FDA 510(k) Premarket Notification

K102339 CONMED LINVATEC SEQUENT MENISCAL REPAIR DEVICE

CLEARED

Nov 23, 2010
97 days to decision

K102339 · 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