

**K222690 SmartGUIDE deflectable hydrophilic guidewire**Apr 24, 2023  
230 days to decisionK222690 · Product code: **MOF** · Neurology  
Source: <https://www.510kdatabase.net/k222690/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Guide, Wire, Catheter, Neurovasculature (MOF)
Date received	Sep 6, 2022
Decision date	Apr 24, 2023
Days to decision	230 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Artiria Medical SA</b>
Location	Geneva, CH
Contact	Guillaume Petit-Pierre
510(k) history	1 submissions · 1 cleared · 2023-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k222690/> 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