FDA 510(k) Premarket Notification

K883806 ROCKET INTRAUTERINE SOUND

CLEARED

Oct 31, 1988
54 days to decision

K883806 · Product code: **HHM** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k883806/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sound, Uterine (HHM)
Date received	Sep 7, 1988
Decision date	Oct 31, 1988
Days to decision	54 days
Third-party review	No

APPLICANT

Company	**A & A Medical, Inc.**
Location	Branford, CT, US
Contact	K BERNBERG
510(k) history	23 submissions · 23 cleared · 1988-2001