FDA 510(k) Premarket Notification

K014260 TISSUELINK SOLID CYLINDER MONOPOLAR DEVICE

CLEARED

Mar 27, 2002
90 days to decision

K014260 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k014260/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Dec 27, 2001
Decision date	Mar 27, 2002
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Tissuelink Medical, Inc.**
Location	Dover, NH, US
Contact	VICKI ANASTASI
510(k) history	12 submissions · 12 cleared · 2000-2008