

NHSGGC Joint Wound Care Formulary

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Foreword

This NHSGGC Joint Wound Care Formulary, and the accompanying wound management data sheets have been developed as a guide to aid Healthcare Professionals in selecting the most appropriate dressings/products. The Wound Management Formulary was updated to reflect the changes made to the National Procurement (NP) Contract.

The formulary was developed by the Wound Formulary Development and Implementation Group (WFDIG), this group has representation from all specialties involved with wound management in Acute and Primary Care Sectors. The WFDIG included products to ensure that the majority of wound management goals can be achieved by prescribing products from this formulary.

To ensure appropriate person centred wound care it is essential that wounds that have not progressed in a two week period must be referred to the appropriate wound care specialist: Dermatology, Podiatry, Vascular, Burns and Plastics, Tissue Viability.

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Prescribing Guidance

- Wounds that have not progressed in a two week period should be referred to the appropriate wound care specialist: Podiatry, Vascular, Burns and Plastics, Tissue Viability.
- Any patient who is receiving ongoing wound management must have an approved NHSGGC wound assessment chart completed and updated at least weekly.
- Practitioners should select products included in the formulary and only use nonformulary products when there is sound clinical rationale for doing so.
- Prescribers should take into account the volume and duration of products prescribed and maintain a two week challenge/review/reassessment of wounds where appropriate.
- Basic wound dressings should be considered for non-complex wounds or for use as secondary dressings. If frequent dressing changes are required a cost effective dressing should be used or seek advice from wound care specialist
- Dressing price can rise significantly with increasing size, so the smallest size dressing that is appropriate to the wound (allowing for any necessary overlap onto healthy skin) should be selected.
- Wound dressings containing an antimicrobial should only be used on the small number of patients who need them – the Health Technology Assessment Report 13 (Dec 2015) on the use of antimicrobial wound dressings for chronic wounds highlighted the lack of evidence for their routine use.
- Wear time varies between products, and will also depend on the both patient and wound related factors.
- If clinicians wish to have a new/different product considered for inclusion on the formulary (or to provide feedback on current products) a non-formulary/product evaluation form must be completed.

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NHSGGC Joint Wound Care Formulary Product List

The list of formulary products are listed by category. Not all products are available to both Acute and Primary care and are listed as:

PC - available in Primary Care (community)

Acute - available in Acute

(S) - These products can only be initiated after review by a wound care specialist (Dermatology, Vascular, Podiatry, or Tissue Viability Nurse (TVN))

Category	Product	Page	PC	Acute
Adhesive Tape	CLINIPORE	11	х	
Adhesive Tape	PRIMAFIX PERMEABLE 10M	12	х	
Antimicrobial	ACTIVHEAL AQUAFIBRE AG	13		х
Antimicrobial	AQUACEL AG+ EXTRA/RIBBON	14	х	
Antimicrobial	CUTIMED SORBACT SWABS/RIBBON	16	х	х
Antimicrobial	INADINE	17	х	х
Antimicrobial	IODOFLEX PASTE	18	х	
Antimicrobial	IODOSORB OINTMENT	19	х	
Antimicrobial	PRONTOSAN GEL X	20	х	х
Antimicrobial	URGOTUL SILVER	21	х	х
Burns basic dressing	JELONET	22	х	x
Cleansing	CLINIPODS	23	x	x
Cleansing	PRONTOSAN SOLUTION	24	x	X
Cleansing	STERICLENS AEROSOL 240ML	25	х	
Debridement Dressing	ACTIFORM COOL	26	x	x
Debridement Dressing	HYDROCLEAN ADVANCE	27	х	х
Debridement Dressing	URGOCLEAN PAD	28	х	х
Debridement Gel	FLAMINAL FORTE	29	х	х
Debridement Gel	FLAMINAL HYDRO	30	х	х
Debridement Hydrogel	ACTIVHEAL HYDROGEL	31	х	х
Debridement /Modulating matrix	ACTIVHEAL AQUAFIBRE EXTRA	32	х	х
Debridement / Occlusive	DUODERM EXTRA THIN	33	х	х
Debridement Physical	DEBRISOFT	34	х	х
Debridement Physical	UCS DEBRIDEMENT	35	х	Х
Dressing pack	DRESSIT	36	x	
Dressing pack	NURSE IT	36	х	
Haemostat	COVAWOUND ALGINATE	37	x	x
Honey -antibacterial	ACTILITE	38		

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Honey -antibacterial	ACTIVON TUBE	39	х	
Honey -antibacterial	ACTIVON TULLE	40	х	
Honey-antimicrobial	L-MESITRAN	41		х
Honey-antimicrobial	L-MESITRAN SOFT	42		X
Modulating Matrix (S)	URGOSTART	43	XS	XS
Modulating Matrix (S)	PROMOGRAN PRISMA	45	XS	xs
Non-adherent contact layer	ATRAUMAN	47	Х	х
Non-adherent contact layer	N-A ULTRA	47	x	х
Odour control	CARBOFLEX	49	x	x
Odour control	CLINISORB	50	X	X
Radiotherapy RTOG damage	POLYMEM	51	х	х
Secondary dressing	ABSOPAD	52	х	
Secondary dressing adhesive	ACTIVHEAL SILICONE FOAM ADHESIVE	53	х	
Secondary dressing non adhesive	ACTIVHEAL SILICONE FOAM BORDERLESS	54	х	
Secondary dressing	CELLUDRESS	55	х	
Secondary dressing tubular bandage	COMFIFAST TUBULAR BANDAGE	56	х	
Secondary dressing adhesive film	HYDROFILM	57	х	
Secondary dressing adhesive film	HYDROFILM PLUS	58	х	
Secondary dressing adhesive silicone foam	KLINIDERM FOAM SILICONE BORDER	59	х	х
Secondary dressing non-adhesive silicone foam	KLINIDERM FOAM SILICONE NON-BORDER	60	х	х
Secondary dressing lightweight bandage	KNIT BAND	61	х	
Secondary sub bandage absorbent	K SOFT	62		
non-woven wadding	PREMIERPORE	63	X	
Secondary dressing		64	X	
Secondary dressing non adhesive	TELFA		X	
Silicone foam (S)	BIATAIN SILICONE	65	XS	XS
Silicone wound contact layer	KLINIDERM WOUND CONTACT LAYER	67	х	х
Superabsorber (S)	CUTIMED SORBION SACHET EXTRA	68	XS	XS
Superabsorber	KLINIDERM SUPER ABSORBENT	69	х	х
Superabsorber	VLIWASORB PRO	70	х	
Superabsorber	ZETIVIT E NON-STERILE	71	х	х
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Drug Tariff Pricing for Primary Care and Pecos code for Acute

