EUROPEAN INSPECTION AND CERTIFICATION COMPANY S.A.

CERTIFICATE OF CONFORMITY

FULLNESS EXAMINATION OF TECHNICAL FILE

Certificate No.	:	MDLVD.0010
Issue Date	:	21/06/2017
Expiry Date	:	20/06/2022
Applicant (Name & Address)	:	LIFELINE PHARMA 623, AVIOR NIRMAL GALAXY, L.B.S. MARG, MULUND (W), MUMBAI - 400 080, MAHARASHTRA, INDIA
Manufacturer (Name & Address)	:	SAME AS ABOVE
Test Report Ref	:	ITC/TEST/NS/1705/04
TCF No.	:	LL/TCF/01
Product Description	:	BIO-MEDICAL WASTE TREATMENT SYSTEM
Model(s)	:	STERIWELL-20
Directive(s)	:	Machinery Directive(2006/42/EC) and Low Voltage Directive (2014/35/EU)
Standard(s)	:	EN ISO 12100:2010 & EN 60204-1:2006+A1:2009

This is to certify that, upon the relevant application of the above-mentioned company, EUROCERT S.A as Third Party Authority has reviewed the Technical Construction File of the described product which found to fulfill the basic health and safety prerequisites of above mentioned Directive(s).

Note:

- *The manufacturer should issue a Declaration of Conformity according to the basic requirements of the applicable and relevant directives.*
- *The CE marking can be affixed on the above mentioned product with the manufacturer's responsibility, if all relevant and applicable directives are complied with.*
- *All modifications to the Technical File should be first submitted to the Third Party Inspection Authority to ensure further validity of this attestation.*
- *This certificate is valid only for the product and configuration described and in conjunction with the technical data detailed above.*

Please check the validity of the certificate from our website using the password 51w3P54z

Third Party Authority Stamp

On Behalf of EUROCERT

George N.Sifonios
Director of Development

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