

K810247 MBI-4Mar 10, 1981
40 days to decisionK810247 · Product code: **IPF** · Physical MedicineSource: <https://www.510kdatabase.net/k810247/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Jan 29, 1981
Decision date	Mar 10, 1981
Days to decision	40 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k810247/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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