





ROMA[™] MATTRESS COVER – STANDARD / COTTON EDGE / NIMBUS

This guide covers all products with the prefix RRMC in the product code.

With or without assistance this product makes moving and/or repositioning in bed that much easier. The user can lie on the **Roma™ Mattress Cover (Roma™)** directly or use with a bed sheet placed over it. Model shown below is **Roma™** with Cotton Edge on a divan bed but they fit standard hospital beds also. The techniques shown apply to all types of our **Roma™**.

USER INSTRUCTIONS





PERSON UNASSISTED

Remove the **Roma™** from its packaging and place over the bed mattress, as shown.

The user can now sit on the edge of the bed, prior to rotating on the **Roma**^{\mathbb{M}}.



The person can now reposition themselves accordingly and sleep comfortably without risk of tissue damage or sweating. The **Roma™** fabric enables them to turn more easily.





PERSON ASSISTED (a Flat Slide Sheet is also required) With the Roma[™] placed over the mattress, carers can reposition the person by inserting a Flat Slide Sheet, as shown (on top of the Roma[™] but under the patient).

If lying on a bed sheet, insert the **Flat Slide Sheet** between the **Roma™** and bed sheet and remove after repositioning.

Unroll the **Flat Slide Sheet** under the patient without disturbing them. The **Flat Slide Sheet** can then be used to reposition the person sliding on the **Roma™** below.







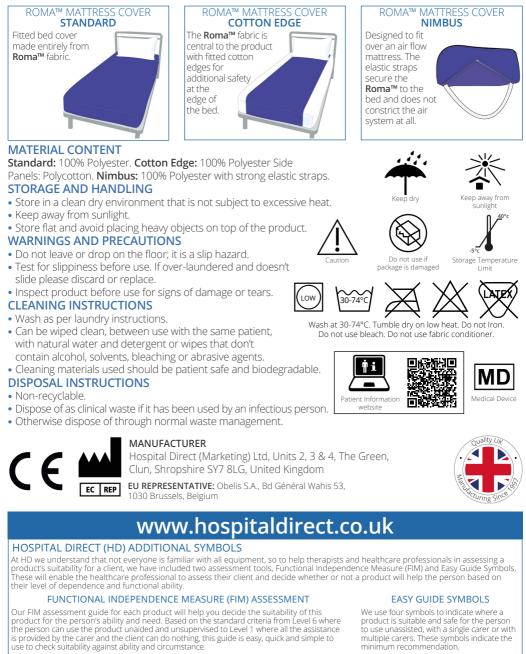
Then, when the person is in the desired position the **Flat Slide Sheet** removes effortlessly...



...leaving the person in exactly the position you want them to be in.

Alternatively use under a standard bed sheet and use a **Flat Slide Sheet** under the bed sheet for all repositioning needs. Remove the **Flat Slide Sheet** once the person is in the desired position.







HD has a policy of continuous development and, as such, reserves the right to alter specifications (including measurements, materials and colours) without prior notice. If any serious incident has occurred in relation to the device it should be reported to the manufacturer and the competent authority in which the user and/or patient is established.