

**K210344 inVisionOS**Nov 10, 2021  
278 days to decisionK210344 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k210344/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Feb 5, 2021
Decision date	Nov 10, 2021
Days to decision	278 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Precisionos Technology, Inc.</b>
Location	Vancouver, CA
Contact	Danny Goel
510(k) history	1 submissions · 1 cleared · 2021-2021

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k210344/> 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 26, 2026