

Ref. No. VXD0261A/2025

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VERIFICATION OF COMPLIANCE

Issue Date: February 13, 2025

Applicant: DFI Inc.

Address: 10F., No. 97, Sec. 1, Xintai 5th Rd., Xizhi Dist.,

New Taipei City, Taiwan, R.O.C

Manufacturer: DFI Inc.

Address: 10F., No. 97, Sec. 1, Xintai 5th Rd., Xizhi Dist.,

New Taipei City, Taiwan, R.O.C

Product: Medical Edge Al inference System

Brand Name/Trade Mark: DFI; ITOX MOdel/Type: MPC350-RPS

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Applicable Standards: EN 60601-1-2: 2015 + A1: 2021

IEC 60601-1-2: 2014 + A1: 2020

CISPR 11: 2015 + A1: 2016 + A2: 2019 (Ed 6.2) IEC 61000-3-2: 2018 + A1: 2020 (Ed. 5.1)

IEC 61000-3-3: 2013 + AMD1: 2017 + AMD2: 2021 + COR1: 2022

IEC 61000-4-2: 2008

IEC 61000-4-3: 2020 (Ed. 4.0)

IEC 61000-4-4: 2012

IEC 61000-4-5: 2014 + A1: 2017

IEC 61000-4-6: 2023 IEC 61000-4-8: 2009

IEC 61000-4-11: 2020 + COR1: 2020 + COR2: 2022 (Ed. 3.0)

IEC 60601-1-2: 2014 + A1: 2020 Subclause 8.11 (IEC 61000-4-39 for §8.10)

Test Laboratory: Compliance Certification Services Inc.

Xindian Lab

No. 163-1, Jhongsheng Rd., Xindian Dist., New Taipei City, Taiwan

Test Report No.: TMXD2501000236DE, dated on February 13, 2025

Conclusion: Based upon a review of the Test Report(s), the tested sample of the product

mentioned above is deemed to comply with the requirements of the above standards

Note: This verification is only valid for the product and configuration described and in conjunction

with the test report as detailed above.

Authorised Signatory:

Compliance Certification Services Inc.

Jason Lee Section Manager

FM-023A-R02

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