PROCEDURE APPROVALS

Note: Date of last approval will serve as the Approval Date for this document.

Name	Function	Signature	Date
Dionne Sanders	Regulatory	dsanders	07/06/2021

Digitally signed by Șarco Alexandru
Date: 2022.07.28 22:55:34 EEST
Reason: MoldSign Signature
Location: Moldova

Declaration of Conformity

Manufacturer: CONMED Corporation
Address: 525 French Road
Utica, New York 13502 USA

European Auth. Rep.: MDSS GmbH
Schiffgraben 41
D-30175 Hannover
Germany

Notified Body Address: BSI Group The Netherlands B.V.
Say Building, John M. Keynesplein 9, 1066EP
Amsterdam
Netherlands

NB Identification #: N/A

Conformity Assessment: Annex II and III , of the Regulation (EU) 2017/745 on Medical Devices

EC Certificate Number: Self-Certified

Device / Risk Classification: Class I

Rule per Annex VIII: 1 and 13

Product Family: SD – Surgical Instruments and Accessories
Intended Purpose: accessories used to display alphabetical and numerical data

Devices:

Reference Number	Product Description	Date 1st CE Marked	Basic UDI-DI	Technical Document #
VP4826	HD LCD Monitor, 26 in. 66 cm	Oct 2014	084585464945FD	AV-SD-002
VP4832	HD LCD Monitor, 32 in. 81 cm	Dec 2014	084585464945FD	AV-SD-002

This declaration is issued under the sole responsibility of the manufacturer.

We, the manufacturer, hereby declare that the medical devices listed on this declaration conform with Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 concerning medical devices.

Place of issue of the DoC: ConMed Corporation
525 French Road
Utica, NY- 13502
USA

Date of issue of the DoC: See Windchill approver Coversheet

Revision History

Revision #	Reason for Change
1	Creation of EU MDR Declaration of Conformity from MDD (V10-36612)