FDA 510(k) Premarket Notification

K233930 MRCP+ version 2 (MRCP+ v2)

CLEARED

Mar 13, 2024
90 days to decision

K233930 · Product code: **LLZ** · Radiology
Source: https://www.510kdatabase.net/k233930/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Dec 14, 2023
Decision date	Mar 13, 2024
Days to decision	90 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Perspectum**
Location	Oxford, GB
Contact	Keri Hildick
510(k) history	3 submissions · 3 cleared · 2022-2024