Category	Product	Sizes					
Adhesive tape (PC only)	CLINIPORE	1.25 x 5	2.5 x 5	5 x 5			
	CLINIPORE	£0.36	£0.61	£1.02			
Drug tariff Adhesive tape (PC only)	PRIMAFIX PERMEABLE x 10m	5	10	15	20		
Drug tariff		£1.70	£2.49	£3.67	£4.52		
Antimicrobial (Acute only)	ACTIVHEAL AQUAFIBER AG	5x5	10 x 10	15 x 15	2.7 x 32		
PECOS Code	ACTIVITEAL AQUAFIBER AG	5 X 5	10 X 10	15 X 15	2.7 X 32		
FLCOS CODE	AQUACEL AG+						
Antimicrobial (PC only)	EXTRA/RIBBON	5 x 5	10 x 10	15 x 15	1 x 45		
Drug tariff		£2.10	£4.99	£9.40	£3.26		
	CUTIMED SORBACT			3cm round			
Antimicrobial	SWAB/RIBBON	4 x 6	7 x 9	x5	2 x 50		
Drug tariff		£1.76	£3.75	£3.51	£4.30		
PECOS Code		178988	178995	178971	178964		
			9.5 x				
Antimicrobial	INADINE	5 x 5	9.5				
Drug tariff		£0.34	£0.50				
PECOS Code		39654	39661	17			
Antimicrobial (PC only)	IODOFLEX (paste)	5g	10g	17g			
Drug tariff		£4.39	£8.77	£13.89			
Antimicrobial (PC only)	IODOSORB (ointment)	10g					
Drug tariff		£4.85	250				
Antimicrobial	PRONTOSAN GEL X	50g	250g				
Drug tariff		£12.44	£33.30				
PECOS Code		193585	171743				
Antimicrobial	URGOTUL SILVER	10 x 12	15 X 20 £9.30				
Drug tariff PECOS Code		£3.28	242542				
		242559	242542				
Burns basic dressing Drug tariff	JELONET	10 X 10 £0.44					
PECOS Code		17409					
FECOS CODE		20ml x					
Cleansing (PC only)	CLINIPODS	25					
Drug tariff		£4.40					
Cleansing	PRONTOSAN SOLUTION	350ml					
Drug tariff		£5.09					
PECOS Code		133895					
Cleansing (PC only)	STERICLENS AEROSOL	240ml					
Drug tariff		£3.19					
Debridement dressing	ACTIFORM COOL	5 x 6.5	10 x 10	10 x 15	20 x 20		
Drug tariff		£1.91	£2.81	£4.03	£8.45		
PECOS Code		117123	84159	93205	D		
				5.5	7.5 x		
Debridement dressing	HYDROCLEAN ADVANCE	3 round	4 round	round	7.5	10 x 10	
Drug Tariff		£2.38	£4.06	£4.82	£5.33	£6.04	
PECOS code		264674	264636	264643	264650	264667	
Debridement dressing	URGOCLEAN PAD	6 x 6	10 x 10	15 x 15			
Drug Tariff		£2.01	£2.24	£4.05			
PECOS code		214761	210510				
			210310				
Debridement gel	FLAMINAL FORTE	15g					

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Drug Tariff		£8.16						
PECOS code		138067						
	FLAMINAL HYDRO							
Debridement gel		15g						
Drug Tariff		£8.16						
PECOS code		138074						
Debridement Hydrogel	ACTIVHEAL HYDROGEL	15g						
Drug tariff		£1.43						
PECOS Code		130474						
Debridement / Modulating matrix	ACTIVHEAL AQUAFIBER EXTRA	5 x 5	10 x 10	15 x 15	2.5 x 30.5			
Drug tariff		£0.61	£1.46	£2.74	£1.76			
PECOS Code		136322						
Debridement and	DUODERM EXTRA THIN	7.5 x						
Occlusive		7.5	5 x 10	10 x 10	9 x 15	9 x 25	9 x 35	15 x 15
Drug Tariff		£0.85	£0.80	£1.41	£1.88	£3.00	£4.20	£3.03
PECOS code		34840	34857	31658	165445	165452	165469	31665
Debridement Physical	DEBRISOFT	10 x 10						
Drug Tariff		£6.86						
PECOS code		217106						
Debridement Physical	UCS DEBRIDEMENT	pad						
Drug Tariff		£3.41						
PECOS code		217106						
Dressing pack (PC only)	DRESSIT	pack						
Drug tariff		£0.69						
Dressing pack (PC only)	NURSE IT	pack						
Drug tariff		£0.81						
Haemostat	COVAWOUND ALGINATE	5 x 5	10 x 10	10 x 20	15 x 15	15 x 20	2 x 30	
Drug Tariff		£0.44	£0.74	£1.88	£2.51	£3.18	£1.42	
PECOS code		268030	268047	268054			268061	
Honey -antibacterial	ACTILITE	5 x 5	10 x 10	10 x 20	20 x 30	30 x 30	30 x 60	
Drug Tariff		£0.72	£1.25	£2.41	£6.78	£11.22	£19.39	
PECOS code		197392	138104	182435				
Honey -antibacterial (PC only)	ACTIVON TUBE	25g						
Drug tariff		£2.80						
Honey -antibacterial (PC								
only)	ACTIVON TULLE	5 x 5	10 x 10					
Drug tariff Honey-antimicrobial		£2.28	£3.76					
(Acute only)	L-MESITRAN OINTMENT	20g	50g					
PECOS Code		£3.91	£9.90					
Honey-antimicrobial	L-MESITRAN SOFT							
(Acute only)		15g						
PECOS Code		£3.62	40.10	45 00				
Modulating Matrix (S)	URGOSTART	6 x 6	10 x 10	15 x 20				
Drug Tariff		£4.70	£6.50	£11.68				
PECOS code		20.011	216789					
Modulating Matrix (S)	PROMOGRAN PRISMA	28CM	123CM					
Drug Tariff		£6.51	£18.54					
PECOS code Non-adherent contact		117697						
layer	ATRAUMAN	5 x 5	7.5 x 10	10 x 20	20 x 30			
Drug Tariff		£0.36	£0.37	£0.84	£2.31			
PECOS code		230495	233182	52301				

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Non-adherent contact	N-A ULTRA	9.5 x						
layer		9.5	9.5 x 19					
Drug Tariff		£0.34	£0.65					
PECOS code		25374						
Odour control	CARBOFLEX	10 x 10	8 x 15	15 x 20				
Drug Tariff		£3.42	£4.10	£7.78				
PECOS code		117277	49646	34871				
Odour control	CLINISORB	10 x 10	10 x 20	15 x 25				
Drug Tariff		£2.01	£2.68	£4.31				
PECOS code		24728	24735	173389				
Radiotherapy RTOG damage only	POLYMEM NON-ADHESIVE	10 x 10	17 x 19	10 x 61 roll				
Drug Tariff		£2.52	£6.20	£13.36				
		242290						
PECOS code Secondary dressing (PC		242290	242313	242320				
only)	ABSOPAD	5 x 5	10 x 10	20 x 10				
Drug tariff		£0.70	£0.13	£0.28				
Secondary dressing	ACTIVHEAL SILICONE FOAM	7.5 x		12.5 x				
adhesive (PC only)	ADHESIVE	7.5	10 x 10	12.5	15 x 15	20 x 20		
Drug tariff		£1.08	£1.60	£2.35	£2.87	£4.97		
Secondary dressing non adhesive (PC only)	ACTIVHEAL SILICONE FOAM BORDERLESS	5 x 5	7.5 x 7.5	10 x 10	10 x 20	15 x 15		
Drug tariff	DONDENELOS	£0.84	£0.98	£1.45	£2.69	£2.67		
Secondary dressing (PC		10.84	10.98	11.45	12.09	12.07		
only)	CELLUDRESS	10 x 10	10 x 15	10 x 20	15 x 20	20 x 25		
Drug tariff		£0.19	£0.20	£0.22	£0.30	£0.40		
Secondary dressing tubular bandage	COMFIFAST TUBULAR BANDAGE	3.5 x 1	E v 1	7.5 x1	10.75 x 1	17 E v 1		
	BANDAGE		5 x 1			17.5 x 1		
		red	green	blue	yellow	beige		
Drug tariff		£0.56	£0.58	£0.77	£1.20	£1.83		
Secondary dressing adhesive film	HYDROFILM	6 x 7	10 x 12.5	10 x 15	10 x 25	12 x 25	15 x 20	20 x 30
Drug tariff		£0.24	£0.45	£0.56	£0.87	£0.92	£1.04	£1.71
PECOS code		216727	216734					216758
Secondary dressing								
adhesive film (PC only)	HYDROFILM PLUS	7.2 x 5	9 x 10	9 x 15	10 x 20	10 x 25	10 x 30	
Drug tariff		£0.19	£0.29	£0.32	£0.49	£0.65	£0.73	
PECOS code		171217		171231	171255		171262	
Secondary dressing adhesive silicone foam	KLINIDERM FOAM SILICONE	7.5 x	10 x 10	12.5 x	15 x 15	10 × 20	10 x 20	15 x 20
	BORDER	7.5	10 x 10	12.5	15 x 15	10 x 20	10 x 30	15 x 20
Drug Tariff		£0.94	£1.23	£1.79	£2.70	£3.15	£4.76	£4.65
PECOS code Secondary dressing non		216659	216604	216673	216680	216697	237388	216703
adhesive	KLINIDERM FOAM SILICONE NON BORDER	5 x 5	10 x 10	10 x 20	15 x 15	20 x 20		
Drug Tariff		£0.78	£1.68	£2.58	£3.11	£4.44		
PECOS code		216598	216604	216611	216628	216635		
Secondary dressing			220007					
lightweight bandage (PC only)	KNIT BAND x 4m	5 x 4	7 x 4	10 x 4	15 x 4			
		1						
Drug tariff Secondary absorbent non		£0.11	£0.16	£0.18	£0.32			
woven wadding	K SOFT							
Drug tariff								
PECOS code								
Secondary dressing	PREMIERPORE							
adhesive		5 x 7	10 x 10	10 x 15	10 x 20	10 x 25	10 x 30	10 x 35

