FDA 510(k) Premarket Notification

K810247 MBI-4

CLEARED

Mar 10, 1981
40 days to decision

K810247 · Product code: **IPF** · Physical Medicine
Source: 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