FDA 510(k) Premarket Notification

CLEARED

K900725 RADIO FREQUENCY RECEIVER SUBSYSTEM

Mar 23, 1990
36 days to decision

K900725 · 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	**Myheal Technologies**
Location	Flushing, NY, US
Contact	ORBACH, PH.D.
510(k) history	1 submissions · 1 cleared · 1990-1990