

K010552 CG-900P FETAL/MATERNAL MONITORMay 11, 2001
74 days to decisionK010552 · Product code: **HGM** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k010552/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Monitoring, Perinatal (HGM)
Date received	Feb 26, 2001
Decision date	May 11, 2001
Days to decision	74 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Card Guard Scientific Survival , Ltd.
Location	Israel, IL
Contact	LEONID TRACHTENBERG
510(k) history	49 submissions · 49 cleared · 1988-2011

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k010552/> 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 May 25, 2026