

**K200179 Collaboration Live**Feb 18, 2020  
25 days to decisionK200179 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k200179/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Jan 24, 2020
Decision date	Feb 18, 2020
Days to decision	25 days
Third-party review	Yes
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Philips Health Care</b>
Location	Bothell, WA, US
Contact	Paul Elias
510(k) history	20 submissions · 20 cleared · 2011-2020

**REGULATORY CONSULTANT**

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Consulting firm	<b>Regulatory Technology Services, LLC</b>
Contact	Prithul Bom

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/k200179/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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