FDA 510(k) Premarket Notification

K210344 inVisionOS

CLEARED

Nov 10, 2021
278 days to decision

K210344 · 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	**Precisionos Technology, Inc.**
Location	Vancouver, CA
Contact	Danny Goel
510(k) history	1 submissions · 1 cleared · 2021-2021