

K890328 ADULT RESECTOSCOPEFeb 6, 1990
379 days to decisionK890328 · Product code: **HIH** · Obstetrics & GynecologySource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k890328/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 29, 2026