FDA 510(k) Premarket Notification

K821776 SURGITEK PERCUTANEOUS NEPHROST. STENT

CLEARED

Aug 10, 1982
54 days to decision

K821776 · Product code: **LJE** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k821776/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Nephrostomy (LJE)
Date received	Jun 17, 1982
Decision date	Aug 10, 1982
Days to decision	54 days
Third-party review	No

APPLICANT

Company	**Medical Engineering Corp.**
Location	Mchenry, IL, US
510(k) history	28 submissions · 28 cleared · 1977-1993