

K822217 THERMOTHERAPYSep 24, 1982
60 days to decisionK822217 · Product code: **LKX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k822217/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Thermal, Hemorrhoids (LKX)
Date received	Jul 26, 1982
Decision date	Sep 24, 1982
Days to decision	60 days
Third-party review	No

APPLICANT

Company	Luther Medical Products, Inc.
Location	Walker, MI, US
510(k) history	17 submissions · 16 cleared · 1980-1998

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Device record: <https://www.510kdatabase.net/k822217/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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