

K802024 GRAVIDAPHONEJan 7, 1981
140 days to decisionK802024 · Product code: **KNG** · Obstetrics & GynecologySource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k802024/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 30, 2026