

**K222035 Avatar Medical Software V1**May 24, 2023  
317 days to decisionK222035 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k222035/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Jul 11, 2022
Decision date	May 24, 2023
Days to decision	317 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Avatar Medical</b>
Location	Paris, FR
Contact	Elodie Brient-Litzler
510(k) history	3 submissions · 3 cleared · 2023-2025

**REGULATORY CONSULTANT**

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Consulting firm	<b>Cosm</b>
Contact	Rory Carrillo

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

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