

**K030517 BALL RECORDING ELECTRODE/STIMULATION  
PROBE**Aug 1, 2003  
163 days to decisionK030517 · Product code: **GZK** · Neurology  
Source: <https://www.510kdatabase.net/k030517/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Electrode, Nasopharyngeal (GZK)
Date received	Feb 19, 2003
Decision date	Aug 1, 2003
Days to decision	163 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Viasys Healthcare, Inc.</b>
Location	Stoughton, WI, US
Contact	Gary Syring
510(k) history	3 submissions · 3 cleared · 2003-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k030517/>; 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 18, 2026