

K961194 ANTIMICROBIAL MEDIASTINAL SILICONE DRAINAug 8, 1996
134 days to decisionK961194 · Product code: **GBW** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k961194/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Peritoneal (GBW)
Date received	Mar 27, 1996
Decision date	Aug 8, 1996
Days to decision	134 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Sil-Med Corp.
Location	Mchenry, IL, US
Contact	KAREN K SYLVIA
510(k) history	29 submissions · 28 cleared · 1980-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k961194/> Data sourced from FDA 510(k) public records (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. - Generated June 30, 2026