

**K102051 PAIN BUDDY**Jul 29, 2011  
373 days to decisionK102051 · Product code: **NUH** · Neurology  
Source: <https://www.510kdatabase.net/k102051/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, Over-the-counter (NUH)
Date received	Jul 21, 2010
Decision date	Jul 29, 2011
Days to decision	373 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Biomedical Life Systems, Inc.</b>
Location	Walker, MI, US
Contact	GARY BUSSETT
510(k) history	35 submissions · 35 cleared · 1984-2018

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k102051/> 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 May 30, 2026