

K222083 LimFlow V-CeiverAug 9, 2022
25 days to decisionK222083 · Product code: **MMX** · CardiovascularSource: <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

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