

**K014020 NT MEDITENS PLUS, TYPE 290**Jul 11, 2002  
217 days to decisionK014020 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k014020/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Dec 6, 2001
Decision date	Jul 11, 2002
Days to decision	217 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Bio-Medical Research, Ltd.</b>
Location	Washington Dc, DC, US
Contact	MICHELLE SAWYER
510(k) history	32 submissions · 31 cleared · 1996-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k014020/> 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 15, 2026