FDA 510(k) Premarket Notification

K890328 ADULT RESECTOSCOPE

CLEARED

Feb 6, 1990
379 days to decision

K890328 · Product code: **HIH** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k890328/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hysteroscope (and Accessories) (HIH)
Date received	Jan 23, 1989
Decision date	Feb 6, 1990
Days to decision	379 days
Third-party review	No

APPLICANT

Company	**Circon Video**
Location	Stanford, CT, US
Contact	TAYLOR, PE
510(k) history	14 submissions · 14 cleared · 1990-2002