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Drug tariff		£0.05	£0.12	£0.18	£0.32	£0.36	£0.45	£0.52
Secondary dressing non		7.5 x						
adhesive	TELFA	7.5	10 x 7.5	15 x 7.5	20 x 7.5			
Drug tariff		£0.12	£0.16	£0.18	£0.29			
		7.5 x				12.5 x		
Silicone foam (S)	BIATAIN SILICONE	7.5	10 x 10	10 x 20	10 x 30	12.5	15 x 15	18 x 18
Drug Tariff		£1.55	£2.28	£2.88	£5.54	£2.79	£4.15	£5.81
PECOS code			193448	D		73467		59171
	KLINIDERM SILICONE							
Silicone wound contact	WOUND CONTACT LAYER	5 x 7.5	7.5 x 10	12 x 15	17 x 25	20 x 30		
Drug Tariff		£1.10	£1.63	£4.30	£9.00	£12.45		
PECOS code		267767	267774	267781		267798		
	KLINIDERM SUPER							
Superabsorber	ABSORBENT	10 x 10	10 x 15	10 x 20	20 x 20	20 x 30	20 x 40	
Drug Tariff		£0.49	£0.69	N/A	£0.99	£1.49	£1.99	
PECOS code					217144			
	CUTIMED SORBION SACHET		7.5 x					
Superabsorber (S)	EXTRA	5 x 5	7.5	10 x 10	20 x 10	20 x 20	30 x 20	
Drug Tariff		£1.55	£1.90	£2.40	£3.98	£7.46	£10.65	
PECOS code				264889	264896	264902	264919	
		12.5 x		12.5 x				
Superabsorber (PC only)	VLIWASORB PRO	12.5	22 x 22	22.5	22 x 32			
Drug Tariff		£0.93	£1.97	£1.10	£2.48			
Superabsorber	ZETUVIT E NON-STERILE	10 x 10	10 x 20	20 x 20	20 x 40			
Drug Tariff		£0.07	£0.09	£0.15	£0.29			
				413864				
PECOS code		413860	413861	1	413866			
			Wide					
Waterproof dressing		Short	short	Foot /				
protection	SEAL TIGHT	leg	leg	ankle				
Drug Tariff		£10.63	£10.63	£10.63				

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Product Data Sheets

Clinipore (Clinisupplies)	
Surgical adhesive tape	
Description: Permeable non-woven synt	hetic adhesive tape
Sizes:	
1.25cm x 5m	
2.5cm x 5m	
5cm x 5m	
Indications for use	Retention of dressings and bandages
	Fixing ostomy appliances
	Fastening lightweight tubing
	When repeat usage is required
Contraindications/	Any known sensitivity to adhesive
cautions	Do not apply to broken skin
How to apply/remove	Apply: Direct to required area, avoid large adhesive
	margins to ensure patient comfort and minimise risk of
	discomfort on removal
	Removal: Loosen corner of tape and peel back
	For very fragile skin adhesive remover may be required,
	which is applied over surface of adhesive border (refer to
	Therapeutic Stoma Prescribing Guidance for preferred choice)
Secondary Dressing	NA
Frequency of dressing changes and	As wound dressing regimen dictates
removal	
Prescribing guidance	Consider the use of basic adhesive dressings when
	appropriate to negate need for tape

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Surgical adhesive tape Description: Permeable non-woven synthetic adhesive tape Sizes: 5cm 10cm 15cm 20 cm Indications for use • Retention of dressings and bandages • Fixing ostomy appliances

Primafix (Smith & Nephew)

	 Retention of dressings and bandages Fixing ostomy appliances Fastening lightweight tubing When repeat usage is required
Contraindications/ cautions	Any known sensitivity to adhesive Do not apply to broken skin
How to apply/remove	 Apply: direct to required area, avoid large adhesive margins to ensure patient comfort and minimise risk of discomfort on removal Removal: loosen corner of tape and peel back For very fragile skin adhesive remover may be required, which is applied over surface of adhesive border (refer to Therapeutic Stoma Prescribing Guidance for preferred choice)
Secondary Dressing	NA
Frequency of dressing changes and removal	As wound dressing regimen dictates
Prescribing guidance	Consider the use of basic adhesive dressings when appropriate to negate need for tape

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ActivHeal Aquafiber Ag (Advanced Medical Solutions)

Antimicrobial (Silver)

Description: A sterile, non-woven pad consisting of a high M (mannuronic acid) calcium alginate and carboxymethylcellulose (CMC). Silver ions are released in the presence of wound fluid. As fluid is absorbed, the alginate forms a soft gel. Effective for up to 7 days.

Sizes Available	
5cm x 5cm	
10cm x 10cm	
15cm x 15cm	
2.7cm x 32cm (ribbon)	

Indications for use	Moderate to heavily exuding wounds including pressure ulcers, venous leg ulcers, diabetic ulcers, cavity wounds, post-op surgical wounds, superficial and partial thickness burns, traumatic wounds (dermal lesions, trauma injuries or incisions.
Contraindications/cautions	Not suitable for dry or lightly exuding wounds.
	Do not use on individuals with a known sensitivity to alginates or
	silver.
	Not suitable to control heavy bleeding.
	Not suitable for full thickness burns
How to apply/remove	Application: Sheet – Apply directly to wound bed to ensure maximum contact. Lay loosely into cavity wounds, filling no more than 80% to allow for product swelling
	Ribbon: Loosely pack into cavity to approximately 80% depth to allow for product swelling. Ensure a 'tail-end' is left exposed of each ribbon for ease of removal.
	Both sheet and ribbon will require to be secured with a secondary dressing.
	Removal : If adhering to wound bed, moisten for ease of removal.
Frequency of dressing changes	As exudate level and slough dictate. Wetter wounds will require more frequent dressing changes.
Prescribing guidance	When packing a wound, ensure the number of products used is clearly documented in the dressing regime section of the wound chart. The number of products removed and inserted should be documented at every dressing change to reduce the risk of retained products in a wound.

