

K062510 AESCULAP STERILCONTAINER SYSTEMNov 17, 2006
81 days to decisionK062510 · Product code: **FRG** · General Hospital
Source: <https://www.510kdatabase.net/k062510/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wrap, Sterilization (FRG)
Date received	Aug 28, 2006
Decision date	Nov 17, 2006
Days to decision	81 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Aesculap
Location	Center Valley, PA, US
Contact	MATTHEW M HULL
510(k) history	10 submissions · 10 cleared · 2003-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k062510/> 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 May 19, 2026