

K991592 KSEA DE LA PLAZA BLEPHAROPLASTY SETNov 2, 1999
179 days to decisionK991592 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k991592/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 15, 2026