FDA 510(k) Premarket Notification

K792220 VANCE KIDNEY BIOPSY BRUSHES

CLEARED

Dec 6, 1979
31 days to decision

K792220 · Product code: **KOD** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k792220/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Urological (KOD)
Date received	Nov 5, 1979
Decision date	Dec 6, 1979
Days to decision	31 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k792220/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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