

K890139 TOITU DOPPLER FETUS DETECTOR, FD-400DMar 29, 1989
75 days to decisionK890139 · Product code: **KNG** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k890139/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Ultrasonic, Fetal (KNG)
Date received	Jan 13, 1989
Decision date	Mar 29, 1989
Days to decision	75 days
Third-party review	No

APPLICANT

Company	Toitu of America, Inc.
Location	Wayne, PA, US
Contact	A OGLETREE
510(k) history	2 submissions · 2 cleared · 1989-1989

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k890139/>; 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