

K030380 TISSUELINK SEALING FORCEPS, MODEL 1 21-202-1Mar 3, 2003
26 days to decisionK030380 · Product code: **GEI** · General & Plastic SurgerySource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k030380/> 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 June 26, 2026