PRESSALIT®

United Kingdom Conformity Assessment

United Kingdom RP:
Pressalit Ltd.
100, Longwater Avenue
Green Park
Reading RG2 6GP
United Kingdom
T: 01180923 2696
uk@pressalit.com

This product is produced in compliance with all relevant requirements of:
- Regulation (EU) 2017/745 on medical devices (Risk class I, rule 1)
- Council Directive 2006/42/EC on machinery
- Council Directive 2011/65/EC on RoHS

For the UK:
- Medical Device Regulation (MDR) 2002 on medical devices
- Supply of Machinery (Safety) Regulations 2008
- Restriction of the Use of Certain Hazardous Substances in Electrical and Electronic Equipment Regulations 2012

Medical devices - Application of risk management and quality management in design and production	ISO 14971:2019 ISO 13485:2016

The products are manufactured in accordance with the following standards:	IEC 60601-1:2005 ISO 17966:2016

Ry, 27. maj 2026

Mette Elstrøm Kristoffersen, QA/RA Specialist, Pressalit A/S
Responsible

Manufacturer: Pressalit A/S - Pressalitvej 1 DK-8580 Ry - T: +45 8788 8788 - pressalit@pressalit.com

CT 4000

REF **R8584572000B**

EAN (GTIN):	5708590400563
Nomenclature (GMDN):	43600
Nomenclature (EMDN):	Y093399

	Name
EN	CT 4000 change table, outer measurements 1008 x 1829 mm, lying surface in compact laminate, 813 x 1829 mm, white. With safety rail and body belt, manually foldable, electrically height adjustable with operating buttons in table front, back-up battery, wall-mounted

	Intended purpose
EN	This product is intended for changing a person who is unable to stand or sit. The person is lying horizontally on the table. Expected service life is 10 years for institutional use, provided that maintenance is carried out as described in the user manual.