FDA 510(k) Premarket Notification

K102330 AMSCO V-PRO MAX LOW TEMPERATURE STERILIZATION SYSTEM

CLEARED

Aug 12, 2011
360 days to decision

K102330 · 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