

**K203262 Cannula for Organ Perfusion**Jul 1, 2021  
238 days to decisionK203262 · Product code: **KDN** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k203262/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Perfusion, Kidney (KDN)
Date received	Nov 5, 2020
Decision date	Jul 1, 2021
Days to decision	238 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Bridge TO Life, Ltd.</b>
Location	Columbia, SC, US
Contact	Brett Schultz
510(k) history	4 submissions · 3 cleared · 2019-2026

**REGULATORY CONSULTANT**

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Consulting firm	<b>Wood Burditt Group</b>
Contact	H Carl Jenkins

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k203262/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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