FDA 510(k) Premarket Notification

K220186 Zone Specific AIM

CLEARED

May 13, 2022
109 days to decision

K220186 · Product code: **GAT** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k220186/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Suture, Nonabsorbable, Synthetic, Polyethylene (GAT)
Date received	Jan 24, 2022
Decision date	May 13, 2022
Days to decision	109 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Conmed Corporation**
Location	Utica, NY, US
Contact	Orjada Dervishleri
510(k) history	82 submissions · 82 cleared · 2004-2026