FDA 510(k) Premarket Notification

K032252 MODEL 120 SERIES MATERNAL/FETAL MONITOR

CLEARED

Aug 21, 2003
30 days to decision

K032252 · Product code: **HGM** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k032252/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Monitoring, Perinatal (HGM)
Date received	Jul 22, 2003
Decision date	Aug 21, 2003
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Ge Medical Systems Information Technologies**
Location	Mchenry, IL, US
Contact	MELISSA ROBINSON
510(k) history	136 submissions · 132 cleared · 1978-2012