

AdaptAir™

The Original Adaptive Therapy System

Warranty & Standards

1. Warranty

The AdaptAir™ Air Mattress System carries a comprehensive warranty as detailed below. Please keep proof of purchase for proof of warranty commencement.

2 YEARS WARRANTY

Incl. Pump, mattress, cells

**A 2 Year blanket warranty is provided over all foam components, NE-7 control unit (Pump), mattress cover and cells.*

Studio Care does not warrant against excessive or incorrect use, modification or any situation that could be deemed as fair wear and tear. This a return to base (RTB) warranty and does not cover freight and transport costs pertaining to the return of any items under warranty. For more information please contact Studio Care. Studio Care will not warrant the safety and or correct functioning of products where any original component have been changed or modified by non-Studio Care approved and trained service & maintenance staff or external providers. Safety is not guaranteed where components have been replaced with non-original Studio Care parts. All mattress customisation, modifications, and or alterations made to this product are considered as "Custom" changes. These may cause the product to no longer meet safety standards and should be considered prior to commencement. Any customisation or modification of the air mattress is not warranted by Studio Care and will void warranty. If any faults are detected upon receipt of this product please phone Studio Care. Any faults that are detected during normal use should be reported to Studio Care immediately to determine if warranty conditions apply and if so, the necessary repair or replacement work to be completed. Spare parts lists are available upon request. For servicing, preventative maintenance and any other questions regarding this or any Studio Care product, please contact Studio Care.

2. Standards

The air mattress systems components are manufactured in ISO compliant production facilities in Taiwan and Australia to strict quality control standards.

Compliance:

210681	175810
ISO-13485	ISO-9001

Symbol	Definition of Symbol
⚠	Attention: See Instructions for use
CE	CE Marking indicating conformance to EC Directive No. 2007/47/EC concerning medical devices
██	Type BF Applied Part (patient isolation from electrical shock)
██	Class II Product
██	Operation Instructions
██	Indicates separate collection for electrical and electronic equipment (WEEE)
IP21	Protection against finger and dripping water