FDA 510(k) Premarket Notification

CLEARED

K260207 Multimodality Simulation Workspace (MM Sim) (v1.0.0)

May 27, 2026
124 days to decision

K260207 · Product code: **MUJ** · Radiology
Source: https://www.510kdatabase.net/k260207/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Planning, Radiation Therapy Treatment (MUJ)
Date received	Jan 23, 2026
Decision date	May 27, 2026
Days to decision	124 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	Veronica Padharia
510(k) history	104 submissions · 103 cleared · 2005-2026

REGULATORY CONSULTANT

Consulting firm	**Philips Medical Systems Technologies , Ltd.**
Contact	Carmit Shmuel

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov