FDA 510(k) Premarket Notification

K811833 VNACE FASCIAL DILATOR SET

CLEARED

Jul 10, 1981
10 days to decision

K811833 · Product code: **KOE** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k811833/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Urethral (KOE)
Date received	Jun 30, 1981
Decision date	Jul 10, 1981
Days to decision	10 days
Third-party review	No

APPLICANT

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