



Certificate of Compliance

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Confirms that the Class II complies according to the directive 93/42/EEC Medical Council Directive.

Certificate No.: CE-3653

Manufacturer

Name : Microflow Devices India Pvt. Ltd.

Address : Plot No. 4&5, Wip SIDCO Industrial Estate Thirumudivakkam,

Chennai 600044, Tamilnadu, India

Products

Name : Laminar Air Flow, Modular Operating Room, Biological Safety

Cabinet Class II Type A2, Type B2 and Class III, Dispensing

Booth, Fume Hood, Pass Box and Air Shower Only

The manufacturer's technical documentation as required for Medical Council according to the directive 93/42/EEC has been reviewed and found to comply with the requirements for Class II Medical Devices Directive. Any significant changes in the design or construction of the product, not agreed upon by us, this declaration will lose its validity.

This certificate is issued under the following conditions:

- 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.
- The certificate remains valid until the manufacturing conditions or the quality systems are changed.
- The certificate validity is conditioned by positive results or surveillance audits.
- After fulfilling the relevant EU legislation, the manufacturer shall affix to each device, of the above 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.co.uk/verify

Date of Certification

22nd October 2018

1st Surveillance Audit Due

21st October 2019

2nd Surveillance Audit Due

21st October 2020

Certificate Expiry (subject to the company maintaining its

21st October 2021

system to the required standard)

Authorised Signatory