

**K030853 STERITITE PERFORATED BASE RIGID REUSABLE  
STERILIZATION CONTAINER SYSTEM WITH  
SCF02-POLYPROPYLENE NON-WOVEN DISPOSABLE FI**

Mar 21, 2003  
3 days to decision

K030853 · Product code: **FRG** · General Hospital  
Source: <https://www.510kdatabase.net/k030853/>

**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wrap, Sterilization (FRG)
Date received	Mar 18, 2003
Decision date	Mar 21, 2003
Days to decision	3 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Case Medical, Inc.</b>
Location	Ridgefield, NJ, US
Contact	MARCIA FRIEZE
510(k) history	14 submissions · 14 cleared · 1997-2022

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k030853/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 26, 2026