Lloyd's Register
Notified Body authorised by the MCA

Maritime & Coastguard Agency

EC (MODULE D) CERTIFICATE OF CONFORMITY

LLOYD'S REGISTER VERIFICATION LIMITED (LRV), designated as a "notified body" under the terms of the Merchant Shipping (Marine Equipment) Regulations 2016 (S.I. 2016 No. 1025) did undertake an assessment of the subject manufacturer's Quality System as per requirements of Annex II Conformity to type based on quality assurance of production process (Module D) of Marine Equipment Directive (MED) 2014/90/EU and Commission Implementing Regulation (EU) 2018/773 indicating design, construction and performance requirements and testing standards for marine equipment and was found to conform with the requirements for the Product Types below.

Manufacturer (Applicant)	Xiantao Deming Healthcare Products Co., Ltd
Address	No.198, Pengchang Avenue Xiantao City Hubei Province China
Authorised Representative	Sungo Certification Company Limited
Address	RM101, Maple House 118 High Street Purley London CR8 2AD United Kingdom
Reference	**Regulation (EU) 2018/773**
Regulation Item (No & designation)	**MED/1.7** THERMAL PROTECTIVE AIDS

Approval is subject to continued maintenance of the requirements of the above mentioned Directives and to all products continuing to comply with the standards and conditions of EC Type Examination Certificates issued by Lloyd's Register Verification or other Notified Body when they are of the above Designation(s).

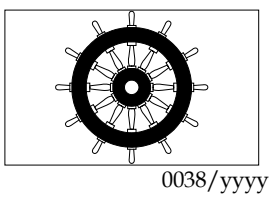
Approval is further subject to continued maintenance of the certified quality management system in accordance with the requirements of CQC certificate number 00112Q26931R1M/4200 or an equivalent replacement thereof.

Authorisation is hereby given to the manufacturer to use the LRV Notified Body Registration Number 0038 in accordance with the requirements of the specified Directives in relation to the described products.

This certificate remains valid unless cancelled or revoked, provided that products manufactured under this Certificate remain satisfactory in service and the above quality management system continues to be approved.

Date of issue	17 February 2019	Expiry date	16 February 2022
Certificate No.	MED 1950037	Signed	
Sheet No	1 of 2	Name	J. Deboer For and on behalf of Lloyd's Register Verification **LRV EC Distinguishing No. 0038**

Note:
This certificate is issued under the authority of the MCA. No product shall be manufactured under this Certificate unless a valid EC Type Examination Certificate (Module B) is held on that product's Technical File. The manufacturer shall advise the Notified Body of all proposed modifications or changes to a product for which an EC Type Examination Certificate (Module B) has been issued, and of proposed changes of manufacturing location or process, and shall retain copy of their written authorisation or Certification of such changes.


0038/yyyy

Subject to the Manufacturer's compliance with the foregoing, and those conditions of the Directive, the Manufacturer or his Authorised Representative is allowed to affix the 'Mark of Conformity' to products of the types shown above.

yyyy = The year in which the mark is affixed.

Lloyd's Register Verification Limited
71 Fenchurch Street, London, EC3M 4BS
Telephone 020 7423 2416
Email med@ lr.org

Document number
MED 1950037

Issue number
1

Date
26 March 2019

Quote this reference on all future communications
UKITSO/TA/QA/MED/JD

EC QUALITY SYSTEM (MODULE D) CERTIFICATE No. MED 1950037

This Certificate is renewal of the Certificate No. MED 1650075/M1

Places of Production
Xiantao Deming Healthcare Products Co. Ltd No. 198 Pengchang Avenue Xiantao City Hubei Province China

Module B No's	Regulation Item number	Date of issue / revision	Notified body
MED 1550036	A.1/1.7	03/02/2015	0038