

K823081 M2100 ANTEPARTUM FETAL MONITORJan 26, 1983
99 days to decisionK823081 · Product code: **HGM** · Obstetrics & GynecologySource: <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

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