

K001314 MEDSCOPEJul 25, 2000
90 days to decisionK001314 · Product code: **HEX** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k001314/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Colposcope (and Colpomicroscope) (HEX)
Date received	Apr 26, 2000
Decision date	Jul 25, 2000
Days to decision	90 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	All-Pro Imaging Corp.
Location	Hicksville, NY, US
Contact	JOSEPH CAREY
510(k) history	4 submissions · 4 cleared · 1991-2000

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k001314/> 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 May 19, 2026