Certificate of Compliance

CE

We hereby declare that the technical files of all the items in each product group of complied with the requirements of the Council Directive on Medical Devices 93/42/EEC as Amended 2007/47/EC, directives 2014/35/EU Low Voltage Directive & directives 2006/42/EC Machinery Directive.

Certificate No.: CE- 1931

Manufacturer

Name : MAAS AIR-N-GAS TECHNOLOGIES PRIVATE LIMITED

Address : Corporate Office: B-1-801, West Gate Business Bay, Markarba, S.G. Highway, Ahmedabad-380051, Gujrat, India

Factory- 38, Panchamrut Industrial Park-3, Dhamatvan, Daskroi, Ahmedabad, Ahmedabad, Gujarat, 382435, India

Products : NITROGEN GENERATION PLANT, PSA OXYGEN GENERATION, MEDICAL OXYGEN PLANT, DEHUMIDIFIER, PRESUSURE VESSELS/ & AIR AND GAS DRYERS

Complies with the requirements applicable to it

The Certification body has performed an audit of the above product quality system covering the design, manufacture and final inspection of the certified product. The quality system has been assessed, approved and is subject to continuous surveillance according to the Council Directive on Medical Devices 93/42/EEC as Amended 2007/47/EC, directives 2014/35/EU Low Voltage Directive & directives 2006/42/EC Machinery Directive.

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.
4. After fulfilling the relevant EU legislation, the manufacturer shall affix to each device, of the referenced models.
5. 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 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

Date of Certification	24th November 2021
1st Surveillance Audit Due	23rd November 2022
2nd Surveillance Audit Due	23rd November 2023
Certificate Expiry (subject to the company maintaining its system to the required standard)	23rd November 2024

Authorised Signatory

UK Certification & Inspection Ltd.