

K233126 aetherSlideMay 30, 2024
246 days to decisionK233126 · Product code: **QKQ** · Pathology
Source: <https://www.510kdatabase.net/k233126/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Digital Pathology Image Viewing And Management Software (QKQ)
Date received	Sep 27, 2023
Decision date	May 30, 2024
Days to decision	246 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Aetherai Co., Ltd.
Location	Taipei City, TW
Contact	Leticia Hong
510(k) history	1 submissions · 1 cleared · 2024-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k233126/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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