

K950282 KARL STORZ MODIFIED VERESS NEEDLE AND CANNULAApr 5, 1995
71 days to decisionK950282 · Product code: **KOC** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k950282/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Blood Circuit, Hemodialysis (KOC)
Date received	Jan 24, 1995
Decision date	Apr 5, 1995
Days to decision	71 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k950282/> 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