



Issue Number: 2

Approved By: H.JOHNSON

Date: 27.05.2014

QD 65 ISSUE 2 MAY 2014

Certificate of Conformity

This is to certify that all Jenx Products
Conform to the EC Medical Devices Directive and have been
marked with the CE mark.

These products fall into Class I Devices: Aids for Daily Living.

Signed.....



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DECLARATION of CONFORMITY

We, **Jenx Limited (Manufacturer)**
Wardsend Rd
Sheffield, S6 1RQ
United Kingdom Tel.+44(0)1142853376
Fax+44(0)1142853528

Hereby declare that the products specified on the products list below, meets the essential health and safety requirements and is in conformance with the provisions of the MEDICAL DEVICES DIRECTIVE (MDD) 93/42/EEC annex 7.

The devices specified on the product list below are classified as Class I, "Aids for Daily Living". The classification is based on the requirements of Rule 1 of annex IX, of the MDD 93/42/EEC

Jenx Limited operate a quality system which meets requirements of ISO 9001:2008 as indicated on certificate No. 417.

The certificate has been granted by BM Trada Limited, United Kingdom.

The CE mark is applied under the guidelines of annex II of the MDD 93/42/EEC.
The CE marking has been affixed on the device according to Article 17 of the MDD 93/42/EEC.

PRODUCT LIST.

All Jenx Products

Signed.....
Date 08/8/2017.....