

K020143 VASOVIEW 5 HARVESTING CANNULAFeb 20, 2002
35 days to decisionK020143 · Product code: **HET** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k020143/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, Gynecologic (and Accessories) (HET)
Date received	Jan 16, 2002
Decision date	Feb 20, 2002
Days to decision	35 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	Guidant Corporation, Cardiac Surgery
Location	S,Mta Clara, CA, US
Contact	ANNE SCHLAGENHAFT
510(k) history	8 submissions · 8 cleared · 2002-2008

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k020143/> 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 25, 2026