

**K052514 CLIRPATH TURBO PLUS EXCIMER LASER
CATHETERS, MODELS 314-151, 314-159, 317-152, 317-156,
320-006, 320-159, 323-001**

Oct 19, 2005
35 days to decision

K052514 · Product code: **PDU** · Cardiovascular
Source: <https://www.510kdatabase.net/k052514/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter For Crossing Total Occlusions (PDU)
Date received	Sep 14, 2005
Decision date	Oct 19, 2005
Days to decision	35 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Spectranetics Corp.
Location	Colorado Springs, CO, US
Contact	NEIL BURRIS
510(k) history	24 submissions · 24 cleared · 1999-2014

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k052514/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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