

**K212137 X-Clever**Dec 10, 2021  
155 days to decisionK212137 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k212137/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Jul 8, 2021
Decision date	Dec 10, 2021
Days to decision	155 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Lg Electronics.Inc</b>
Location	Pyeongtaek-Si, KR
Contact	Jin-Hwan Jun
510(k) history	26 submissions · 26 cleared · 2018-2025

**REGULATORY CONSULTANT**

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Consulting firm	<b>Gms Consulting</b>
Contact	BoKyeong Kim

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k212137/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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