# Certificate of Compliance

CE

We hereby declare that the technical files of all the items in each product group complies with the requirements of the Council Directive on **Medical Device Regulation - (Council Regulation 2017/745).**

**Certificate No.: DRAFT**

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| **Manufacturer** | : | **LAKXUS AUTOMATION** |
| **Address** | : | **SHED NO. 27, GROUND FLOOR, SARJAN ESTATE, KATHWADA ROAD, NIKOL, AHMEDABAD - 382350, GUJARAT, INDIA** |
| **Products** | : | **OZONATOR, OZONE GENERATOR, OXYGEN CONCENTRATOR, OZONE AIR STABILIZER, MEDICAL OZONE GENERATOR, OZONE AIR MONITOR, OZONE DISSOLVED MONITOR AND OZONE ANALYSER.** |

**Complies with the requirements applicable to it**

The quality system file has been assessed, approved and is subject to continuous surveillance according to the Council Directive on **Medical Device Regulation -(Council Regulation 2017/745).**

**This certificate is issued under the following conditions:**

1. It applies only to the quality system maintained in the manufacture of above referenced models and it does not substitute the design or type-examination procedures, if requested.
2. The certificate remains valid until the manufacturing conditions or the quality systems are changed.
3. The certificate validity is conditioned by positive results or surveillance audits.

The CE mark as shown above can be used, under the responsibility of the manufacturer, after completion of an EC Declaration of conformity and compliance with all relevant EC Directives. The statement is based on a single evaluation of test report of one sample of above mentioned product. It does not imply an assessment of the whole production.

**Validity of this certificate can be verified at www.ukcertifications.org.uk/verify**

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| Date of Certification | 30th June 2025 |
| 1st Surveillance Audit Due | 29th June 2026 |
| 2nd Surveillance Audit Due | 29th June 2027 |
| Certificate Expiry | 29th June 2028 |

UK Certification & Inspection Ltd.

Authorised Signatory