FDA 510(k) Premarket Notification

K020657 ACCULASER PRO LOW LEVEL LASER THERAPY DEVICE

CLEARED

Jul 29, 2002
151 days to decision

K020657 · Product code: **NHN** · Physical Medicine
Source: https://www.510kdatabase.net/k020657/

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	Feb 28, 2002
Decision date	Jul 29, 2002
Days to decision	151 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Acculaser, Inc.**
Location	Washington, DC, US
Contact	PATSY J TRISLER
510(k) history	2 submissions · 2 cleared · 2002-2004