FDA 510(k) Premarket Notification

K833782 SURGITEK ALL SILICONE INFECTION DOUBJ

CLEARED

Apr 2, 1984
157 days to decision

K833782 · Product code: **FAD** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k833782/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stent, Ureteral (FAD)
Date received	Oct 28, 1983
Decision date	Apr 2, 1984
Days to decision	157 days
Third-party review	No

APPLICANT

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