FDA 510(k) Premarket Notification

K902104 ORION WATCHGUARD

CLEARED

Oct 25, 1990
170 days to decision

K902104 · Product code: **HFM** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k902104/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Uterine Contraction, External (for Use In Clinic) (HFM)
Date received	May 8, 1990
Decision date	Oct 25, 1990
Days to decision	170 days
Third-party review	No

APPLICANT

Company	**Orion Industries**
Location	Fort Myers, FL, US
Contact	BOB GRANTHAM
510(k) history	1 submissions · 1 cleared · 1990-1990