



UK DECLARATION OF CONFORMITY

Declaration of Conformity #: DC01UK

Revision #: 1

This declaration of conformity is issued under the sole responsibility of the manufacturer and is represented by the undersigned.

Manufacturer:

MIP-UK Ltd,
2nd Floor, Clipper House, Leighton Industrial Park
Billington Road, Leighton Buzzard,
Bedfordshire, LU7 4AJ, England

hereby declares that the product

Type of product(s): **Patient Transfer and Slide sheets**

Basic UDI-DI: **7540289R175371639J**

GMDN Code: **37163**

Intended Purpose: A manual device used by healthcare professionals to assist in the physical movement and transfer of a person/patient (e.g., a hospitalized, a person with a disability, or a geriatric) from one position to another. The device typically has no lifting capabilities and utilizes sliding techniques to move the person. It is typically designed using low-friction artificial fibres and may be designed as an open roll, rather like "a sleeping bag", or a flat sheet. It is slid under the person to be moved, the person is gently pushed by the attending staff, and it rolls whilst the person glides on the rotating surface.

Model(s):

CS366-0001	CS441-0001	CS469-0011	CS469-0012
CS469-0013	CS469-0014	CS555-0001	CS567-0001
JC100-0001	JC100-0011	LP004-0007	LP004-RY07
LP010-0004	LP012-0207	LP025-0001	PGF100100
PGF150100	PGF200100	PGF20085	PGF220140

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PGT100100	PGT150100	PGT200100	PGT220140
PSF100100	PSF150100	PSF200100	PSF200140
PSF200140L	PSF20085	PSF20085UH	PSFH100100
PSFH150100	PSFH200100	PSFH200140	PSFH200140L
PSFH20085	PST11585	PST130100	PST14585
PST150100	PST200100	PST20085	PST220140
PST8585	PSTL150120	PSTL200140	PSTL200140L
PSTL20070	PTD-56/BA/LG/UK	PTD-56/BA/Y/UK	PTD-LS/S/BA1/Y/UK
PTD-LS/S/UK	RSF100100	RSF150100	RSF200100
RSF200140	RSF20085	RSFH100100	RSFH150100
RSFH200100	RSFH200140	RSFH20085	RST11585
RST130100	RST14585	RST200100	RST20085
RST220140	RST8585	RSTL200140	

was designed and manufactured in conformity with the following legislation(s)

- UK Medical Devices Regulations 2002
- UK General Product Safety Regulations 2005
- UK REACH Regulations 2021

The product(s) has been classified according to Annex IX of Directive 93/42/EEC, read with Directive 2003/12 and Directive 2005/50 as:

Class I Medical Device (non-invasive devices Rule 1)

Applicable standard(s):

- EN 15223-1:2016: Medical devices — Symbols to be used with medical device labels, labelling and information to be supplied
- BS EN 14971:2019: Medical devices. Application of risk management to medical devices

Company Certifications:

ISO 9001:2015	Certificate # 40662020	Issued by: QMS International
ISO 14001:2015	Certificate # 14130892	Issued by: QMS International

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Authorized Signature:

Jennifer Rivera

A handwritten signature in black ink that reads 'Jennifer Rivera'. The signature is written over a horizontal line that extends across the page.

QA/QC Manager

Leighton Buzzard, England

Place

2022-09-22

Date