

**K790120 SLENDERTONE**May 7, 1979  
104 days to decisionK790120 · Product code: **IPF** · Physical MedicineSource: <https://www.510kdatabase.net/k790120/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Jan 23, 1979
Decision date	May 7, 1979
Days to decision	104 days
Third-party review	No

**APPLICANT**

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Company	<b>Bloomex International, Inc.</b>
Location	Mchenry, IL, US
510(k) history	16 submissions · 14 cleared · 1979-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k790120/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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