FDA 510(k) Premarket Notification

K810368 VANCE PERCUTANEOUS MALECOT NEPHROSTOMY

CLEARED

Mar 20, 1981
36 days to decision

K810368 · Product code: **KOB** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k810368/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Suprapubic (and Accessories) (KOB)
Date received	Feb 12, 1981
Decision date	Mar 20, 1981
Days to decision	36 days
Third-party review	No

APPLICANT

Company	**Vance Products, Inc.**
Location	Mchenry, IL, US
510(k) history	17 submissions · 17 cleared · 1978-1982