

**K880042 ACUTRON**Aug 1, 1988  
209 days to decisionK880042 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k880042/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Jan 5, 1988
Decision date	Aug 1, 1988
Days to decision	209 days
Third-party review	No

**APPLICANT**

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Company	<b>Davis Assoc.</b>
Location	Bridgeport, CT, US
Contact	PAUL DAVIS
510(k) history	1 submissions · 1 cleared · 1988-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k880042/>; 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 July 1, 2026