

**K130843 SAFESEPT NEEDLE FREE TRANSSEPTAL TROCAR
GUIDEWIRE**Jan 8, 2014
287 days to decisionK130843 · Product code: **DRC** · Cardiovascular
Source: <https://www.510kdatabase.net/k130843/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Trocar (DRC)
Date received	Mar 27, 2013
Decision date	Jan 8, 2014
Days to decision	287 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Oscor, Inc.
Location	Palm Harbor, FL, US
Contact	MILA DOSKOCIL
510(k) history	49 submissions · 46 cleared · 1979-2021

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k130843/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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