EC DECLARATION OF CONFORMITY

Basic UDI-DI : 5415174123009AlbatrosCL

Actor ID/SRN : BE-MF-000019750

The manufacturer:

VERMEIREN GROUP

Address :

Vermeirenplein 1/15

2920 Kalmthout

Belgium

declares under his sole responsibility that the CE marked devices :

Product group:	Patient hoist
Intended purpose:	Equipment for transferring and repositioning a person from a sitting position into an upright standing position; the equipment can be moved around freely; the body support consists of slings, a foot support and leg or knee support. (WHO ICF: d450, d460, d469, d498, d499)
ISO 9999:	12 36 04
EMDN:	V0805030102
Brand:	Vermeiren
Type:	Albatros

have been classified as class I, according to annex VIII (EU) MDR 2017/745, rule 13,

and is manufactured in full conformity with the European regulations and/or directives below - including the latest modifications - and with the national law, that organizes these directives:

MDR (EU) 2017/745, 2006/42/EG, RoHS 2011/65/EU, EMC directive 2014/30/EU, LVD 2014/35/EU

and is in conformity with applicable parts of following international standards:

EN ISO 10535:2006

City, Date: Kalmthout, 2024-07-10

Signature:

Name: Patrick Vermeiren

Function: CEO