FDA 510(k) Premarket Notification

K962119 RE-NEW LAPAROSCOPIC INSTRUMENTS

CLEARED

Aug 9, 1996
70 days to decision

K962119 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k962119/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	May 31, 1996
Decision date	Aug 9, 1996
Days to decision	70 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Microline Pentax, Inc.**
Location	Devers, MA, US
Contact	HUGHES DE LAFORCADE
510(k) history	12 submissions · 12 cleared · 1988-2002