FDA 510(k) Premarket Notification

K940750 INNERVISION,INC. LAPAROSCOPIC ELECTRODES

CLEARED

Aug 23, 1994
186 days to decision

K940750 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k940750/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Feb 18, 1994
Decision date	Aug 23, 1994
Days to decision	186 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