















Product Information Sheet

Biatain Ibu Non-Adhesive Foam / Biatain Ibu Soft-Hold Foam			
Classification	Pain Control Dressing: Ibuprofen		
British Columbia Practice	Consult with NSWOC / Wound Clinician / Physician / NP prior to use.		
Key Points	 Polyurethane foam dressing containing ibuprofen (0.5 mg/cm²) dispersed throughout dressing. Soft, conformable and highly absorbent. Provides an optimal moist wound healing environment and effective exudate management. Ibuprofen is released into the wound bed when in contact with wound exudate. Semi-permeable film backing is waterproof and provides a bacterial barrier. Two versions available: Biatain Ibu Non-Adhesive Foam Dressing is suitable for use on fragile skin due to absence of adhesive. Biatain Ibu Soft-Hold Foam Dressing has a partially adherent layer enabling dressing to stay in place while secondary dressing or compression therapy is applied. Must be in contact with wound bed and will conform to wound depth of up to 2 cm. 		
Indications	 Primary dressing but does not require a secondary dressing. Painful wound. Biatain Ibu Non-Adhesive Foam dressing can be used on donor sites. Can be used in combination with compression therapy. 		
Precautions	 Clinically infected wounds must be treated prior to use of Biatain Ibu Foam. Do not exceed 1200 cm² (e.g. 3 dressings of 20 x 20 cm each) at each dressing change. Dressing should not be changed more than twice daily corresponding to maximum daily use of 2400 cm². Can be used for up to 6 weeks as long as clinically indicated. Remove dressing prior to radiation treatment or examinations (including, x-rays, ultrasonic treatment, diathermy, microwaves or MR scanning). Has not been evaluated on lactating individuals, consult with physician / NP prior to using on this population. 		
Contraindications	 Sensitivity or allergy to ibuprofen, any other components of the dressing, acetylsalicylic acid or other NSAIDs. Untreated clinically infected wounds. Do not use Biatain Ibu Soft-Hold Foam dressings on donor sites. Do not use with oxidizing solutions (e.g. hypochlorite and hydrogen peroxide solutions). Pregnant individuals. Children under 12 years of age except under the direction of a physician / NP. 		
Formats & Sizes	Biatain Ibu Non-Adhesive Foam		

Directions	Rationale / Key Points
Selection	
Select dressing size where foam pad overlaps wound edge by minimum of 2 cm. Dressing can be cut to size if needed.	For small sized dressings an overlap of 1 cm may be necessary. Incorrect sizing of dressing will adversely affect dressing function.
Preparation	
Cleanse wound and periwound / surrounding skin with sterile	The safety of other cleansing agents in combination with

















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Directions	Rationale / Key Points
normal saline or water.	Biatain Ibu Foam has not been demonstrated.
Dr. a arive and I average adia a skip	See Wound Cleansing Procedure or QR Code below.
Dry periwound / surrounding skin.	Ensure that solution has dried prior to dressing
	application to minimize risk of periwound maceration.
If required, apply barrier film to periwound skin.	To protect periwound skin from moisture associated
	skin damage and medical adhesive related skin injury.
Application	
If there is minimal exudate, moisten Biatain Ibu Foam with	Ibuprofen is released to wound bed when in contact
sterile saline or apply amorphous wound gel to wound bed.	with wound exudate.
For wounds with depth 2 cm or less place dressing directly	Must be in contact with wound bed and will conform to
over the wound with the plain, non-printed side toward the	wound depth of up to 2 cm.
wound.	The outer side prevents strike through of exudate.
Secure with tape or wrap with kling.	The non-printed side will release Ibuprofen into wound.
Removal	
Consider using adhesive remover to remove adhesives (e.g.	To decrease risk of medical adhesive related skin injury
tape).	(MARSI).
Gently lift corners of dressing away from wound.	To avoid trauma to wound and periwound skin
Frequency of Dressing Change	
May be left in place for up to 7 days if pain is effectively	Dressing change frequency is dependent on amount of
managed. Usual wear-time is 2- 3 days.	wound exudate.
Change dressing when clinically indicated or when visible signs	Absorbed exudate is clearly visible through backing of
of exudate approach edge of dressing.	dressing.
Expected Outcomes	
Wound pain is managed for 2-3 days	If product does not perform as expected notify
Doodlook and a war and a sure at a d	NSWOC/Wound Clinician and consider submitting a
Product performs as expected.	Supply Chain Product Concern form.
OP Codes	

QR Codes



Wound Cleansing

For further information please contact NSWOC/Wound Clinician