1. General

1. Product Description
The Dyna-Tek Intelligent-Air pressure reducing cushion has 4 individually designed internal foam cores, each encapsulated in a soft thermoplastic polyurethane air cell. Three of the cells are joined together to allow the air to flow between them and aid pressure redistribution in reaction to human body movement. The fourth air cell, situated under the sacral bony prominences (ischial tuberosities) has a unique pressure reducing valve system which reduces the internal air pressure in relation to each person’s body anatomy. The cushion requires no external pump inflation. The Covers are made from a water resistant, two way stretch, vapour permeable material that have been designed to reduce the build-up of both heat and moisture. Please see the product literature for further details.

1.1 Symbols

Warnings
In this User Manual warnings are indicated by symbols. The warning symbols are accompanied by a heading that indicates the severity of the danger.

⚠️ WARNING
Indicates a hazardous situation that could result in serious injury or death if it is not avoided.

⚠️ CAUTION
Indicates a hazardous situation that could result in minor or slight injury if it is not avoided.

⚠️ IMPORTANT
Indicates a hazardous situation that could result in damage to property if it is not avoided.

1.2 Symbols guide

- **DO NOT PIERCE OR CUT**
- **LINE DRY**
- **DO NOT DRY CLEAN**
- **DECLARATION OF CONFORMITY**
- **RECOMMENDED 80°C**
- **DO NOT PUT NEAR FLAME**
- **USER WEIGHT LIMIT 152KG**
- **DO NOT IRON**
- **DO NOT BLEACH**
- **TUMBLE DRY ON LOW MAX 40°C**

2. Usage

2.1 Intended use
The Dyna-Tek range of Pressure Redistribution Cushions are designed for both prevention and treatment of pressure ulcers and are intended to be used in conjunction with an appropriately sized chair / wheel chair, as part of an overall pressure ulcer prevention program of care.

⚠️ WARNING Which cushion is suitable for your individual use should be decided by your clinician who will also determine the most appropriate pressure ulcer prevention or treatment plan and the relevant nursing procedures.

⚠️ WARNING The following user manual should be used in conjunction with the product literature and will give you instructions on how to get the best results when using one of our pressure reducing cushions, however should you require any additional information please contact the relevant Sales Office on (+44) 0845 4599831.

2.2 On Delivery of your new cushion

⚠️ IMPORTANT Open the outer packaging a period of a minimum of 1 hour before use to optimise performance. Please dispose of all packaging in line with your relevant environmental policy.

2.3 Using your cushion

⚠️ CAUTION Prior to using the cushion ensure that the chair has no rough edges that may damage the cushion. Ensure the zip is closed and the zip flap is completely covering the zip. Avoid contact with sharp objects that may damage the cushion.

⚠️ WARNING The usage of additional layers between the cushion and the patient, such as sheepskins or other similar items may reduce the effectiveness of the pressure reducing properties of the cushion or indeed lead to an additional increase of body temperature, which may cause skin maceration and is therefore not advisable. Please seek clinical advice should you wish to use any additional covering other than the cover which has been supplied with the cushion by the manufacturer.

Take care when using mechanical items such as bridging boards, PAT slides or other patient transfer aids that damage does not occur to the cushion due to sharp or damaged edges.

⚠️ WARNING The maximum user weight of your cushion is dependent on the cushion correctly fitting the patient. Ensure the patient buttocks do not overhang the edges of the cushion whilst seated as this would mean the patient is too wide for the specific size cushion. Other cushion sizes are available.

⚠️ WARNING Ensure the cushion is placed the correct way up and in the correct rotation. Please follow the labelling instructions on your cushion at all times. Do not use the cushion without the cover that has been supplied by the manufacturer.
2.4 Patient repositioning

IMPORTANT Your Dyna-Tek Pressure Redistributing Cushion will provide not only the best pressure re-distribution properties but also give greater patient comfort and product longevity. However to prevent the build-up of pressure which may lead to tissue damage and potential ulcer formation, it is important that the patient either repositions themselves, or is repositioned on a regular basis. This should be based on the clinical judgement of a qualified healthcare professional who will provide you with a suitable turning regime.

3. Cleaning & Care

The way in which the cushion is cleaned depends upon the nature of the contamination and the susceptibility of the patient.

