

Attestation of Conformity

according to Medical Device Directive (93/42/EEC as amended
by 2007/47/EC)

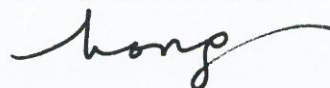
Certificate No.
CA17P1072

<i>Owner of Certificate</i>	JIANGSU ALPHAY MEDICAL DEVICE CO., LTD. No. 95, Zhenxing Road, Economical & Development Zone, Nantong, Jiangsu, China
<i>Manufacturer</i>	JIANGSU ALPHAY MEDICAL DEVICE CO., LTD. No. 95, Zhenxing Road, Economical & Development Zone, Nantong, Jiangsu, China
<i>Trade Mark</i>	N/A
<i>Product</i>	Air-traction Belt, Neck Traction Device
<i>Type/Model</i>	Air-traction Belt : YGAH-1, YGAH-2, YGAH-3 Neck Traction Device : JQAH-I, JQAH-II, JQAH-III
<i>Reference Document</i>	T.F. Document (AH/CH-001-2016, Rev.A/2 3, July.2017) T.F. Document (AH/CH-002-2016, Rev.A/2 3, July.2017)

The product described above complies with the requirements of the Medical Device Directive (93/42/EEC amended by 2007/47/EC) Annex I and Annex VII. The details about the product conformity and applied standards are mentioned in the test report referenced above. This certificate is subject to Kiwa Cermet regulations and it is valid only for the above mentioned medical device(s). This Certificate, different from an EC Certificate, counts as Attestation of conformity to be used by first part.

<i>Issue date</i>	12-07-2017
<i>Last revised date</i>	N/A
<i>Expiry date</i>	11-07-2022
<i>Revision</i>	0

Kiwa Cermet Lead Auditor



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The CE marking may be used if all relevant and effective EC directives are complied with.