FDA 510(k) Premarket Notification

K823081 M2100 ANTEPARTUM FETAL MONITOR

CLEARED

Jan 26, 1983
99 days to decision

K823081 · Product code: **HGM** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k823081/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Monitoring, Perinatal (HGM)
Date received	Oct 19, 1982
Decision date	Jan 26, 1983
Days to decision	99 days
Third-party review	No

APPLICANT

Company	**Huntleigh Technology, Inc.**
Location	Walker, MI, US
510(k) history	23 submissions · 23 cleared · 1981-1999