

K780415 VIVONEX DELIVERY SYSTEMApr 10, 1978
27 days to decisionK780415 · Product code: **FPD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k780415/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tube, Feeding (FPD)
Date received	Mar 14, 1978
Decision date	Apr 10, 1978
Days to decision	27 days
Third-party review	No

APPLICANT

Company	Norwich Pharmacal Co.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1978-1978

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Device record: <https://www.510kdatabase.net/k780415/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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