

**K861031 WYLER SPHENOIDAL ELECTRODE**Aug 25, 1986  
160 days to decisionK861031 · Product code: **GZL** · Neurology  
Source: <https://www.510kdatabase.net/k861031/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Depth (GZL)
Date received	Mar 18, 1986
Decision date	Aug 25, 1986
Days to decision	160 days
Third-party review	No

**APPLICANT**

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Company	<b>Ad-Tech Medical Instrument Corp</b>
Location	Racine, WI, US
Contact	DAVID PUTZ
510(k) history	18 submissions · 17 cleared · 1985-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k861031/>, 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 26, 2026