

**K030140 DUOMED, SERIES 500, (MODELS ID 500, FL 500)**Mar 18, 2003  
63 days to decisionK030140 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k030140/>**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 14, 2003
Decision date	Mar 18, 2003
Days to decision	63 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Duomed, Inc.</b>
Location	West Palm Beach, FL, US
Contact	CHARLES KOKINOS
510(k) history	1 submissions · 1 cleared · 2003-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k030140/> 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 June 30, 2026