

**K110419 AVS ARIA NEUROMONITORING PROBE AND AVS  
ARIA PROBE DILATORS**Jul 28, 2011  
164 days to decisionK110419 · Product code: **GXZ** · Neurology  
Source: <https://www.510kdatabase.net/k110419/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Needle (GXZ)
Date received	Feb 14, 2011
Decision date	Jul 28, 2011
Days to decision	164 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Stryker Spine</b>
Location	Allendale, NJ, US
Contact	TIFFANI D ROGERS
510(k) history	74 submissions · 73 cleared · 2004-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k110419/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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