FDA 510(k) Premarket Notification

K062510 AESCULAP STERILCONTAINER SYSTEM

CLEARED

Nov 17, 2006
81 days to decision

K062510 · 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