FDA 510(k) Premarket Notification

K001808 DORO HEADREST SYSTEM

CLEARED

Aug 4, 2000
50 days to decision

K001808 · Product code: **HBL** · Neurology
Source: https://www.510kdatabase.net/k001808/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Holder, Head, Neurosurgical (skull Clamp) (HBL)
Date received	Jun 15, 2000
Decision date	Aug 4, 2000
Days to decision	50 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Pro-Med Instruments GmbH**
Location	Roswell, GA, US
Contact	ANITA THIBEAULT
510(k) history	10 submissions · 10 cleared · 2000-2024