FDA 510(k) Premarket Notification

K950282 KARL STORZ MODIFIED VERESS NEEDLE AND CANNULA

CLEARED

Apr 5, 1995
71 days to decision

K950282 · 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