

K133843 SABER PERCUTANEOUS TRANSLUMINAL ANGIOPLASTY DILATATION CATHETERJun 27, 2014
191 days to decisionK133843 · Product code: LIT · Cardiovascular
Source: <https://www.510kdatabase.net/k133843/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Angioplasty, Peripheral, Transluminal (LIT)
Date received	Dec 18, 2013
Decision date	Jun 27, 2014
Days to decision	191 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Cordis Corporation
Location	Warren, NJ, US
Contact	KIM FONDA
510(k) history	13 submissions · 12 cleared · 2004-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k133843/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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