FDA 510(k) Premarket Notification

K822217 THERMOTHERAPY

CLEARED

Sep 24, 1982
60 days to decision

K822217 · Product code: **LKX** · Gastroenterology & Urology
Source: 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