

K212988 ONO Retrieval DeviceMay 23, 2022
245 days to decisionK212988 · Product code: **MMX** · CardiovascularSource: <https://www.510kdatabase.net/k212988/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Percutaneous Retrieval (MMX)
Date received	Sep 20, 2021
Decision date	May 23, 2022
Days to decision	245 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Onocor Vascular, LLC
Location	West Chester, PA, US
Contact	Tiffini Wittwer
510(k) history	1 submissions · 1 cleared · 2022-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k212988/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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