

**K242612 Option ELITE Vena Cava Filter System (352506070E
352506100E)**Sep 10, 2024
7 days to decisionK242612 · Product code: **DTK** · Cardiovascular
Source: <https://www.510kdatabase.net/k242612/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Filter, Intravascular, Cardiovascular (DTK)
Date received	Sep 3, 2024
Decision date	Sep 10, 2024
Days to decision	7 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Argon Medical Devices, Inc.
Location	Athens, TX, US
Contact	Scott Bishop
Website	https://www.argonmedical.com
510(k) history	20 submissions · 20 cleared · 2007-2026

Argon Medical Devices, Inc. manufactures specialty interventional medical devices. The company is based in Athens, US. Argon has received FDA 510(k) clearances from total submissions since 2007. The company specializes in Cardiovascular devices, which represent the dominant focus of its regulatory portfolio. The latest clearance was in 2026, reflecting continued active development and market engagement. Argon's cleared device portfolio includes thrombectomy systems, vascular access devices, biopsy instruments, and retrieval kits. The company also offers custom product sol...