

**K082535 COVIDIEN SPORTS SURGERY AS MENISCAL REPAIR
DEVICE**Dec 1, 2008
90 days to decisionK082535 · Product code: **GAT** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k082535/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Suture, Nonabsorbable, Synthetic, Polyethylene (GAT)
Date received	Sep 2, 2008
Decision date	Dec 1, 2008
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Surgical Devices, A Global Business Unit Covidien
Location	North Haven, CT, US
Contact	TIM M LOHNES
510(k) history	3 submissions · 3 cleared · 2008-2009

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k082535/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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