FDA 510(k) Premarket Notification

K020143 VASOVIEW 5 HARVESTING CANNULA

CLEARED

Feb 20, 2002
35 days to decision

K020143 · Product code: **HET** · Obstetrics & Gynecology
Source: 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