

K222254 Gunther Tulip Vena Cava Filter Retrieval SetMay 16, 2023
293 days to decisionK222254 · Product code: **MMX** · CardiovascularSource: <https://www.510kdatabase.net/k222254/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Percutaneous Retrieval (MMX)
Date received	Jul 27, 2022
Decision date	May 16, 2023
Days to decision	293 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Cook Incorporated
Location	Bloomington, IN, US
Contact	Daniel Corbin
510(k) history	175 submissions · 153 cleared · 2006-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k222254/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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