FDA 510(k) Premarket Notification

K940718 INNERVISION INC. LAPAROSCOPIC HAND INSTRUMENTS

CLEARED

Jul 22, 1994
154 days to decision

K940718 · Product code: **HET** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k940718/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, Gynecologic (and Accessories) (HET)
Date received	Feb 18, 1994
Decision date	Jul 22, 1994
Days to decision	154 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Innervision, Inc.**
Location	Memphis, TN, US
Contact	FRANK M LEWIS
510(k) history	6 submissions · 6 cleared · 1994-1999