decision

K131120 · Product code: MLR · General Hospital
Source: <https://www.510kdatabase.net/k131120/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sterilizer, Chemical (MLR)
Date received	Apr 22, 2013
Decision date	Jul 16, 2013
Days to decision	85 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k131120/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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