FDA 510(k) Premarket Notification

K012003 DRG REACTION CHAMBER/SAFETY TIP

CLEARED

Feb 12, 2002
230 days to decision

K012003 · Product code: **HEI** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k012003/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Heart-valve Movement, Fetal, Ultrasonic (HEI)
Date received	Jun 27, 2001
Decision date	Feb 12, 2002
Days to decision	230 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Doctor'S Research Group, Inc.**
Location	Wolcott, CT, US
Contact	RICHARD DESLAURIERS
510(k) history	13 submissions · 13 cleared · 1998-2005