

**K811833 VNACE FASCIAL DILATOR SET**Jul 10, 1981  
10 days to decisionK811833 · Product code: **KOE** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k811833/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k811833/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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