FDA 510(k) Premarket Notification

K232127 21HQ513D, 32HL512D, 31HN713D, 32HQ713D

CLEARED

Aug 15, 2023
29 days to decision

K232127 · 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