

K223789 21HQ513DJan 9, 2023
21 days to decisionK223789 · Product code: **PGY** · Radiology
Source: <https://www.510kdatabase.net/k223789/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Display, Diagnostic Radiology (PGY)
Date received	Dec 19, 2022
Decision date	Jan 9, 2023
Days to decision	21 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Lg Electronics.Inc
Location	Pyeongtaek-Si, KR
Contact	Jinhwan Jun
510(k) history	26 submissions · 26 cleared · 2018-2025

REGULATORY CONSULTANT

Consulting firm	Lg Electronics
Contact	Daseul An

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k223789/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 13, 2026