FDA 510(k) Premarket Notification

K912138 HOMEVUE MONITORING SYS.MODEL TD 2000

CLEARED

Mar 19, 1992
310 days to decision

K912138 · Product code: **HGM** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k912138/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Monitoring, Perinatal (HGM)
Date received	May 14, 1991
Decision date	Mar 19, 1992
Days to decision	310 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Homevue Health Systems, Inc.**
Location	Laguna Beach, CA, US
Contact	GARY R.MOUNTS
510(k) history	2 submissions · 2 cleared · 1992-1995