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Aquacel Ag+ Extra and Aquacel Ag+ Ribbon (Convatec)

Antimicrobial Dressings / Hydrocolloid dressings

Description: Primary hydrofibre wound contact layer composed of hydrocolloid fibre (sodium carboxymethylcellulose) impregnated with silver and surfactant. High absorbency. Converts to gel on contact with moisture (i.e. wound exudate).

Sizes Aquacel Ag+ I 5 x 5cm	4 x 10cm	
10 x 10cm	4 x 20cm	
15 x 15cm	4 x 30cm	
Sizes Aquacel Ag+ I		
1 x 45cm	2 x 45cm	
Indications for use Contraindications	infection is susper • Moderate to heavily • Debridement of mo • Do not use on part dressing compon • Do not use where	ist slough tients with a known sensitivity to silver or other ents e the presence of metals is contraindicated e.g.
	patients receiving	g radiotherapy or having MRI
	Do not use on pre	egnant or breast feeding women
How to apply/remove	 the wound and apply it g Apply to wound bed ledge Ensure maximum cod Lay loosely into cavit product swelling Overlap surrounding Ribbon: Loosely pack into cavit product swelling Ribbon can be cut ledusing on narrow cavit Removal: Lift carefully full Irrigate to facilitate moist bed. Re-assessment of wound to continue should be apply to wound to wound to continue should be apply to wound to	leaving small overhang around the entire wound ntact with wound bed by wounds filling no more than 80% to allow for peri-wound skin wity to approximately 80% of depth to allow for ngthwise. Use 4 x 20cm sheet and cut to size if ty rom wound bed using area of overhang. sure and ease of removal if adherence to wound and to determine if silver containing dressing undertaken at least two weekly
Frequency of dressing		p to 7 days. As exudate and slough dictates –
changes		ridement management guidance (appendices 1
	& 2)	
Prescribing guidance	management of uncomConsideration should alsMechanically lifts sloug	o be given to the following when prescribing: Ih and bacteria from wound bed
	 Do not use with enzym 	atic debriders (eg Flaminal Forte)

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 Reduces risk of maceration and excoriation of peri-wound and surrounding tissues Avoid in dry or low exuding wounds as it can dry out and adhere to wound bed
 In deep cavities requiring multiple dressings consider alternative Can be used as secondary dressing with honey in tracking wounds

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Cutimed Sorbact Swab and Ribbon (BSN)

Other Antimicrobials

Description: Low-adherence dressing made from fabric coated with dialkylcarbamoyl chloride, a hydrophobic substance is designed to bind microorganisms in the presence of moisture.

	1
Sizes	
Swabs 4x6cm	
Swabs 7x9cm	
Ribbon 2x50cm	
Ribbon 5x200cm	
Round swabs 3cm	
Indications for use	Chronic and acute wounds that are critically colonised
	Where an antimicrobial dressing is indicated in moderately to highly exuding wounds
Contraindications	Do not use in combination with ointments and creams as the binding effect is impaired
How to apply/remove	Place directly onto the wound surface
	Swabs can be used folded or unfolded and applied to achieve
	maximum contact with the wound bed
Frequency of dressing changes	As exudate dictates – refer to exudate management guidance, can be left in place for up to 7 days
	Re-assessment of wound to determine if antimicrobial dressing to continue should be undertaken at least two weekly.
Prescribing guidance	 Requires a moist wound condition to be effective
	Ribbon must not be cut due to shedding

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Inadine (Systagenix)

Antimicrobial dressings, lodine

Antimicrobial dressings, lodine		
Sizes 5 x 5 cm 9.5 x 9.5 cm		
Indications for use	 Reduce bacterial burden in superficial low exuding wounds with signs of local infection Can be used under compression 	
Contraindications	 Renal/thyroid impairment Lithium therapy NB for full list of cautions/contraindications refer to product literature and BNF 	
How to apply/remove	 Apply: Iodine based products can stain or irritate surrounding skin therefore ensure products do not have large border out with wound bed. "Bumpy" side should be in contact with wound bed Products can be cut to size Centre the dressing on the wound and apply directly onto wound bed Removal: Lift corner of dressing and peel back from wound Irrigate with sterile saline to facilitate moisture and ease of removal if adherence to wound bed 	
Frequency of dressing changes	 1-7 days depending upon exudate levels Pale colour of rayon indicates uptake of iodine Re-assessment of wound to determine if antimicrobial dressing to continue should be undertaken at least two weekly. 	
Prescribing guidance	 Volume of products prescribed should reflect short term use e.g. two week supply in first instance and review treatment plan. Antimicrobials should only be used on the small number of patients who need them and educate those who don't. Recent Health Technology Assessment Report 13 (Dec 2015) on the use of antimicrobial wound dressings for chronic wounds highlighted the lack of evidence for their routine use. Dressings have little absorbency capacity 	

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Iodoflex Paste (Smith and Nephew)

Antimicrobials, lodine

Description: A paste basis containing iodine 0.9% as cadexomer-iodine with a gauze backing that releases free iodine on exposure to wound exudate.

