

K232127 21HQ513D, 32HL512D, 31HN713D, 32HQ713DAug 15, 2023
29 days to decisionK232127 · Product code: **PGY** · Radiology
Source: <https://www.510kdatabase.net/k232127/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Display, Diagnostic Radiology (PGY)
Date received	Jul 17, 2023
Decision date	Aug 15, 2023
Days to decision	29 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k232127/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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