

K161113 QXLinkAug 5, 2016
107 days to decisionK161113 · Product code: **LLZ** · Radiology
Source: <https://www.510kdatabase.net/k161113/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Apr 20, 2016
Decision date	Aug 5, 2016
Days to decision	107 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Vieworks Co., Ltd.
Location	Gyeonggi-Do, KR
Contact	YOONJAE IM
510(k) history	19 submissions · 19 cleared · 2007-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k161113/> 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 26, 2026