

**K041623 REUSABLE STERILCONTAINER FILTER MODEL  
#JK090**Dec 13, 2004  
180 days to decisionK041623 · Product code: **FRG** · General Hospital  
Source: <https://www.510kdatabase.net/k041623/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wrap, Sterilization (FRG)
Date received	Jun 16, 2004
Decision date	Dec 13, 2004
Days to decision	180 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k041623/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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