

K212894 PhasTIPPMar 31, 2022
202 days to decisionK212894 · Product code: **DWQ** · CardiovascularSource: <https://www.510kdatabase.net/k212894/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stripper, Vein, External (DWQ)
Date received	Sep 10, 2021
Decision date	Mar 31, 2022
Days to decision	202 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	LeMaitre Vascular, Inc.
Location	Burlington, MA, US
Contact	Anurag Gadgil
510(k) history	32 submissions · 32 cleared · 2003-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k212894/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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