FDA 510(k) Premarket Notification

K991444 HERMES CONTROL CENTER, WITH CONTROL OF THE AESOP 3000HR (HERMES READY).

CLEARED

Jun 25, 1999
60 days to decision

K991444 · Product code: **GCJ** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k991444/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, General & Plastic Surgery (GCJ)
Date received	Apr 26, 1999
Decision date	Jun 25, 1999
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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