

**K211802 GoBack Crossing Catheter**Mar 1, 2022  
264 days to decisionK211802 · Product code: **PDU** · CardiovascularSource: <https://www.510kdatabase.net/k211802/>**SUBMISSION DETAILS**

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|                       |  |
|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)           |
| Submission type       | Traditional                                  |
| Device classification | Catheter For Crossing Total Occlusions (PDU) |
| Date received         | Jun 10, 2021                                 |
| Decision date         | Mar 1, 2022                                  |
| Days to decision      | 264 days                                     |
| Third-party review    | No   |
| Combination product   | No   |
| PCCP authorized       | 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         |

**REGULATORY CONSULTANT**

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|-----------------|-----------------------------|
| Consulting firm | <b>Hogan Lovells US LLP</b> |
| Contact         | Janice M Hogan              |

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k211802/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 27, 2026