

**K241620 ChestView US**Feb 27, 2025  
267 days to decisionK241620 · Product code: **MYN** · Radiology  
Source: <https://www.510kdatabase.net/k241620/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Medical Image (MYN)
Date received	Jun 5, 2024
Decision date	Feb 27, 2025
Days to decision	267 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Gleamer Sas</b>
Location	Paris, FR
Contact	Antoine Tournier
510(k) history	2 submissions · 2 cleared · 2025-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k241620/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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