FDA 510(k) Premarket Notification

K150262 Crux Vena Cava Filter System, femoral

CLEARED

Apr 3, 2015
58 days to decision

K150262 · Product code: **DTK** · Cardiovascular
Source: 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