

**K903161 MC II (TM), MODEL 7200S**Mar 27, 1991  
252 days to decisionK903161 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k903161/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Jul 18, 1990
Decision date	Mar 27, 1991
Days to decision	252 days
Third-party review	No

**APPLICANT**

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Company	<b>Medical Devices, Inc.</b>
Location	Mchenry, IL, US
Contact	BRUCE MACFARLANE
510(k) history	49 submissions · 47 cleared · 1977-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k903161/>; 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 21, 2026