

**K822649 DEVICE FOR THE TREATMENT OF HEMORRHOIDS**Nov 3, 1982  
63 days to decisionK822649 · Product code: **LKX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k822649/>**SUBMISSION DETAILS**

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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**

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

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k822649/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 3, 2026