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 Medical Device Directive EU Directive 93/42/EEC

Certificate No.: CE-3715

Manufacturer

Name : **AAR KAY ENTERPRISES**

Address : **E 25, Jawahar Park, Laxmi Nagar, Delhi 110092, India**

Products : **Hospital Furniture, Medical Equipments, Physiotherapy Equipments, Healthcare Products and Disposable**

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 Medical Device Directive EU Directive 93/42/EEC

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 not changed.
3. The certificate validity is conditioned by positive results of 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 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	10th September 2018
1st Surveillance Audit Due	09th September 2019
2nd Surveillance Audit Due	09th September 2020
Certificate Expiry (subject to the company maintaining its system to the required standard)	09th September 2021

Authorised Signatory

UK Certification & Inspection Ltd.