FDA 510(k) Premarket Notification

K933653 MS 2500

CLEARED

Mar 23, 1994
238 days to decision

K933653 · Product code: **IPF** · Physical Medicine
Source: https://www.510kdatabase.net/k933653/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Jul 28, 1993
Decision date	Mar 23, 1994
Days to decision	238 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Lsi Intl., Inc.**
Location	Overland Park, KS, US
Contact	STEVE PAPA
510(k) history	5 submissions · 5 cleared · 1994-2000