

K974540 SIS HERNIA REPAIR DEVICEMay 20, 1998
168 days to decisionK974540 · Product code: **FTL** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k974540/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mesh, Surgical, Polymeric (FTL)
Date received	Dec 3, 1997
Decision date	May 20, 1998
Days to decision	168 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Sentron Medical, Inc.
Location	Cincinnati, OH, US
Contact	ROBERT J MORFF
510(k) history	1 submissions · 1 cleared · 1998-1998

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