

K002582 PILLING E-FRAZIER BIPOLAR SUCTION TUBENov 3, 2000
77 days to decisionK002582 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k002582/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Aug 18, 2000
Decision date	Nov 3, 2000
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Pilling
Location	Fort Washington, PA, US
Contact	ELIZABETH LAZARO
510(k) history	1 submissions · 1 cleared · 2000-2000

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k002582/> 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 3, 2026