FDA 510(k) Premarket Notification

K240385 VINT

CLEARED

Oct 15, 2024
250 days to decision

K240385 · Product code: **LLZ** · Radiology
Source: https://www.510kdatabase.net/k240385/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Feb 8, 2024
Decision date	Oct 15, 2024
Days to decision	250 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Mediimg Corporation**
Location	Gangdong-Gu, KR
Contact	Byeon-Uk Jeon
510(k) history	1 submissions · 1 cleared · 2024-2024

REGULATORY CONSULTANT

Consulting firm	**Mediguide, Inc.**
Contact	Miso Choi

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov