ACTIVON BARRIER CREAM PRODUCT EVALUATION

Please follow the Information for Use (IFU) for indications for use, including Activon Barrier Cream NOT being used on patients allergic to bee venom, honey or cocoa butter.

Contra-Indications:

Do NOT use on infected area of skin, open wounds or deep puncture wounds, where there are any known allergies or sensitivities to the ingredients or on neonates (less than one year old).

Warnings and Precautions:

Do NOT use on uncovered areas of skin during phototherapy treatment.

PART 1 - Pre Treatment

Date:

Healthcare Professional Name:

Healthcare Professional Job Title:

Place of Work (please select): Acute Community Nursing Home Hospice

Other - please specify area:

Data Protection

The information obtained from this evaluation will be entered into a database which be managed by Advancis. All data storage will follow all legal obligations relevant to Advancis. No data will be disclosed or transferred to a third party and if you have any further questions please contact your local Advancis representative.

Consent gained from patient: Yes No (please obtain)

Patient ID Number: Age:

Sex: Male Female Prefer not to say

PART 2 - Initial Assessment

Is the patient allergic to bee venom, honey or cocoa butter?

Yes No

If yes, do not continue with this evaluation.

Co-Morbidities and Relevant Clinical History

Please provide details:

Initial Skin Status

Is MASD (Moisture Associated Skin Damage) present?

Yes No

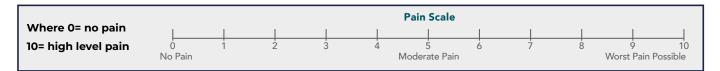
Is the skin irritated?

Yes No

Is the skin vulnerable?

Yes No.

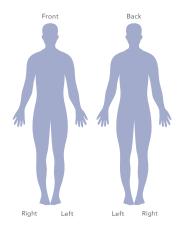
Did the patient experience pain on application of previous regime? Yes No If the patient experiences pain in the at risk area, please record the level of pain on a scale of 1-10:



Location of the At Risk Area

Using the body outlines on the right hand side, please mark the location of the at risk area.

Please specify in words the location of the at risk area:



Previous Management

Product name:

How many daily applications:

PART 3 – Regime Change

Instructions for Use:

1. Ensure that the skin is clean and dry prior to application. 2. Apply a thin, uniform coating over the whole area to be protected, or where the ostomy pouch, adhesive device or dressing will be applied. 3. Pea to Palm application: Use a pea sized amount of cream to cover an area approximately the size of your palm. Do NOT over-apply. 4. Remove any excess with a clean tissue. 5. If application, once fully dry, apply the ostomy pouch, adhesive device or dressing as normal. 6. Reapplication is recommended as and when required. When used with adhesive dressings or devices, reapply at each dressing or device change.

How many applications per day:

Did the patient experience pain on application or during use? Yes No

If the patient experiences pain related to the wound, please record the level of pain on a scale of 1-10:



Please rate Activon Manuka Honey Barrier Cream

Criteria	Excellent	Good	Fair	Poor
Ease of application				
Drying time				
Longevity on skin				
Patient satisfaction				
Skin condition of application area (before using Activon Manuka Honey Barrier Cream)				
Skin condition of application area (after using Activon Manuka Honey Barrier Cream)				
Overall performance				

Please describe any skin changes below:

Did you experience any incidences of faecal incontinence while using this product?		Did the skin feel moisturised after using this product?			
Yes	No	Yes	No		
Did you experience any stoma leakage on at risk area while using this product? Yes No		Did you experience any, or worsening of, Moisture Associated Skin Damage while using this product?			
		Yes	No No		
Did you experience any incidences of urinal incontinence while using this product?		Did you experience any skin maceration while using this product?			
Yes	No	Yes	No		
		res	NO		
Did you experience any incidences of sweat on at risk area while using this product?		Did this product provide an effective moisture barrier?			
Yes	No	Yes	No		
• .	perience any incidences of at risk area while using this				
Yes	No				
Please describe any further comments below:					
PART 4 – C	onsent				
Please sign and date below to confirm consensus.					
Clinician					
Print name:					
Signature:					
Date:					
Thank you f	for your participation in the evaluat	ion.			