Sizes
5g
10g
17g

Indications for use	 Treatment of wound infection and debridement of moist, superficial slough in chronic wounds Maximum single application of 50g; Maximum weekly application of 150g; Maximum duration up to 3 months in any single course of treatment 		
Contraindications	Should not be used on:		
	Dry, necrotic tissue		
	 Known sensitivity to any of its ingredients 		
	Children		
	 Pregnant or lactating women 		
	 People with thyroid disorders or renal impairment 		
	 Patients prescribed lithium 		
	If bone or tendon visible		
How to apply/remove	1. Peel back gauze backing		
	2. Remove suitable amount and mould to wound surface area,		
	ensuring in full contact with wound bed		
	Removal:		
Frequency of dressing	Removal:• by irrigation with saline or water		
Frequency of dressing changes	Removal:• by irrigation with saline or waterRegularly monitor for reduction in exudate to ensure wound bed does not		
Frequency of dressing changes	Removal:• by irrigation with saline or water		
	Removal: • by irrigation with saline or water Regularly monitor for reduction in exudate to ensure wound bed does not dry out. Refer to exudate and debridement management guidance		
changes	Removal: • by irrigation with saline or water Regularly monitor for reduction in exudate to ensure wound bed does not dry out. Refer to exudate and debridement management guidance (appendices 1&2)		
	Removal: • by irrigation with saline or water Regularly monitor for reduction in exudate to ensure wound bed does not dry out. Refer to exudate and debridement management guidance (appendices 1&2) Re-assessment of wound to determine if antimicrobial dressing to		
changes	Removal: • by irrigation with saline or water Regularly monitor for reduction in exudate to ensure wound bed does not dry out. Refer to exudate and debridement management guidance (appendices 1&2) Re-assessment of wound to determine if antimicrobial dressing to continue should be undertaken at least two weekly.		
changes	 Removal: by irrigation with saline or water Regularly monitor for reduction in exudate to ensure wound bed does not dry out. Refer to exudate and debridement management guidance (appendices 1&2) Re-assessment of wound to determine if antimicrobial dressing to continue should be undertaken at least two weekly. Iodine may be absorbed, particularly from large wounds or during 		
changes	Removal: • by irrigation with saline or water Regularly monitor for reduction in exudate to ensure wound bed does not dry out. Refer to exudate and debridement management guidance (appendices 1&2) Re-assessment of wound to determine if antimicrobial dressing to continue should be undertaken at least two weekly. • lodine may be absorbed, particularly from large wounds or during prolonged use		
changes	 Removal: by irrigation with saline or water Regularly monitor for reduction in exudate to ensure wound bed does not dry out. Refer to exudate and debridement management guidance (appendices 1&2) Re-assessment of wound to determine if antimicrobial dressing to continue should be undertaken at least two weekly. Iodine may be absorbed, particularly from large wounds or during prolonged use Suitable for smaller wound surface areas. 		
changes	 Removal: by irrigation with saline or water Regularly monitor for reduction in exudate to ensure wound bed does not dry out. Refer to exudate and debridement management guidance (appendices 1&2) Re-assessment of wound to determine if antimicrobial dressing to continue should be undertaken at least two weekly. Iodine may be absorbed, particularly from large wounds or during prolonged use Suitable for smaller wound surface areas. Not suitable for large surface areas. 		
changes	 Removal: by irrigation with saline or water Regularly monitor for reduction in exudate to ensure wound bed does not dry out. Refer to exudate and debridement management guidance (appendices 1&2) Re-assessment of wound to determine if antimicrobial dressing to continue should be undertaken at least two weekly. lodine may be absorbed, particularly from large wounds or during prolonged use Suitable for smaller wound surface areas. Not suitable for large surface areas. Some patients may find pain on application; if pain in wound continues/cannot be tolerated discontinue use and irrigate 		
changes	 Removal: by irrigation with saline or water Regularly monitor for reduction in exudate to ensure wound bed does not dry out. Refer to exudate and debridement management guidance (appendices 1&2) Re-assessment of wound to determine if antimicrobial dressing to continue should be undertaken at least two weekly. lodine may be absorbed, particularly from large wounds or during prolonged use Suitable for smaller wound surface areas. Not suitable for large surface areas. Some patients may find pain on application; if pain in wound 		

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Iodosorb Ointment (Smith and Nephew)

Antimicrobials, lodine

Description: ointment containing 0.9% iodine as cadexomer-iodine. Free iodine is released from ointment on exposure to wound exudate.

Size	
10g	

Indications for use	Treatment of wound infection and debridement of moist, superficial slough in chronic wounds
Contraindications	 Should not be used for: Dry, necrotic tissue Known sensitivity to any of its ingredients Children Pregnant or lactating women People with thyroid disorders or renal impairment Patients taking lithium If bone or tendon exposed
How to apply/remove	 Ensure in full contact with wound surface area Removal: Irrigation with saline or water
Frequency of dressing changes	Re-assessment of wound to determine if antimicrobial dressing to continue should be undertaken at least two weekly.
Prescribing Guidance	 lodine may be absorbed, particularly from large wounds or during prolonged use Less likely to dry wound bed out when slough removed and bacterial burden reduced due to ointment preparation Not suitable for large surface areas Some patients may find pain on application; if pain in wound continues/cannot be tolerated discontinue use and irrigate Seek specialist advice in diabetic foot conditions and arterial insufficiency Maximum single application of 50g Maximum weekly application of 150g Maximum duration up to 3 months in any single course of treatment

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Prontosan Gel X (B Braun)

Other antimicrobials

Description: A hydrogel wound gel containing betaine surfactant (disrupts biofilm) and polihexanide (an antiseptic).

Size	
30ml	

Indications for use Contraindications	 Biofilm disruption, cleansing, decontamination and moisturising of: Acute wounds Chronic wounds First and second degree burns If known sensitivity to any of the gel's ingredients. NB In very rare cases
	there may be a mild burning sensation after application of Prontosan wound gel but this should disappear after a few minutes.
How to apply/remove	Apply directly to wound bed
Frequency of dressing	N/A
Changes	
Prescribing Guidance	Has a shelf life of 28 days after opening - no refrigeration required

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Urgotul AG/Silver

Antimicrobial Dressings Description: Urgotul AG/silver is a polyester mesh impregnated with hydrocolloid particles, Vaseline, cohesion polymers and salts. Non -adhesive and non-occlusive dressing

Size 10cm x 12cm

	-	
Indications for use	Provides a contact layer directly onto the wound surface.	
	 Infected or at risk of infection wound 	
	 Non to low exuding wounds 	
	Acute wounds	
	Chronic wounds	
Contraindications/cautions	 Do not use if know sensitivities to silver or other ingredients of the dressing 	
	Do not use on patients undergoing MRI	
	Do not use in a hyperbaric chamber	
	 Do no use on pregnant or breast feeding women, new born or premature babies 	
	Avoid contact with electrodes or conductive gels during electronic measurement , e.g. EEG and ECG	
How to apply/remove	Apply: Moisten gloves with saline in order to hold the dressing easier	
	Removal : Raise corner and peel back off wound. Should lift off wound with no adherence.	
Frequency of dressing changes	Dependent on the nature of the wound, can be left in place for up to7 days; however may require more frequent changes if there is a risk of desiccation or unexpected increase in exudate with need to review dressing regimen or more frequent changes.	
	Maximum use 1 month	
	Refer to exudate and debridement management guidance (appendix 1 & 2)	
Prescribing guidance	Usually used for low exuding wounds	
	1	

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Jelonet (Smith & Nephew)

Specialist burns dressing Low adherence dressings Description: Knitted polyester primary dressing impregnated with neutral triglycerides, conforms to wound bed.

Size 10 x 10cm

Indications for use	 Provides a contact layer directly onto the wound surface. Basic wound dressing for non-complex wounds: Minor burns Abrasions Superficial wounds As a contact layer under compression bandage on leg ulcers A cost effective alternative to silicone contact layer products when dressings are changed more than once a week.
Contraindications/cautions	 Can be used under compression; however risk of adherence to wound bed if minimal exudate present. Use with caution on chronic low exuding wounds with viscous exudate which may result in pooling and restricted drainage through dressing pores.
How to apply/remove	 N-A Ultra Do not use if allergic to silicone. Apply: Place flat onto the wound surface with 2.5cm border May be applied in multiple layers "fluffed" up to reduce risk of adherence and frequency of dressing changes. Can be cut or folded to size. Removal: Raise corner and peel back off wound. Should lift off wound with no adherence.
Frequency of dressing changes	 Dependent on the nature of the wound, can be left in place for up to 7 days; however may require more frequent changes if there is a risk of desiccation or unexpected increase in exudate with need to review dressing regimen and/or more frequent changes. If secondary dressing allows strike through e.g. bandages or dry dressings there is a risk of bacterial ingress with requirement for review of dressing regimen or more frequent changes.
	Refer to exudate and debridement management guidance (appendix 1 & 2)
Prescribing guidance	Usually used for low exuding wounds Atrauman is a cost effective alternative to silicone contact layer dressings, if greater than once a week dressing changes are required.

