

**K001808 DORO HEADREST SYSTEM**Aug 4, 2000  
50 days to decisionK001808 · 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	<b>Pro-Med Instruments GmbH</b>
Location	Roswell, GA, US
Contact	ANITA THIBEAULT
510(k) history	10 submissions · 10 cleared · 2000-2024

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k001808/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 26, 2026