FDA 510(k) Premarket Notification

K233069 Removal System Large Bore 60 cc Syringe

CLEARED

Mar 26, 2024
182 days to decision

K233069 · Product code: **PUR** · General Hospital
Source: https://www.510kdatabase.net/k233069/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Vacuum Syringe (PUR)
Date received	Sep 26, 2023
Decision date	Mar 26, 2024
Days to decision	182 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Inari Medical, Inc.**
Location	Aliso Viejo, CA, US
Contact	Kaitlyn Weinkauf
510(k) history	30 submissions · 30 cleared · 2015-2025