FDA 510(k) Premarket Notification

K902190 FETAL PULSE DETECTOR MODEL DOP-1

CLEARED

Oct 22, 1990
159 days to decision

K902190 · Product code: **HEK** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k902190/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Heart Sound, Fetal, Ultrasonic (HEK)
Date received	May 16, 1990
Decision date	Oct 22, 1990
Days to decision	159 days
Third-party review	No

APPLICANT

Company	**Advanced Medical Systems, Inc.**
Location	Hamden, CT, US
Contact	ANTHONY CALDERONI
510(k) history	10 submissions · 10 cleared · 1988-1996