FDA 510(k) Premarket Notification

K890379 UTERINE TENTACULUM

CLEARED

Feb 13, 1989
21 days to decision

K890379 · Product code: **HDC** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k890379/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tenaculum, Uterine (HDC)
Date received	Jan 23, 1989
Decision date	Feb 13, 1989
Days to decision	21 days
Third-party review	No

APPLICANT

Company	**Kinetic Medical Products**
Location	Erie, PA, US
Contact	JAMES I LAUGHNER
510(k) history	63 submissions · 63 cleared · 1989-1989