FDA 510(k) Premarket Notification

K143652 Vector PTA Balloon Dilatation Catheter

CLEARED

Feb 26, 2015
65 days to decision

K143652 · Product code: **LIT** · Cardiovascular
Source: https://www.510kdatabase.net/k143652/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Angioplasty, Peripheral, Transluminal (LIT)
Date received	Dec 23, 2014
Decision date	Feb 26, 2015
Days to decision	65 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Vector Corporation, LLC**
Location	New York, NY, US
Contact	Jay Sturm
510(k) history	1 submissions · 1 cleared · 2015-2015