REGAL REGISTER (UK) LIMITED

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# CERTIFICATE

## *Certificate of Compliance*

We confirm that the technical documentation for the below mentioned products according conformity to according to the council Directive 93/42/EEC Medical Devices Directive (MDD).

Products:

**Anesthesia Machine, Hydraulic O.T. Table, Remote O.T. Table, Ceiling Mounted Led O.T. Light, Ceiling Mounted Llalogen O.T. Light, Medical Gas Pipeline Accessories.**

Manufactured by company:

**United Health Care Products India**

**8/4 Ho Chi Min Sarani, (Biren Roy Road West) Behala Chowrasta, Kolkata-700008, West Bengal, India.**
is complying to the applicable essential requirements of according to the council Directive 93/42/EEC Medical Devices Directive (MDD).
The Regal Register (UK) Ltd. has reviewed manufacturer's technical documentation & Test reports and found it in compliance with above mentioned Directive (s).

This certificate is issued under the following conditions:

1. It applies only to the above referenced set of products. The manufacturer is obligated to assure that all products of the respective model confirm to the type assessed for this certificate.
2. The Certificate remains valid until the manufacturing conditions, the quality systems or relevant legislation are changed subjected to maximum Validity of 3 years.
3. The Certificate validity is conditioned by the positive results of the surveillance audits.
4. After fulfilling the relevant EU legislation requirements, the manufacturer shall affix to each product of the above referenced models, CE Marking according to the following example.

| | |
|---|---|
| *Certificate No.* | : CE- 2912 |
| *Date of registration* | : 02nd November |
| *Date of this certificate* | : 02nd November |
| **Expiry Date* | : 01st November |
| *Recertification Due Date* | : 01st November |

**Validity of certificate is subjected to the continued satisfactory performance during surveillance audit*

CE

*Authorized Signatory*
**Regal Register (UK) Ltd.**

**Statement:**
This certificate of conformity based on the evaluation of a sample of the above mentioned products. It does not imply an assessment of the mass-production of the product. The certification body should be informed (revision of technical file) for any modification or alterations made to the aforementioned product type(s). The manufacturer is responsible for the product and ensuring that all manufactured products are in compliance with the specifications declared in the technical construction file.
Further clarifications regarding the scope of this certificate and the applicability of the management system requirements may be obtained by contacting the organization. Check www.regalregistrars.co.uk for current status of the certificate. Whilst all due care and skill was exercised in carrying out this assessment, RRUL accepts responsibility only for proven gross negligence. This certificate remains property of RRUL to whom it must be returned upon request.