

# EC DECLARATION OF CONFORMITY

**Basic UDI Nr : 5415174 183015692 99**

**Date Doc : 2020-08-06**

**Valid Date : 2025-08-06**

**The manufacturer or his authorized representative:**

VERMEIREN GROUP

**Address :**

Vermeirenplein 1/15

2920 Kalmthout

Belgium

**declares under his sole responsibility that the CE marked devices :**

Product group:	Ramps
ISO 9999:	18 30 15
Brand:	Vermeiren
Type:	692

**have been classified as class I, according to annex VIII (EU) MDR 2017/745, rule 1,**

**and is manufactured in full conformity with the European instructions below - including the latest modifications - and with the national law, that organizes this directions:**

Medical devices directive MDR 2017/745 Annex VIII

**and is in conformity with the relevant European harmonized standards:**

en 12182:2012

**City, Date:** Kalmthout, 2020-08-06

**Signature:**

A handwritten signature in black ink, appearing to read 'Vermeiren', written over a horizontal line.

**Name:** Patrick Vermeiren

**Function:** CEO