FDA 510(k) Premarket Notification

# K030240 MODIFICATION TO HERMES OPERATING ROOM CONTROL CENTER

**CLEARED**

Feb 21, 2003
29 days to decision

K030240 · Product code: **GCJ** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k030240/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Special |
| Device classification | Laparoscope, General & Plastic Surgery (GCJ) |
| Date received | Jan 23, 2003 |
| Decision date | Feb 21, 2003 |
| Days to decision | 29 days |
| Third-party review | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **Computer Motion, Inc.** |
| Location | Washington, DC, US |
| Contact | CATHY STUPAK |
| 510(k) history | 26 submissions · 26 cleared · 1993-2003 |