

**K020891 NAKAO SNARE II AND NAKAO SNARE III**Jun 17, 2002  
90 days to decisionK020891 · Product code: **FDI** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k020891/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Snare, Flexible (FDI)
Date received	Mar 19, 2002
Decision date	Jun 17, 2002
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Granit Medical Innovations</b>
Location	New York, NY, US
Contact	NAOMI L NAKAO
510(k) history	1 submissions · 1 cleared · 2002-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k020891/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 30, 2026