3.1 Light Soiling

IMPORTANT In the absence of gross contamination or unusual risk, the removal of dirt and spills using clean, warm water, neutral detergent and manual dexterity should be sufficient. However if required the cover can be wiped down with a 0.1% Hypochlorite Solution (1,000ppm). Ensure that the cover is rinsed with clean water using a single use non-abrasive cloth, and thoroughly dried. Ensure adequate ventilation.

3.2 Heavy Soiling

IMPORTANT In cases of gross contamination the cushion cover should be cleaned first with detergent and water then with a diluted solution of 1% Hypochlorite (10,000ppm). Rinse well with clean water and a damp single use non-abrasive cloth. Ensure adequate ventilation. Disposable gloves and aprons should be worn. If splashing could occur, eye/face protection should also be worn. On removal of gloves, hands should be washed.

3.3 Drying

IMPORTANT Ensure cushion cover is thoroughly dried before placing it back onto the cushion. Damp humid environments have been shown to lead to the continuation of bacterial growth. Within the community setting, it may be desirable to launder the cover after removing from the foam core. In such cases the cover should be laundered at no more than 80 degrees C. To avoid shrinkage of the cover only tumble dry on a low heat setting not exceeding 40 degrees C and not for longer than 10 minutes. Covers must be thoroughly dried before re-fitting to the cushion.

3.4 Inspection

WARNING Inspect the inner and outer surfaces of covers and their zip fasteners regularly for signs of damage. If the cover is stained, soiled or torn, the air cells should also be examined for contamination. Damaged/soiled covers and air cells should be reported to the ward/department manager. If the core of the cushion is wet or badly stained, the cushion should be withdrawn from service. Check that air cells are not punctured or damaged. Check that the patient is on a layer of air whilst seated by pushing down on the outer edges of each air cell. We recommend that the chair should also be cleaned in between each change of patient.

4. MAINTENANCE

4.1 Audit

CAUTION It is recommended that a minimum of a 6 monthly or annual audit of all cushions within a ward/department should be adhered to. In the community setting, an audit should also be considered at similar periods and between each change of patient. Details on how to audit your cushion, including covers, air cells and pressure valves are available from Direct Healthcare Services.

5. AFTER USE

5.1 Storage

CAUTION Cushions can become damaged when stored incorrectly. It is advisable to store your cushion off the floor in a clean dry environment. Do not store your cushion near heating devices such as electrical fires or radiators. Wherever possible, cushions should be stored within a protective cover.

5.2 Re-Use

IMPORTANT A cleaning record must be kept as part of the cleaning system. The product is suitable for repeated use. The number of times it can be used depends on how often and in which way the product is used. Before reuse, clean the product thoroughly in accordance with the guidelines.

5.3 Disposal

IMPORTANT The disposal and recycling of used devices and packaging must comply with the applicable legal regulation in each country.

6. QUALITY & TESTING

6.1 Service Life

We estimate a life expectancy of four years for these products, provided they are used in strict accordance with the intended use as set out in this document and all maintenance and service requirements are met.

The estimated life expectancy can be exceeded if the product is carefully used and properly maintained, and provided technical and scientific advances do not result in technical limitations. The life expectancy can also be considerably reduced by extreme or incorrect usage. The fact that we estimate a life expectancy for these products does not constitute an additional warranty.

6.2 Product Warranty Periods

Please refer to the product literature for your cushion warranty period. The warranty of your Dyna-Tek cushion is valid from time of shipping. If a defect or fault is discovered please contact relevant Sales Office.

6.3 Fire Testing


6.4 Quality Standards

Quality is fundamental to the company’s operation, working within the disciplines of BS EN ISO 9001:2008 and ISO 13485:2012. All Direct Healthcare Cushion products feature the CE mark, in compliance with the Medical Device Directive 93/42/EEC.

Every effort has been made to ensure that the contents of this publication are fully up-to-date at the time of going to print. As part of our continuous improvement process, Direct Healthcare Services reserves the right to modify existing models at any time.

Sales Offices

UK & Europe
Direct Healthcare Services Ltd.
6 – 10 Withey Court, Western Industrial Estate,
Lon-y-Llyn, Caerphilly, CF83 1BF, UK
T: +44 (0) 845 459 9831
info@directhealthcareservices.co.uk
www.directhealthcareservices.co.uk

Asia Pacific
Direct Healthcare Services PTY Ltd.
PO Box 562, Wembley, Western Australia 6913
T: +61 (0) 423 852 810
info@directhealthcareservices.com.au
www.directhealthcareservices.com.au