

**K063721 NIPRO SAFETOUCH II GAMMA**Jan 12, 2007  
28 days to decisionK063721 · Product code: **FIE** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k063721/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Needle, Fistula (FIE)
Date received	Dec 15, 2006
Decision date	Jan 12, 2007
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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

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