

K232456 RW-1Aug 11, 2025
728 days to decisionK232456 · Product code: **LLZ** · Radiology
Source: <https://www.510kdatabase.net/k232456/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Aug 14, 2023
Decision date	Aug 11, 2025
Days to decision	728 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Mediott, Inc.
Location	Shinjuku-Ku, JP
Contact	Akio Oki
510(k) history	1 submissions · 1 cleared · 2025-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k232456/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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