

**K030240 MODIFICATION TO HERMES OPERATING ROOM
CONTROL CENTER**Feb 21, 2003
29 days to decisionK030240 · Product code: **GCJ** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k030240/>**SUBMISSION DETAILS**

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k030240/> Data sourced from FDA 510(k) public records (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. - Generated July 3, 2026