FDA 510(k) Premarket Notification

K801747 TERUMO FETAL MONITOR # FHM-602

CLEARED

Aug 13, 1980
16 days to decision

K801747 · Product code: **HGM** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k801747/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Monitoring, Perinatal (HGM)
Date received	Jul 28, 1980
Decision date	Aug 13, 1980
Days to decision	16 days
Third-party review	No

APPLICANT

Company	**Terumo Medical Corp.**
Location	Elkton, MD, US
510(k) history	143 submissions · 143 cleared · 1980-2011