FDA 510(k) Premarket Notification

K991592 KSEA DE LA PLAZA BLEPHAROPLASTY SET

CLEARED

Nov 2, 1999
179 days to decision

K991592 · Product code: **HNI** · Ophthalmic
Source: https://www.510kdatabase.net/k991592/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Retractor, Ophthalmic (HNI)
Date received	May 7, 1999
Decision date	Nov 2, 1999
Days to decision	179 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**KARL STORZ Endoscopy-America, Inc.**
Location	Mchenry, IL, US
Contact	KEVIN KENNAN
510(k) history	361 submissions · 361 cleared · 1980-2024