FDA 510(k) Premarket Notification

K812256 MODEL AP 180 MINI-MONITOR

CLEARED

Oct 20, 1981
69 days to decision

K812256 · Product code: **HGM** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k812256/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Monitoring, Perinatal (HGM)
Date received	Aug 12, 1981
Decision date	Oct 20, 1981
Days to decision	69 days
Third-party review	No

APPLICANT

Company	**Sonicaid, Inc.**
Location	Mchenry, IL, US
510(k) history	18 submissions · 17 cleared · 1977-1988