

K221984 InterView XPMar 29, 2023
266 days to decisionK221984 · Product code: **LLZ** · Radiology
Source: <https://www.510kdatabase.net/k221984/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Jul 6, 2022
Decision date	Mar 29, 2023
Days to decision	266 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	InterView FUSION

APPLICANT

Company	Mediso Medical Imaging Systems, Lttd.
Location	Budapest, HU
Contact	Daniel Cser
510(k) history	1 submissions · 1 cleared · 2023-2023

REGULATORY CONSULTANT

Consulting firm	Cro Group, Inc.
Contact	William McLain

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)**510k Database** - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k221984/> 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 27, 2026