

K810371 VANCE URETHAL DILATOR SETMar 20, 1981
36 days to decisionK810371 · Product code: **KOE** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k810371/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Urethral (KOE)
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

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Device record: <https://www.510kdatabase.net/k810371/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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