

**K880426 ELECTROSTIMULATOR MODEL 808**Sep 19, 1988  
231 days to decisionK880426 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k880426/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Feb 1, 1988
Decision date	Sep 19, 1988
Days to decision	231 days
Third-party review	No

**APPLICANT**

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Company	<b>Cornell Enterprises Co., Inc.</b>
Location	San Diego, CA, US
Contact	CORNELL N FUNG
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/k880426/>; 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