

**K052274 GUIDANT VASOVIEW HEMOPRO ENDOSCOPIC VESSEL HARVESTING SYSTEM, MODEL VH-3000**

Sep 21, 2005  
30 days to decision

K052274 · Product code: **GEI** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k052274/>

**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Aug 22, 2005
Decision date	Sep 21, 2005
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Guidant Corporation</b>
Location	St. Paul, MN, US
Contact	M. LAURIE WONG
510(k) history	15 submissions · 12 cleared · 2003-2006

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k052274/> 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 30, 2026