

K973273 XOMED T-STENT, FRONTAL SINUS STENTSep 29, 1997
27 days to decisionK973273 · Product code: **KAM** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k973273/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Sinus (KAM)
Date received	Sep 2, 1997
Decision date	Sep 29, 1997
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Xomed, Inc.
Location	Arnold, MO, US
Contact	DEBRA B CORTNER
510(k) history	82 submissions · 81 cleared · 1976-2000

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k973273/> 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 May 21, 2026