

K801747 TERUMO FETAL MONITOR # FHM-602Aug 13, 1980
16 days to decisionK801747 · Product code: **HGM** · Obstetrics & GynecologySource: <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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k801747/> Data sourced from FDA 510(k) public records (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. - Generated May 27, 2026