FDA 510(k) Premarket Notification

K071925 MAGNETOM ESSENZA

CLEARED

Aug 14, 2007
33 days to decision

K071925 · Product code: **LNH** · Radiology
Source: https://www.510kdatabase.net/k071925/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Jul 12, 2007
Decision date	Aug 14, 2007
Days to decision	33 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Siemens Medical Solutions USA, Inc.**
Location	Hoffman Estates, IL, US
Contact	JUDITH CAMPBELL
510(k) history	778 submissions · 778 cleared · 1980-2026

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k071925/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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