FDA 510(k) Premarket Notification

K822649 DEVICE FOR THE TREATMENT OF HEMORRHOIDS

CLEARED

Nov 3, 1982
63 days to decision

K822649 · Product code: **LKX** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k822649/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Thermal, Hemorrhoids (LKX)
Date received	Sep 1, 1982
Decision date	Nov 3, 1982
Days to decision	63 days
Third-party review	No

APPLICANT

Company	**Dunmore Corp.**
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1982-1982