FDA 510(k) Premarket Notification

K160351 CX50N, CX50YQS

CLEARED

Apr 7, 2016
59 days to decision

K160351 · 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	**Wide Corporation**
Location	Denton, TX, US
Contact	YeoJin Yun
510(k) history	20 submissions · 20 cleared · 2003-2023