

K233069 Removal System Large Bore 60 cc SyringeMar 26, 2024
182 days to decisionK233069 · 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 Weinkauff
510(k) history	30 submissions · 30 cleared · 2015-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k233069/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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