FDA 510(k) Premarket Notification

K241879 vRad Viewer

CLEARED

Jul 18, 2024
20 days to decision

K241879 · Product code: **LLZ** · Radiology
Source: https://www.510kdatabase.net/k241879/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Jun 28, 2024
Decision date	Jul 18, 2024
Days to decision	20 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Virtual Radiologic Corporation**
Location	Eden Prairie, MN, US
Contact	Wade Steigauf
510(k) history	3 submissions · 3 cleared · 2009-2024

REGULATORY CONSULTANT

Consulting firm	**Proxima Clinical Research**
Contact	Ellie Reynolds

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k241879/ 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 13, 2026