Certificate of Compliance

CE

We hereby declare that an examination of the under mentioned full quality assurance system has been carried out the following requirements of the national legislation to which the undersigned is subjected, transposing annex I- II (with the exemption of section 4) of the directive 93/42/EEC, Low Voltage Directive 2014/35/EU & Electromagnetic Compatibility Directive 2014/30/EU on medical devices. We UKCert Certify that the full quality assurance system confirms with the relevant provisions of the aforementioned directive. Identification of the products covered by this certificate is given in the Appendix

Certificate No.: CE-5957

Manufacturer

Name : **Medicaid Systems**

Address : **Plot: 667, JLPL Industrial Park, Sector: 82, Mohali (SAS Nagar) Punjab - 140306 India**

Product Name:
- **NEUROMAX (NMX-32 V) 32 Channel EEG Long Term Video Monitoring System with Brain Mapping**
- **NEUROMAX (NMX-32) 32 Channel Portable EEG System with Brain Mapping**
- **NEUROMAX - 64 channel EEG with Brain Mapping & Long Term Video Monitoring System Model : (NMX-64)**

Classification of Medical Device, Low Voltage Directive, Electromagnetic Compatibility

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 Directive 93/42/EEC Medical Device Directive, 2014/35/EU Low Voltage Directive & 2014/30/EU Electromagnetic Compatibility 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 of Class 1-II in medical directive, the manufacturer shall affix to each device, of the referenced models.

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 in Class I-II product. The statement is based on a single evaluation of one sample of above mentioned products. 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 initial registration	16th August 2024
Date of this certificate	16th August 2024
Certificate Expiry	15th August 2027
Recertification due (subject to the company maintaining its system to the required standard)	15th August 2027

Authorised Signatory

This certificate is the property of UK Certification & Inspection Limited and shall be returned immediately on request.
71-75 Shelton Street, Covent Garden, London, WC2H 9JQ, United Kingdom
Website:- www.ukcertifications.org.uk, email:- info@ukcertifications.org.uk
Company No. 11847851