

**K122543 AZUR PURE PERIPHERAL COIL SYSTEM, PUSHABLE
18 AZUR PURE PERIPHERAL COIL SYSTEM, PUSHABLE 35**Nov 28, 2012
99 days to decisionK122543 · Product code: **KRD** · Cardiovascular
Source: <https://www.510kdatabase.net/k122543/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Vascular, For Promoting Embolization (KRD)
Date received	Aug 21, 2012
Decision date	Nov 28, 2012
Days to decision	99 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	MicroVention, Inc.
Location	Aliso Viejo, CA, US
Contact	LARAIN PANGELINA
510(k) history	85 submissions · 85 cleared · 2001-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k122543/> 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 June 9, 2026