

**K190234 ODI HD Dental Sensor**Mar 1, 2019  
23 days to decisionK190234 · Product code: **MUH** · Radiology  
Source: <https://www.510kdatabase.net/k190234/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Extraoral Source, Digital (MUH)
Date received	Feb 6, 2019
Decision date	Mar 1, 2019
Days to decision	23 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cnd Global Technologies, Inc.</b>
Location	Cumberland, RI, US
Contact	Neeti Rathor
510(k) history	1 submissions · 1 cleared · 2019-2019

**REGULATORY CONSULTANT**

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Consulting firm	<b>Regulatory Science Solutions</b>
Contact	Lisa Jager

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/k190234/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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