

K981398 ANATOMARK NONINVASIVE HEAD REFERENCE SYSTEMJun 1, 1998
45 days to decisionK981398 · Product code: **LNH** · Radiology
Source: <https://www.510kdatabase.net/k981398/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Apr 17, 1998
Decision date	Jun 1, 1998
Days to decision	45 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Interneuron Pharmaceuticals, Inc.
Location	Lexington, MA, US
Contact	SONJA LOAR
510(k) history	1 submissions · 1 cleared · 1998-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k981398/> Data sourced from FDA 510(k) public records (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. - Generated May 19, 2026