FDA 510(k) Premarket Notification

K200179 Collaboration Live

CLEARED

Feb 18, 2020
25 days to decision

K200179 · Product code: **LLZ** · Radiology
Source: https://www.510kdatabase.net/k200179/

SUBMISSION DETAILS

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

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

REGULATORY CONSULTANT

Consulting firm	**Regulatory Technology Services, LLC**
Contact	Prithul Bom

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