FDA 510(k) Premarket Notification

K003222 MODIFICATION TO HERMES OPERATING ROOM CONTROL CENTER

CLEARED

Nov 3, 2000
18 days to decision

K003222 · Product code: **GCI** · General & Plastic Surgery
Source: 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