

**K022145 OVER THE WIRE EMBOLECTOMY CATHETER, MODEL
2302**Jan 14, 2003
196 days to decisionK022145 · Product code: **DXE** · Cardiovascular
Source: <https://www.510kdatabase.net/k022145/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Embolectomy (DXE)
Date received	Jul 2, 2002
Decision date	Jan 14, 2003
Days to decision	196 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	LeMaitre Vascular, Inc.
Location	Burlington, MA, US
Contact	JAMES ASHBY
510(k) history	32 submissions · 32 cleared · 2003-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k022145/> Data sourced from FDA 510(k) public records (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. - Generated May 15, 2026