FDA 510(k) Premarket Notification

K800095 MGD-3

CLEARED

Feb 11, 1980
28 days to decision

K800095 · Product code: **KNG** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k800095/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Ultrasonic, Fetal (KNG)
Date received	Jan 14, 1980
Decision date	Feb 11, 1980
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Kt Medical, Inc.**
Location	Mchenry, IL, US
510(k) history	13 submissions · 13 cleared · 1980-1980