FDA 510(k) Premarket Notification

K802024 GRAVIDAPHONE

CLEARED

Jan 7, 1981
140 days to decision

K802024 · Product code: **KNG** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k802024/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Ultrasonic, Fetal (KNG)
Date received	Aug 20, 1980
Decision date	Jan 7, 1981
Days to decision	140 days
Third-party review	No

APPLICANT

Company	**Solar Electronics**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981