

K171603 Cope Nephroureterostomy Stent and Amplatz Ureteral Stent SetApr 16, 2018
319 days to decisionK171603 · Product code: **FAD** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k171603/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - K
Submission type	Traditional
Device classification	Stent, Ureteral (FAD)
Date received	Jun 1, 2017
Decision date	Apr 16, 2018
Days to decision	319 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Cook Incorporated
Location	Bloomington, IN, US
Contact	Erum B. Nasir
510(k) history	175 submissions · 153 cleared · 2006-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k171603/> 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 30, 2026