FDA 510(k) Premarket Notification

K171012 Veraview X800

CLEARED

Dec 19, 2017
259 days to decision

K171012 · Product code: **OAS** · Radiology
Source: https://www.510kdatabase.net/k171012/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	X-ray, Tomography, Computed, Dental (OAS)
Date received	Apr 4, 2017
Decision date	Dec 19, 2017
Days to decision	259 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**J. Morita USA, Inc.**
Location	Irvine, CA, US
Contact	Keisuke Mori
510(k) history	52 submissions · 52 cleared · 1988-2022