

**K200205 Reprocessed Advisor FL Circular Mapping Catheter, Sensor Enabled**

May 13, 2020  
106 days to decision

K200205 · Product code: **NLH** · Cardiovascular  
Source: <https://www.510kdatabase.net/k200205/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Recording, Electrode, Reprocessed (NLH)
Date received	Jan 28, 2020
Decision date	May 13, 2020
Days to decision	106 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Stryker Sustainability Solutions</b>
Location	Tempe, AZ, US
Contact	Mia McCorkel
510(k) history	31 submissions · 31 cleared · 2011-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k200205/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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