FDA 510(k) Premarket Notification

K901849 LUMISCOPE EMS 8000

CLEARED

Aug 24, 1990
122 days to decision

K901849 · Product code: **IPF** · Physical Medicine
Source: https://www.510kdatabase.net/k901849/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Apr 24, 1990
Decision date	Aug 24, 1990
Days to decision	122 days
Third-party review	No

APPLICANT

Company	**Lumiscope Co., Inc.**
Location	Edison, NJ, US
Contact	MORGENIER, III
510(k) history	13 submissions · 13 cleared · 1988-2003