FDA 510(k) Premarket Notification

K110151 ACHIEVA R4 1.5T AND ACHIEVA R4 3.0T

CLEARED

Mar 22, 2011
63 days to decision

K110151 · Product code: **LNH** · Radiology
Source: https://www.510kdatabase.net/k110151/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Jan 18, 2011
Decision date	Mar 22, 2011
Days to decision	63 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Philips Health Care**
Location	Bothell, WA, US
Contact	LYNN HARMER
510(k) history	20 submissions · 20 cleared · 2011-2020

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k110151/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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