FDA 510(k) Premarket Notification

K222083 LimFlow V-Ceiver

CLEARED

Aug 9, 2022
25 days to decision

K222083 · Product code: **MMX** · Cardiovascular
Source: https://www.510kdatabase.net/k222083/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Percutaneous Retrieval (MMX)
Date received	Jul 15, 2022
Decision date	Aug 9, 2022
Days to decision	25 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**LimFlow, Inc.**
Location	San Jose, CA, US
Contact	Zachary Woodson
510(k) history	6 submissions · 6 cleared · 2022-2026