FDA 510(k) Premarket Notification

K901718 KLEIN LAMPREY-CANNULA(TM)

CLEARED

May 24, 1990
41 days to decision

K901718 · Product code: **GEA** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k901718/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Surgical, General & Plastic Surgery (GEA)
Date received	Apr 13, 1990
Decision date	May 24, 1990
Days to decision	41 days
Third-party review	No

APPLICANT

Company	**Jeff Klein Surgical, Inc.**
Location	San Clemente, CA, US
Contact	JACQUELINE KLEIN
510(k) history	5 submissions · 5 cleared · 1990-1990