FDA 510(k) Premarket Notification

K823188 M2000 INTRAPARTUM FETAL MONITOR

CLEARED

Jan 26, 1983
92 days to decision

K823188 · Product code: **HGM** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k823188/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Monitoring, Perinatal (HGM)
Date received	Oct 26, 1982
Decision date	Jan 26, 1983
Days to decision	92 days
Third-party review	No

APPLICANT

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