FDA 510(k) Premarket Notification

K952973 THE TULIP MONO-POLAR CAUTERY SPATULA

CLEARED

Sep 11, 1995
76 days to decision

K952973 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k952973/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 27, 1995
Decision date	Sep 11, 1995
Days to decision	76 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**The Tulip Mfg. Co.**
Location	Mesa, AZ, US
Contact	RICHARD S HUNTER
510(k) history	24 submissions · 24 cleared · 1995-1996