

**K160351 CX50N, CX50YQS**Apr 7, 2016  
59 days to decisionK160351 · Product code: **PGY** · Radiology  
Source: <https://www.510kdatabase.net/k160351/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Display, Diagnostic Radiology (PGY)
Date received	Feb 8, 2016
Decision date	Apr 7, 2016
Days to decision	59 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k160351/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 16, 2026