FDA 510(k) Premarket Notification

K854267 HUNTLEIGH DOPCORD, D140

CLEARED

Jul 9, 1986
259 days to decision

K854267 · Product code: **KNG** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k854267/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Ultrasonic, Fetal (KNG)
Date received	Oct 23, 1985
Decision date	Jul 9, 1986
Days to decision	259 days
Third-party review	No

APPLICANT

Company	**Huntleigh Technology, Inc.**
Location	Walker, MI, US
Contact	JAMES BRITTON
510(k) history	23 submissions · 23 cleared · 1981-1999