California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Inari Medical - Oak Canyon Artix MT Thrombectomy Device

Recall Date	Product Description	Recalling Firm	Recall Reason
05/06/2025	Artix MT Thrombectomy Device REF: 32-102	Inari Medical - Oak Canyon	Thrombectomy device specifies vessel range of 2.5-6 mm on the pouch, carton, and Instructions for Use rather than the correct vessel range of 3-6 mm, which could incrementally increase the radial force on the artery, which has the potential of vascular complications.

Recall Class	Product Identification	Distribution	Affected Dates
II	UDI-DI: 00810123710584 Lot: 24100082 Expiration: 14 April 2025 IFU: IU-01087 Rev. B	8 units in California	April and prior

For additional information, please visit the FDA Website.