K091022 MODIFICATION TO RESERT XL HLD HIGH LEVEL DISINFECTANT

CLEARED

Apr 12, 2010
368 days to decision

K091022 · Product code: **MED** · General Hospital
Source: https://www.510kdatabase.net/k091022/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sterilant, Medical Devices (MED)
Date received	Apr 9, 2009
Decision date	Apr 12, 2010
Days to decision	368 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**STERIS Corporation**
Location	Mentor, OH, US
Contact	JOHN SCOVILLE JR.
510(k) history	204 submissions · 202 cleared · 1997-2026