

**K912419 CORDGUARD BLOOD COLLECTION DEVICE**Oct 5, 1992  
493 days to decisionK912419 · Product code: **HFW** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k912419/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Clamp, Umbilical (HFW)
Date received	May 31, 1991
Decision date	Oct 5, 1992
Days to decision	493 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>O.B. Tech., Inc.</b>
Location	Huntington Beach, CA, US
Contact	STEPHEN R HESSEL
510(k) history	1 submissions · 1 cleared · 1992-1992

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k912419/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 30, 2026