CERTIFICATE

ATTESTATION CERTIFICATE FOR MEDICAL DEVICE DIRECTIVE

Technical file of the company mentioned below has been observed.

MDD 93/42/EEC Medical Device Directive Annex VII has been taken as references for these processes.

Company Name	: Zhangjiagang Tengda Machinery Manufacturer Co., Ltd.
Company Address	: No.3. District, Changyinsha Farm, Zhangjiagang City, Jiangsu Province, China
Related Directives and Annex	: MDD 93/42/EEC Medical Device Directive/Annex VII Class I Non-Sterile
Related Standards	: EN ISO 14971-2012; EN ISO 15223-1:2016; EN 1865-1:2010+A1:2015
Product Name	: Stretcher
Report No and Date	: TENGDA-190612
Product Brand/Model/Type	: MLF999-A, MLF999-D, MLF999-B, MLF999-B1, MLF999-C, MLF999-C1, MLF999-C2, MLF999-C3, TD01016, TD010161, TD010163, TD010162-A, TD010162-B, TD010162-C, MLF999-F1, MLF999-F2, MLF999-A3-1, MLF999-A3-2, MLF999-A3-3, TD03016, TD03017, TD01080, TD01081, DYC-87A, DYC-87B, DYC-87C

Certificate Number	: M.2019.206.C1469
Initial Assessment Date	: 14.06.2019
Registration Date	: 17.06.2019
Reissue Date/No	: -
Expiry Date	: 16.06.2024

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