FDA 510(k) Premarket Notification

K023623 NOVAVISION, MODEL 2.0

CLEARED

Apr 22, 2003
175 days to decision

K023623 · Product code: **HPT** · Neurology
Source: https://www.510kdatabase.net/k023623/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Perimeter, Automatic, Ac-powered (HPT)
Date received	Oct 29, 2002
Decision date	Apr 22, 2003
Days to decision	175 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Novavision, Inc.**
Location	Washington, DC, US
Contact	SANDRA J.P. DENNIS
510(k) history	3 submissions · 3 cleared · 1996-2003