

**K143652 Vector PTA Balloon Dilatation Catheter**Feb 26, 2015  
65 days to decisionK143652 · 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	<b>Vector Corporation, LLC</b>
Location	New York, NY, US
Contact	Jay Sturm
510(k) history	1 submissions · 1 cleared · 2015-2015

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k143652/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 5, 2026