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Clinipods (Mayers Pharma)

Sterile Saline Solution

Description: Sterile normal saline solution 0.9% w/v ph Eur

Size 20ml vials in packs of 25

Indications for use	 For topical irrigation of wounds to remove loose slough, 	
	debris and chronic wound fluid from wound bed	
	Social cleansing of peri wound margins	
Contraindications/cautions	Do not mix with other fluids for irrigation unless directed	
	Do not use for injection	
How to apply/remove	Remove twist off seal then use vial to direct the flow of saline	
	over the wound.	
	Use gauze to soak up irrigation waste.	
	Product should be warmed to body temperature before use	
Frequency of dressing	At each wound intervention only if wound requires cleansing as	
changes	not all wounds require cleansing at dressing change.	
Prescribing guidance	Clean granulating wounds do not require routine cleansing	
	Patient may irrigate wound in shower, which can negate need	
	for saline irrigation	
	Refer to NHSGGC wound cleansing guidance	

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Prontosan solution (B Braun)

Other antimicrobials

Description: An aqueous wound irrigation solution containing betaine surfactant (disrupts biofilm) and polihexanide (an antiseptic).

Size
350ml
40ml

Indications for use	 Biofilm disruption, cleansing, decontamination and moisturising of: Acute wounds Chronic wounds First and second degree burns
Contraindications	If known sensitivity to any of the solutions ingredients
How to apply/remove	apply as a soak for at least 10 minutes
Frequency of dressing changes	N/A
Prescribing guidance	 Use only if indicated by wound cleansing guidance (See links) and debridement guidance (Appendix 2) Wound cleansing product for use in wounds showing signs of critical colonisation or for removal of biofilm Has a shelf life of 28 days after opening - no refrigeration required One bottle should allow for approximately 8 dressing changes (based on average size 10 x 10 cm wound size) Apply as a soak at every dressing change as per wound cleansing guidance (See links)

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Stericlens Aerosol (CD Medical)

Irrigation solutions (NaCl 0.9%) Description: Sterile sodium- chloride solution in a spray

Sizes
100mls
240 mls

	-
Indications for use	 For topical irrigation of wounds to remove loose slough, debris and chronic wound fluid from wound bed
	Social cleansing of peri wound margins
Contraindications/cautions	Do not mix with other fluids for irrigation unless directed
	Do not use for injection
	 Aerosol is pressurised container and should not be exposed to high temperatures, punctured or burnt. Local disposal regulations and requirements apply.
How to apply/remove	Apply : Direct nozzle to area requiring irrigation and spray approx 10cm from wound surface to reduce risk of spray back and allow maximum coverage of wound bed.
	Can be used through 360 degrees or upside down for awkward to irrigate areas
	Replace cap after use and store in clean area
	 Refer to manufacturer's instructions for further details if required
Frequency of dressing changes	At each wound intervention.
Prescribing guidance	 Aerosol design allows all saline to be used with no waste. Consider number of interventions and volume required at each dressing change to reduce waste
	 Clean granulating wounds do not require routine cleansing Patient may irrigate wound in shower, which can negate need for saline irrigation Refer to NHSGGC wound cleansing guidance

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ActiFormCool (Activa)

Hydrogel dressings Description: Ionic non adherent hydrogel sheet to debride devitalised tissue

Sizes 5 x 6.5cm 10 x 10cm 10 x 15cm 20 x 20cm	
10 x 10cm 10 x 15cm	
10 x 15cm	
20 x 20cm	
Indications for use	Dry eschar or slough
	Painful wounds
	Burns
	Radiation burns
	 Fungating wounds
	 Under compression for light to moderate exuding wounds
Contraindications	Deep cavity wounds
	Narrow cavity wounds
	Sinus wounds
	Bleeding wounds
	Infected wounds
	Poorly perfused wounds
How to	Position on wound bed and smooth into place
apply/remove	Removal: Lift one corner and gently peel off dressing
	If dressing has dried out, soak with water or saline to rehydrate and peel
	off.
Frequency of	As exudate and slough dictates – refer to exudate and debridement
dressing changes	management guidance (appendix 1 & 2)
	Dressing should be changed when dressing becomes discoloured or
	opaque.
Prescribing	Can stay in place for 7 days depending on levels of exudate
guidance	 Monitor peri-wound skin for maceration
	Consider cutting dressing to size of wound if maceration evident

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HYDROCLEAN ADVANCE (Hartmanns)

Debridement

Description: Hydroclean advance is a hydroactive wound dressing that contains, as a core component, a superabsorbent, polyacrylate (SAP) embedded in cellulose fibres and activated with Ringer's solution. The wound contact layer consists of a polypropylene knitted fabric to which silicone strips have been applied. Hydroclean advance delivers Ringer's solution to the wound for up to three days. During this time, interactive and continuous and continuous wound irrigation takes place and wound exudate is absorbed. The SAP inactivates metalloproteinases which impair wound healing, as a result, stagnating healing in chronic wounds can be reactivated.

Sizes	
3cm round	10cm x 10cm
4cm round	4cm round cavity
5.5cm round	7.5cm x 7.5cm cavity
7.5cm x 7.5 cm	

Indications for use	Suitable for wet/moist treatment of acute and chronic wounds, in particular wounds requiring debridement or with impaired healing tendency. It can be used on infected wounds and stagnating wounds.
Contraindications/cautions	Do not use on patients with an intolerance to any of the components in the dressing. Consult wound specialist before using on infected wounds. In patients with risk of excessive bleeding remove with caution or irrigate wound bed prior to removal with appropriate wound cleansing solution.
How to apply/remove	Apply: DO NOT CUT. Select dressing size to fit wound without overlap as hyper-hydration may occur (swelling of cells that will resolve, this is not maceration). Apply white side down so that the blue writing is visible on the outside. Secondary dressing should be a simple film and not a foam or silicone dressing.
	Removal : Hydroclean advance should be able to be removed easily, however if there is adherence then the dressing can be removed by irrigating with appropriate wound cleansing solution
Frequency of dressing changes	Three days
Prescribing guidance	Hydroclean advance debrides rapidly and often debridement is achieved within a few days. Do not prescribe more than one week supply of dressings.

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UrgoClean Pad (Urgo)

Hydrocolloid dressings Description: Hydrocolloid fibre that converts to gel on contact with moisture (i.e. wound exudate). Pad has soft-adherent lipocolloidal contact layer.

Sizes: pad	
6 x 6cm	
10 x 10cm	
20 x 15cm	

Indications for use	 Moderate to heavily exuding wounds 	
	Debridement of moist slough	
Contraindications	Any known sensitivities	
How to apply/remove	Pad:	
	Select a dressing larger than the wound area. Centre the dressing on	
	the wound and apply it gently to wound site.	
	1. Apply to wound bed leaving small overhang around the entire	
	wound edge	
	2. Ensure maximum contact with wound bed	
	3. Lay loosely into cavity wounds filling no more than 80% to allow for	
	product swelling	
	4. Overlap surrounding peri wound skin	
	Removal: Lift carefully from wound bed using area of overhang	
	Irrigate to facilitate moisture and ease of removal if adherence to wound bed	
Frequency of	As exudate and slough dictates – refer to exudate and debridement	
dressing changes	management guidance (appendices 1 & 2)	
Prescribing	Mechanically lifts slough and bacteria from wound bed	
guidance	 Reduces risk of maceration and excoriation of peri-wound and 	
	surrounding tissues	
	-	
	 Avoid in dry or low exuding wounds as it can dry out and adhere to wound bed 	
	 In deep cavities requiring multiple dressings consider alternative 	
	1	

