

**K020180 RELIEFBAND DEVICE, MODELS RB-DL, RB-EL, RB-RL**Mar 21, 2002  
62 days to decisionK020180 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k020180/>**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 18, 2002
Decision date	Mar 21, 2002
Days to decision	62 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Woodside Biomedical, Inc.</b>
Location	Lake Forest, CA, US
Contact	CAROL PATTERSON
510(k) history	6 submissions · 6 cleared · 1998-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k020180/> 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