

**K003222 MODIFICATION TO HERMES OPERATING ROOM
CONTROL CENTER**Nov 3, 2000
18 days to decisionK003222 · Product code: **GCI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k003222/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Laryngoscope, Endoscope (GCI)
Date received	Oct 16, 2000
Decision date	Nov 3, 2000
Days to decision	18 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Computer Motion, Inc.
Location	Washington, DC, US
Contact	DAVID U THOMAS
510(k) history	26 submissions · 26 cleared · 1993-2003

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k003222/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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