FDA 510(k) Premarket Notification

K002582 PILLING E-FRAZIER BIPOLAR SUCTION TUBE

CLEARED

Nov 3, 2000
77 days to decision

K002582 · Product code: **GEI** · General & Plastic Surgery
Source: 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