

K800098 MINIFETON 200Feb 11, 1980
28 days to decisionK800098 · Product code: **KNG** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k800098/>**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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k800098/> 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 May 22, 2026