FDA 510(k) Premarket Notification

K241871 Philips IntelliSite Pathology Solution

CLEARED

Dec 2, 2024
158 days to decision

K241871 · Product code: **PSY** · Pathology
Source: https://www.510kdatabase.net/k241871/

SUBMISSION DETAILS

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

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