

K061258 TRABECTOME HIGH FREQUENCY GENERATOR/LPMay 18, 2006
13 days to decisionK061258 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k061258/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	May 5, 2006
Decision date	May 18, 2006
Days to decision	13 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	Neomedix Corp.
Location	San Juan Capistrano, CA, US
Contact	THOMAS SCHROEDER
510(k) history	4 submissions · 4 cleared · 2003-2006

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k061258/> 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 30, 2026