

**K900725 RADIO FREQUENCY RECEIVER SUBSYSTEM**Mar 23, 1990  
36 days to decisionK900725 · Product code: **LNH** · Radiology  
Source: <https://www.510kdatabase.net/k900725/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Feb 15, 1990
Decision date	Mar 23, 1990
Days to decision	36 days
Third-party review	No

**APPLICANT**

---

Company	<b>Myheal Technologies</b>
Location	Flushing, NY, US
Contact	ORBACH, PH.D.
510(k) history	1 submissions · 1 cleared · 1990-1990

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900725/>; 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 21, 2026