

**K932513 LABELING FOR MYO/WIRE II STERNOTOMY SUTURES**

Jan 7, 1994  
227 days to decision

K932513 · Product code: **GAO** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k932513/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Suture, Nonabsorbable (GAO)
Date received	May 25, 1993
Decision date	Jan 7, 1994
Days to decision	227 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Alto Development Corp.</b>
Location	Mchenry, IL, US
Contact	TIM WOJCIECHOWICZ
510(k) history	10 submissions · 10 cleared · 1982-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k932513/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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