FDA 510(k) Premarket Notification

K030380 TISSUELINK SEALING FORCEPS, MODEL 1 21-202-1

CLEARED

Mar 3, 2003
26 days to decision

K030380 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k030380/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Feb 5, 2003
Decision date	Mar 3, 2003
Days to decision	26 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