

K894036 TOITU FETAL ACTOCARDIOGRAPH, MT-332Nov 6, 1989
153 days to decisionK894036 · Product code: **HGM** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k894036/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Monitoring, Perinatal (HGM)
Date received	Jun 6, 1989
Decision date	Nov 6, 1989
Days to decision	153 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/k894036/>; 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