

**K220699 Prestige Coil System**Apr 8, 2022  
29 days to decisionK220699 · Product code: **KRD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k220699/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Device, Vascular, For Promoting Embolization (KRD)
Date received	Mar 10, 2022
Decision date	Apr 8, 2022
Days to decision	29 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Bait USA, LLC</b>
Location	Irvine, CA, US
Contact	Michael Peters
510(k) history	1 submissions · 1 cleared · 2022-2022

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