FDA 510(k) Premarket Notification

K894036 TOITU FETAL ACTOCARDIOGRAPH, MT-332

CLEARED

Nov 6, 1989
153 days to decision

K894036 · Product code: **HGM** · Obstetrics & Gynecology
Source: 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