FDA 510(k) Premarket Notification

K001314 MEDSCOPE

CLEARED

Jul 25, 2000
90 days to decision

K001314 · Product code: **HEX** · Obstetrics & Gynecology
Source: 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