

**K241871 Philips IntelliSite Pathology Solution**Dec 2, 2024  
158 days to decisionK241871 · Product code: **PSY** · Pathology  
Source: <https://www.510kdatabase.net/k241871/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Whole Slide Imaging System (PSY)
Date received	Jun 27, 2024
Decision date	Dec 2, 2024
Days to decision	158 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Philips Medical Systems Nederland B.V.</b>
Location	Best, NL
Contact	Donna Peled
510(k) history	103 submissions · 102 cleared · 2005-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k241871/> 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 13, 2026