

**K920152 BUBBLE TOCO**May 26, 1993  
499 days to decisionK920152 · Product code: **HFM** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k920152/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Uterine Contraction, External (for Use In Clinic) (HFM)
Date received	Jan 13, 1992
Decision date	May 26, 1993
Days to decision	499 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Utah Medical Products, Inc.</b>
Location	Mchenry, IL, US
Contact	EDWIN O GOODMAN
510(k) history	38 submissions · 38 cleared · 1979-2015

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k920152/> 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 May 23, 2026