FDA 510(k) Premarket Notification

# K883737 VFQ HEAD COIL

**CLEARED**

Oct 13, 1988
42 days to decision

K883737 · 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 |