

K150262 Crux Vena Cava Filter System, femoralApr 3, 2015
58 days to decisionK150262 · Product code: **DTK** · CardiovascularSource: <https://www.510kdatabase.net/k150262/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filter, Intravascular, Cardiovascular (DTK)
Date received	Feb 4, 2015
Decision date	Apr 3, 2015
Days to decision	58 days
Third-party review	No
Summary / Statement	Summary
Other names	Crux Vena Cava Filter System, jugular

APPLICANT

Company	Volcano Corporation
Location	Rancho Cordova,, CA, US
Contact	Elaine Alan
510(k) history	55 submissions · 55 cleared · 2005-2021

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k150262/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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