FDA 510(k) Premarket Notification

CLEARED

K232370 Percutaneous Nephroscope System

May 1, 2024
267 days to decision

K232370 · Product code: **FGA** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k232370/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Nephroscope (FGA)
Date received	Aug 8, 2023
Decision date	May 1, 2024
Days to decision	267 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Karl Storz SE & CO. KG**
Location	Tuttlingen, DE
Contact	Alita McElroy
Website	https://www.karlstorz.com
510(k) history	23 submissions · 23 cleared · 2018-2026

Karl Storz SE & CO. KG is a medical device manufacturer headquartered in Tuttlingen, Germany. The company specializes in endoscopic instruments and visualization systems for surgical and diagnostic procedures. The company has received FDA 510(k) clearances from total submissions since 2018. Karl Storz devices span multiple surgical specialties, with particular strength in Gastroenterology & Urology applications. The latest FDA 510(k) clearance was granted in 2026, confirming the company's active regulatory engagement. Recent cleared devices include flexible video endoscop...