

**K063872 KINESIA**Apr 6, 2007  
98 days to decisionK063872 · Product code: **GYD** · Neurology  
Source: <https://www.510kdatabase.net/k063872/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Tremor (GYD)
Date received	Dec 29, 2006
Decision date	Apr 6, 2007
Days to decision	98 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cleveland Medical Devices, Inc.</b>
Location	Cleveland, OH, US
Contact	JOSEPH GIUFFRIDA
510(k) history	8 submissions · 8 cleared · 1995-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k063872/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 21, 2026