FDA 510(k) Premarket Notification

# K020891 NAKAO SNARE II AND NAKAO SNARE III

**CLEARED**

Jun 17, 2002
90 days to decision

K020891 · Product code: **FDI** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k020891/

## SUBMISSION DETAILS

| | |
|---|---|
| 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

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