FDA 510(k) Premarket Notification

K790120 SLENDERTONE

CLEARED

May 7, 1979
104 days to decision

K790120 · Product code: **IPF** · Physical Medicine
Source: https://www.510kdatabase.net/k790120/

SUBMISSION DETAILS

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

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