

**K182937 Upstream GoBack Crossing Catheter**May 23, 2019  
213 days to decisionK182937 · Product code: **DQY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k182937/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Oct 22, 2018
Decision date	May 23, 2019
Days to decision	213 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Upstream Peripheral Technologies, Ltd.</b>
Location	Caesarea, IL
Contact	Dan Rottenberg
510(k) history	2 submissions · 2 cleared · 2019-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k182937/> 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 May 27, 2026