

**K062697 SIS HERNIA REPAIR DEVICE, SURGISIS GOLD
HERNIA REPAIR GRAFT**Oct 13, 2006
32 days to decision

K062697 · Product code: FTL · General & Plastic Surgery

Source: <https://www.510kdatabase.net/k062697/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Mesh, Surgical, Polymeric (FTL)
Date received	Sep 11, 2006
Decision date	Oct 13, 2006
Days to decision	32 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Cook Biotech, Inc.
Location	West Lafayette, IN, US
Contact	MARY A FADERAN
510(k) history	31 submissions · 31 cleared · 1998-2014

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k062697/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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