

**K171012 Veraview X800**Dec 19, 2017  
259 days to decisionK171012 · Product code: **OAS** · Radiology  
Source: <https://www.510kdatabase.net/k171012/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k171012/> 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 14, 2026