

**K944874 ZEROID HEMORRHOIDAL DEVICE**Aug 7, 1995  
308 days to decisionK944874 · Product code: **LKX** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k944874/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Thermal, Hemorrhoids (LKX)
Date received	Oct 3, 1994
Decision date	Aug 7, 1995
Days to decision	308 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Medical Appliance Research Corp.</b>
Location	North Miami Beach, FL, US
Contact	MURRAY CERAMI
510(k) history	1 submissions · 1 cleared · 1995-1995

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k944874/> 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 1, 2026