

**K102339 CONMED LINVATEC SEQUENT MENISCAL REPAIR
DEVICE**Nov 23, 2010
97 days to decisionK102339 · Product code: **GAT** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k102339/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Suture, Nonabsorbable, Synthetic, Polyethylene (GAT)
Date received	Aug 18, 2010
Decision date	Nov 23, 2010
Days to decision	97 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Conmed Linvatec
Location	Largo, FL, US
Contact	JAN FLEGEAU
510(k) history	21 submissions · 21 cleared · 2005-2011

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k102339/> 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 May 14, 2026