

K233930 MRCP+ version 2 (MRCP+ v2)Mar 13, 2024
90 days to decisionK233930 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k233930/> Data sourced from FDA 510(k) public records (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. - Generated May 14, 2026