FDA 510(k) Premarket Notification

K812868 FETAL PULSE DETECTORS, F50/F60

CLEARED

Nov 25, 1981
42 days to decision

K812868 · Product code: **KNG** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k812868/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Ultrasonic, Fetal (KNG)
Date received	Oct 14, 1981
Decision date	Nov 25, 1981
Days to decision	42 days
Third-party review	No

APPLICANT

Company	**South Carolina Medical, Inc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981