

**K093985 NIPRO SAFETOUCH LOCKTAIL SAFETY FISTULA
NEEDLE**Jan 22, 2010
29 days to decisionK093985 · Product code: **FIE** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k093985/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Needle, Fistula (FIE)
Date received	Dec 24, 2009
Decision date	Jan 22, 2010
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Nipro Medical Corp.
Location	Miami, FL, US
Contact	JESSICA OSWALD
510(k) history	27 submissions · 27 cleared · 1997-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k093985/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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