

K053389 AESCULAP STERILCONTAINER SYSTEMFeb 15, 2006
72 days to decisionK053389 · Product code: **FRG** · General Hospital
Source: <https://www.510kdatabase.net/k053389/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wrap, Sterilization (FRG)
Date received	Dec 5, 2005
Decision date	Feb 15, 2006
Days to decision	72 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Aesculap, Inc.
Location	Burlingame, CA, US
Contact	MATTHEW M HULL
510(k) history	207 submissions · 201 cleared · 1991-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k053389/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 28, 2026