Declaration of Conformity

This Declaration of Conformity is issued under the sole responsibility of the manufacturer.

MANUFACTURER		
Name of Company	**Address**	**SRN**
Ferno Canada Inc. **Origin of Manufacture:** Guangzhou Linbi Medical Devices Co., Ltd.	2460 Tedlo Street Mississauga, Ontario, Canada Room 805, 2nd Building, Headquarters Center, Tian An Hi-tech Ecological Park, No. 555 North Panyu Road, Panyu District, Guangzhou, China	TBD

EU AUTHORIZED REPRESENTATIVE			
Name of Company	**Address**	**SRN**	**Telephone/Email**
Ferno S.r.l. CE	via B.Zallone, n.26,40066 Pieve di Cento, Bologna, Italy	TBD	+39.051.6860028 www.ferno.it

PRODUCT IDENTIFICATION	
Product Brand Name	**Photo**
CAN CPR Board	
Intended Purpose:	
Professional Use Fist Aid	

Item / Catalog #	Item Description	Basic UDI-DI
CAN CPR	CPR Board	00754016222078

RISK CLASS FOR MEDICAL DEVICES	
Device Classification	**Common Specifications**
Class I Self Certified Rule 1	Not applicable

Ferno Canada Inc., declares that the above-mentioned products meet the provision of the following legislation:

- (EU) 2017/745 Medical Device Regulation (effective 5/26/2021)
- List other legislation/Directives as applicable:
  - MDD 93/42/EEC
  - ISO13485:2016 Medical Devices Quality Management System.

COMPANY REPRESENTATIVE: Sophia Hatsisavvas

TITLE: Director of Regulatory **SIGNATURE:** Sophia Hatsisavvas

PLACE: Mississauga Ontario, Canada **DATE:** February 25, 2022