FDA 510(k) Premarket Notification

K840059 SILITEK URINARY DIVERSION STENT

CLEARED

Feb 4, 1984
26 days to decision

K840059 · Product code: **FAD** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k840059/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stent, Ureteral (FAD)
Date received	Jan 9, 1984
Decision date	Feb 4, 1984
Days to decision	26 days
Third-party review	No

APPLICANT

Company	**Surgitek**
Location	Mchenry, IL, US
510(k) history	29 submissions · 28 cleared · 1983-1995