

K230845 32HL512D, 31HN713D, 32HQ713DApr 27, 2023
30 days to decisionK230845 · Product code: **PGY** · Radiology
Source: <https://www.510kdatabase.net/k230845/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Display, Diagnostic Radiology (PGY)
Date received	Mar 28, 2023
Decision date	Apr 27, 2023
Days to decision	30 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

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