COMPLIANCE DOCUMENT

South Asia

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This Compliance Document is issued to:

LIFELINE PHARMA
623, Avior, Nirmal Galaxy, L.B.S. Marg,
Mulund West, Mumbai – 400080,
Maharashtra (India)

FOR

Product	: Biomedical Waste Treatment System (Non-Burn Technology)
Model No.	: STERIWELL-20 & STERIWELL-40
Test Standards	: EN ISO 12100:2010, EN 60204-1:2018, EN 61000-6-2:2005 & EN 61000-6-4:2007/A1:2011
Test Report No.	: BLR/ENE/SAF/22/1926748 & BLR/ENE/EMC/22/1926748

This is to confirm that a sample of machine was tested to the safety test requirements of Machinery Directive (2006/42/EC), Low voltage directive (2014/35/EU) and Electromagnetic Compatibility Directive (2014/30/EU) by us at the manufacturing location.

CE

After implementation of the requirements as per test report no. BLR/ENE/SAF/22/1926748 and BLR/ENE/EMC/22/1926748 issued to the manufacturer, preparation of the necessary technical documentation and signing of the declaration of conformity, CE marking can be affixed on the product. Other relevant directives have to be observed.

The document holder is responsible for the consistent manufacturing of the product in compliance with the test sample submitted to us.

TÜV SÜD SOUTH ASIA PVT. LTD.

Sr GM-PRODUCT SERVICE – ENE

Ref No.	Date of Original Issue	Date of revision:
CD/1926748	14th July,2022	

Note: Products entering the European Union are subject to the requirements set forth in the European Directives. It is the responsibility of the person placing the product on the European market to ensure compliance with applicable Directives. The requirements of directives and standards change from time to time and it is the responsibility of the manufacturer to ensure compliance. Our services are intended to provide you with the data needed to determine compliance, but the person placing the product on the European market still has the responsibility to ensure each product complies with the applicable Directives. The product liability rests with the manufacturer or his representative in accordance with Council Directive 85/374/EEC.

This document applies only to the particular model provided for testing and CE certification and does not permit the use of a TÜV PRODUCT SERVICE certification mark. It becomes ineffective in the event of changes relating to products, applicable standards and the Directives.