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Flaminal Forte (Flen Hea	althcare)
	ntimicrobial Alginate Gel
	e alginate gel containing dual enzymes (glucose oxidase and
lactoperoxidase) to reduc	e bioburden and debride devitalised tissue
Size	
15g (coverage 40cm)	
15g (coverage 40cm)	
Indications for use	Moderate to heavily exuding, critically colonised or infected
	Sloughy critically colonised or infected wounds
	 Critically colonised or infected cavity wounds
Controindiactions	
Contraindications	Dry or low exuding wounds
	Clean wounds with no signs or risks of clinical infection
	Known sensitivities
How to apply/remove	1. Apply directly to wound bed ensuring protection of surrounding
	skin
	2. A syringe may be used to insert into cavity wounds
	Removal: By gentle irrigation with sterile water or saline
Frequency of dressing	1 - 4 days depending upon exudate levels. Requires changing when
Changes	gel structure disappears
	Re-assessment of wound to determine if antimicrobial dressing
	to continue should be undertaken at least two weekly.
Prescribing Guidance	No fibre shed in cavities
5	Should only be used for two week periods

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Flaminal Hydro (Flen Health)

Antimicrobial Alginate Gel

Description: Alginate gel containing two antimicrobial enzymes (glucose oxidase and lactoperoxidase) which exert their effect without damaging healthy skin cells. Debrides the wound and manages moisture balance. Contains lower proportion of alginate than Flaminal Forte so absorbs less exudate.

Sizes

15g Tube (coverage 40cm)

Indications for use	Low to modertately exuding wounds	
	Sloughy critically colonised or infected wounds	
	Critically colonised or infected cavity wounds	
Contraindications/cautions	Highly exuding wounds (use Flaminal Forte instead)	
	Clean wounds with no signs or risks of clinical infection	
	Known sensitivities	
How to apply/remove	Apply: Apply directly to wound bed ensuring protection of	
	surrounding skin. A syringe may be used to insert into cavity	
	wounds.	
	Removal : By gentle irrigation with sterile water or saline.	
Frequency of dressing	1-4 days depending on exudate levels. Requires changing	
changes	when gel structure disappears. Reassessment of wound to	
-	determine if antimicrobial dressing to continue should be	
	undertaken at least 2 weekly.	
Prescribing guidance	If exudate levels increase, consider switch to Flaminal Forte.	

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ActivHeal Hydrogel (Advanced Medical Solutions)

Hydrogel application Description: Gel (composed of guar gum and propylene glycol) containing 85% water. No animal derived ingredients.



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Indications for use	 Necrotic and sloughy wounds with nil to low exudate
Contraindications	 Surgical implantations Full thickness burns
How to apply/remove	Direct to wound bed, half fill cavity to reduce risk of maceration to
now to apply/remove	surrounding skin and number of dressing changes required.
Frequency of dressing changes	As exudate and slough dictates – refer to exudate and debridement management guidance (appendix 1 & 2)
Prescribing guidance	Seek specialist advice in diabetic foot conditions and arterial insufficiency

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ActivHeal Aquafiber Extra (Advanced Medical Solutions)

Gelling Fibre Dressing

Description: A soft, conformable, highly absorbent dressing that converts into a soft clear gel when in contact with wound exudate. It contains a re-inforcing layer between 2 layers of absorbent fibres to improve the integrity of the dressing when it is wet, so it can be removed intact.

Sizes
5cm x 5cm
10cm x 10cm
15cm x 15cm
2cm x 46cm (ribbon)

Indications for use	Moderate to heavily exuding wounds including pressure ulcers, venous leg ulcers, diabetic ulcers, cavity wounds, post-op surgical wounds, superficial and partial thickness burns. Can also be used to control minor bleeding in superficial wounds.
Contraindications/cautions	Not indicated to control heavy bleeding
How to apply/remove	Apply: Sheet – Apply directly to wound bed to ensure maximum contact. Lay loosely into cavity wounds, filling no more than 80% to allow for product swelling
	Ribbon: Loosely pack into cavity to approximately 80% depth to allow for product swelling. Ensure a 'tail-end' is left exposed of each ribbon for ease of removal.
	Removal : If adhering to wound bed, moisten for ease of removal.
Frequency of dressing changes	As exudate level and slough dictate. Wetter wounds will require more frequent dressing changes.
Prescribing guidance	When packing a wound, ensure the number of products used is clearly documented in the dressing regime section of the wound chart. The number of products removed and inserted should be documented at every dressing change to reduce the risk of retained products in a wound.

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Duoderm Extra Thin (Convatec)

Polyurethane matrix dressing without adhesive border Description: Semi-permeable conformable low absorbency hydrocolloid adherent occlusive dressing.

Sizes		
5 x 10cm	9 x 15cm	
7.5 x 7.5cm	9 x 25cm	
10 x 10cm	9 x 35cm	
15 x 15cm		
Indications for use	 Superficial low exuding wounds To debride low levels of slough by autolysis Primary dressing on clean granulating/epithelialising wound Secondary dressing over Aquacel Extra (Jubilee technique) on pos operative incisions in highly exuding wounds Secondary dressing to provide showerproof, bacterial barrier To protect peri-wound margins when using NPWT or Larvae therap 	
Contraindications/ cautions	 Known sensitivities to carboxymethylcellulose, gelatin, pectin Heavily exuding wounds when used direct to wound bed Known wound infection Should not be applied to exposed muscle or bone 	
How to apply/remov	Apply: dry surrounding peri wound skinPeel backing layer and place directly on wound bed, allowing a 3cmborder around wound bed.	
	Can be cut to size	
	Removal: Press down gently on skin and lift corner of dressing stretching each edge until free.	
Frequency of dressing changes	 Semi transparent qualities and will allow for viewing of wound bed. Change when gelling of 80% of dressing has taken place or if wour assessment is required. If Jubilee technique is used, change when underlying Aquacel Extra has fully gelled or if wound assessment is required. 	
Prescribing guidance	Frequency of dressing change when prescribing volume	

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Debrisoft (Activa)

Physical debridement pads

Description: Debrisoft is a polyacrylate coated pad made up of polyester fibres with bound edges.

NB: this is a debridement pad and **NOT** a wound dressing

Sizes	
10cm x 10cm	
Indications for use	 To debride loose superficial slough and debris to reveal underlying granulating wound bed Removal of softened loose hyperkeratotic skin from peri wound margins
Contraindications	 Wound bed with granulating base Dry slough or necrosis Deep slough Pain despite analgesia
How to use	 Fully moisten pad with water before use and shake off excess – do not squeeze out Apply rotational movements over wound bed and margins with pad, with fibre side contacting the wound bed to loosen and remove slough and debris. Procedure may take a few minutes, as tolerated, to debride and expose granulating wound bed. During procedure if less hydrated slough is exposed, further hydration with wound dressings is required to soften and liquefy slough to be removed at following dressing change with Debrisoft. Check pad at end of intervention – if pad is clean this may be due to technique in using pad (seek further advice on correct use)
Frequency/ Prescribing guidance	 May only require a "one off" treatment or follow up depending on chronicity of wound At follow up dressing change if slough which was removed is apparent again, this may indicate poor perfusion with vascular referral required; or biofilm formation requiring cleansing with surfactant (For further information on range of debridement techniques refer to appendix 2)

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UCS Debridement (medi UK)

Physical debridement clothDescription: Sterile, pre-moistened soft debridement cloth for single use.

