

**K220186 Zone Specific AIM**May 13, 2022  
109 days to decisionK220186 · Product code: **GAT** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k220186/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k220186/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 13, 2026