FDA 510(k) Premarket Notification

K010175 MICROLIGHT 830 LASER SYSTEM

CLEARED

Feb 6, 2002
384 days to decision

K010175 · Product code: **NHN** · Physical Medicine
Source: https://www.510kdatabase.net/k010175/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Light Based Laser Non-thermal Instrument With Non-heating Effect For Adjunctive Use In Pain Therapy (NHN)
Date received	Jan 18, 2001
Decision date	Feb 6, 2002
Days to decision	384 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Microlight Corporation of America**
Location	Missouri City, TX, US
Contact	MICHAEL BARBOUR
510(k) history	3 submissions · 3 cleared · 2002-2008