

K895670 SURGICAL SUCTION APPARATUSSep 27, 1989
8 days to decisionK895670 · Product code: **GEA** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k895670/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Surgical, General & Plastic Surgery (GEA)
Date received	Sep 19, 1989
Decision date	Sep 27, 1989
Days to decision	8 days
Third-party review	No

APPLICANT

Company	Baker-Sundt Research, Inc.
Location	Tucson, AZ, US
Contact	SUNDT, JR
510(k) history	1 submissions · 1 cleared · 1989-1989

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k895670/>, 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 July 7, 2026