

K883737 VFQ HEAD COILOct 13, 1988
42 days to decisionK883737 · Product code: **LNH** · Radiology
Source: <https://www.510kdatabase.net/k883737/>**SUBMISSION DETAILS**

| | |
|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | System, Nuclear Magnetic Resonance Imaging (LNH) |
| Date received | Sep 1, 1988 |
| Decision date | Oct 13, 1988 |
| Days to decision | 42 days |
| Third-party review | No |

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k883737/>; 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 20, 2026