

**K931923 DANTEC MAGLITE MAGNETIC STIMULATOR  
(MAGLITE)**Nov 9, 1993  
207 days to decisionK931923 · Product code: **GWF** · Neurology  
Source: <https://www.510kdatabase.net/k931923/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Apr 16, 1993
Decision date	Nov 9, 1993
Days to decision	207 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Dantec Medical, Inc.</b>
Location	Mahwah, NJ, US
Contact	RICHARD D MANTHEI
510(k) history	25 submissions · 25 cleared · 1990-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k931923/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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