FDA 510(k) Premarket Notification

K920152 BUBBLE TOCO

CLEARED

May 26, 1993
499 days to decision

K920152 · Product code: **HFM** · Obstetrics & Gynecology
Source: 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	**Utah Medical Products, Inc.**
Location	Mchenry, IL, US
Contact	EDWIN O GOODMAN
510(k) history	38 submissions · 38 cleared · 1979-2015