

**K920835 PRESATURATION**May 22, 1992  
88 days to decisionK920835 · Product code: **LNH** · Radiology  
Source: <https://www.510kdatabase.net/k920835/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Feb 24, 1992
Decision date	May 22, 1992
Days to decision	88 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Resonex, Inc.</b>
Location	Sunnyvale, CA, US
Contact	BRUCE FLOYD
510(k) history	21 submissions · 21 cleared · 1988-1994

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k920835/> 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 20, 2026