HANS DINSLAGE GmbH	Title	**EC Declaration of Conformity**
	Date	2025-10-23

Manufacturer:	Hans Dinslage GmbH (see address in footer)
SRN:	DE-MF-000005488
Product category:	TENS devices
Product type:	SEM 39
Intended use:	The device is intended to relieve pain using TENS (transcutaneous electrical nerve stimulation) technology.

The product specified above is in conformity with the following specifications.

(EU) 2017/745	Medical device regulation (MDR)
Basic-UDI-DI:	4066299SEM39XM
Classification/applied rule(s):	Class IIa/rule 9
Conformity assessment procedure:	Annex IX, Chapter I

The notified body mdc medical device certification GmbH, located at Kriegerstr. 6, 70191 Stuttgart, Germany, identification number 0483, issued in the course of the mentioned conformity assessment procedure the following certificate:

Certificate no. and validity:	D1326900049, valid to 2026-04-08

2011/65/EU	Restrictions of the use of certain hazardous substances in electrical and electronic equipment (RoHS) EN IEC 63000:2018

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

Signed for and on behalf of:	Hans Dinslage GmbH
Place, date of issue:	Ulm, 2025-10-23
Name, function, signature, stamp:	Werner Meternek, Director Quality Management & Regulatory Affairs ppa. Hans Dinslage GmbH Riedlinger Str. 28 • 88524 Uttenweiler

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