

K002805 KENDRO HERACELL CO2 INCUBATORNov 9, 2000
62 days to decisionK002805 · Product code: **MQG** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k002805/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessory, Assisted Reproduction (MQG)
Date received	Sep 8, 2000
Decision date	Nov 9, 2000
Days to decision	62 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Kendro Laboratory Products, L.P.
Location	Great Neck, NY, US
Contact	SUSAN D GOLDSTEIN-FALK
510(k) history	2 submissions · 2 cleared · 2000-2003

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k002805/> 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 June 29, 2026