

K812567 RELIEFNov 10, 1981
62 days to decisionK812567 · Product code: **LKX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k812567/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Thermal, Hemorrhoids (LKX)
Date received	Sep 9, 1981
Decision date	Nov 10, 1981
Days to decision	62 days
Third-party review	No

APPLICANT

Company	American Pharmaceutical Co., Inc.
Location	Walker, MI, US
510(k) history	2 submissions · 2 cleared · 1981-1995

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Device record: <https://www.510kdatabase.net/k812567/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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