FDA 510(k) Premarket Notification

K160433 V-PRO® 1 Plus and V-PRO® maX Low Temperature

CLEARED

Jul 6, 2016
141 days to decision

K160433 · Product code: **MLR** · General Hospital
Source: https://www.510kdatabase.net/k160433/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sterilizer, Chemical (MLR)
Date received	Feb 16, 2016
Decision date	Jul 6, 2016
Days to decision	141 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