

**K012603 REPROCESSED AUTOSUTURE UNIPOLAR  
LAPAROSCOPIC/ENDOSCOPIC INSTRUMENTS**Nov 7, 2001  
86 days to decisionK012603 · Product code: **NUJ** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k012603/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation Accessories, Laparoscopic & Endoscopic, Reprocessed (NUJ)
Date received	Aug 13, 2001
Decision date	Nov 7, 2001
Days to decision	86 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Alliance Medical Corp.</b>
Location	Phoenix, AZ, US
Contact	DON SELVEY
510(k) history	36 submissions · 36 cleared · 2001-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k012603/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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