

**K220632 Boss Crossing Support Catheter**Jun 2, 2022  
90 days to decisionK220632 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k220632/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Mar 4, 2022
Decision date	Jun 2, 2022
Days to decision	90 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Marvao Medical Devices, Ltd.</b>
Location	Northborough, MA, US
Contact	Chris Davey
510(k) history	9 submissions · 4 cleared · 2011-2022

**REGULATORY CONSULTANT**

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Consulting firm	<b>Avania, Ltd.</b>
Contact	Mary P. LeGraw

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

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