

K080553 DIGITAL RADIOGRAPHY SYSTEM, MODEL QXR-16Apr 16, 2008
48 days to decisionK080553 · Product code: **KPR** · Radiology
Source: <https://www.510kdatabase.net/k080553/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Feb 28, 2008
Decision date	Apr 16, 2008
Days to decision	48 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k080553/> 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