

**K920353 ULTRACELL INSTRUMENT WIPE**Sep 15, 1992  
232 days to decisionK920353 · Product code: **HEP** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k920353/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Blood-flow, Ultrasonic (HEP)
Date received	Jan 27, 1992
Decision date	Sep 15, 1992
Days to decision	232 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Ultracell Medical Technologies, Inc.</b>
Location	North Stonington, CT, US
Contact	GEORGE P KORTEWEG
510(k) history	24 submissions · 24 cleared · 1992-2001

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k920353/> 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 29, 2026