

**K922029 REMUSK MP**Oct 6, 1992  
158 days to decisionK922029 · Product code: **LBB** · Neurology  
Source: <https://www.510kdatabase.net/k922029/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dynamometer, Ac-powered (LBB)
Date received	May 1, 1992
Decision date	Oct 6, 1992
Days to decision	158 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Equipement Biomedical Specialise, Inc.</b>
Location	Canada, CA
Contact	SYLVAIN BOUCHER
510(k) history	1 submissions · 1 cleared · 1992-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k922029/> 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 July 1, 2026