

**K222719 CX50N**May 9, 2023  
243 days to decisionK222719 · Product code: **PGY** · Radiology  
Source: <https://www.510kdatabase.net/k222719/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Display, Diagnostic Radiology (PGY)
Date received	Sep 8, 2022
Decision date	May 9, 2023
Days to decision	243 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Wide Corporation</b>
Location	Denton, TX, US
Contact	YeoJin Yun
510(k) history	20 submissions · 20 cleared · 2003-2023

**REGULATORY CONSULTANT**

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Consulting firm	<b>Ot Consulting, Inc.</b>
Contact	Josh Baker

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/k222719/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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