NB: this is a debridement cloth and $\ensuremath{\textbf{NOT}}$ a wound dressing

Size	
10 x ⁻	10cm

Indications for use	 To debride loose superficial slough and debris to reveal underlying granulating wound bed Removal of softened loose hyperkeratotic skin from peri wound margins
Contraindications	 Wound bed with granulating base Dry slough or necrosis Deep slough Pain despite analgesia
How to use	 Apply rotational movements over wound bed and margins with cloth to loosen and remove slough and debris. Procedure may take a few minutes, as tolerated, to debride and expose granulating wound bed. During procedure if less hydrated slough is exposed, further hydration with wound dressings is required to soften and liquefy slough to be removed at following dressing change with UCS cloth.
Frequency/ Prescribing guidance	 May only require a "one off" treatment or follow up depending on chronicity of wound At follow up dressing change if slough which was removed is apparent again, this may indicate poor perfusion with vascular referral required; or biofilm formation requiring cleansing with surfactant (For further information on range of debridement techniques refer to appendix 2)

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Dress-It (Richardson) Nurse-It (Medicareplus International)

Dressing packs

Description: Procedure dressing pack for use in primary care, to provide a sterile working surface with contents to support aseptic technique when carrying out wound management.

Sizes of gloves: Small/medium and medium/large gloves available NB variation in contents of packs

Dress-It	Nurse-It	
 Vitrex gloves x one pail Softswabs 4 ply x 4 Absorbent pad x 1 Sterile field x 1 Paper towel x 1 Large apron x 1 Disposable bag 	 Latex Free, Powder Free, Nitrile Gloves x one pair Non-Woven Swabs x 7 Laminated Paper Sterile Fields x 2 Paper Towel x 1 Large Apron x 1 White Polythene Disposable Bag x 1 Compartment Tray x 1 Disposable Forceps x 1 Laminated Paper Sterile Field x1 Paper Measuring Tape x 1 	
dications for se Dressing pack for patients to support aseptic wound management in domiciliary setting.		
Contraindications/ • Nor cautions	None noted	
guidance • Pac	leration should be given to the following when prescribing: ize 10; take this into account when prescribing to avoid waste dent on number of dressing changes	

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Covawound Alginate (Covalon Technologies Ltd)

Alginate

Description: A primary wound dressing made from the calcium salt of alginic acid. When in contact with wound exudate, the calcium ions in the dressing are exchanged for sodium ions to produce a highly absorbent, soft gelling dressing which maintains its integrity. The soft gel facilitates moist wound healing and de-sloughing, settling into the contours of the wound to minimise the dead spaces where exudate can pool and bacteria grow.

Sizes		
5cm x 5cm		
10cm x 10cm		
2cm x 30cm (rope)		
Indications for use	 Moderate to heavily exuding wounds such as partial thickness burns, donor sites, leg ulcers, pressure ulcers, diabetic foot ulcers, post-op incision and trauma wounds, The rope dressing can be applied to moderate to heavily exuding cavity wounds. This dressing has haemostatic properties, therefore can be used on bleeding wounds. 	
	This product can be used under compression	
Contraindications/cautions	Sensitivities to calcium alginate	
	Dry or lightly exuding wounds.	
How to apply/remove	Apply: Select a dressing that is slightly larger than the wound. Alginate rope dressing should be used in cavity wounds. Product may be trimmed to fit the wound size. Apply directly to wound bed. For cavity wounds, loosely pack to 80% depth, ensuring an end of product is left exposed for removal. Cover and secure with a non-occlusive secondary dressing.	
	Removal : Hydrate with saline or sterile water if dry to ensure ease of removal.	
Frequency of dressing changes	As exudate levels dictate.	
Prescribing guidance	If using to pack wound ensure number of products inserted and removed are clearly documented on a wound chart to avoid the risk of retained products.	

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Actilite (Advancis Medical)

Antimicrobial Dressings, Honey sheet dressing

Description: Medical grade manuka honey 99% and manuka oil 1% knitted viscose sheet dressing 99% manuka honey and 1% manuka

Sizes	
5 x 5cm	20 x 30cm
10 x 10cm	30 x 30cm
10 x 20cm	30 x 60cm
Indications for use	 Reduce bacterial burden in superficial low exuding wounds with signs of local infection Actilite may be used for patients whom iodine based products are contraindicated or alternative honey and iodine based products are no tolerated by patient Can be used under compression
Contraindications/ cautions	 Any known sensitivities to bee venom NB for full list of cautions/contraindications refer to product literature and BNF
How to apply/remove	Apply: "Bumpy" side should be in contact with wound bed Products can be cut to size Centre the dressing on the wound and apply directly onto wound bed. Removal: lift corner of dressing and peel back from wound
Frequency of dressing changes	Can be left in place up to one week; however if antimicrobial dressing is required more frequent assessment may be required
Prescribing guidance	 Volume of products prescribed should reflect short term use e.g. two week supply in first instance and review treatment plan. Antimicrobials should only be used on the small number of patients who need them and educate those who don't. Health Technology Assessment Report 13 (Dec 2015) on the use of antimicrobial wound dressings for chronic wounds highlighted the lack of evidence for their routine use

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Activon Tube (Advancis)		
Antimicrobial Dressings, Honey-based topical application		
	al grade manuka honey ointment.	
Sizes 25g tube		
Indications for use	Debridement	
	Helps control odour	
	 Provides a moist wound healing environment for all types of acute and chronic wounds including; 	
	 pressure ulcers 	
	o burns	
	o graft sites	
	 fungating tumours 	
	Has antimicrobial properties suitable for use on infected wounds or	
	where bacterial resistance is suspected	
Contraindications	 Can be used in cavities <u>DO NOT</u> use if the patient has a known allergy to bee venom 	
Contraindications	 <u>DO NOT</u> use if the patient has a known allergy to bee venom <u>Not recommended on leg ulcers (SIGN 120)</u> 	
How to apply/remove	Apply directly to wound bed or insert into cavity.	
	Refer to wound cleansing guidelines (see links)	
Frequency of dressing changes	<i>Re-assessment of wound to determine if antimicrobial dressing to continue should be undertaken at least two weekly.</i>	
Prescribing guidance	 Can make wound bed very moist and may lead to maceration if 	
r reserising guidance	exudate not managed adequately	
	• A short lived stinging sensation may be experienced when applying	
	the honey, if pain in wound continues / cannot be tolerated	
	discontinue use and irrigate with saline solution	
	 Activon contains a high level of glucose, although no incidents of increased blood sugar levels due to use of honey in wounds has 	
	been reported, it is advisable to monitor blood sugar level in patients with diabetes	
	 Seek specialist advice in diabetic foot conditions and arterial insufficiency 	
	• Tube can be used for up to 90 days after opening (single patient use only)	

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Activon Tulle (Advanc	is)		
Antimicrobial Dressin	gs, Honey sheet dressing		
	scose impregnated with medical grade honey.		
Sizes			
10 x 10cm			
Indications for use	Debridement		
	Helps control odour		
	• Provides a moist wound healing environment for all types of acute and		
	 chronic wounds including; pressure ulcers 		
	o burns		
	 o graft sites 		
	 fungating tumours Has antimicrobial properties suitable for use on infected wounds or 		
	 Has antimicrobial properties suitable for use on infected wounds or where bacterial resistance is suspected 		
Contraindications	• <u>DO NOT</u> use if the patient has a known allergy to bee venom.		
	Not recommended on leg ulcers (SIGN 120)		
How to apply/remove	Apply directly to wound bed (can be opened out to cover larger surface		
	area).		
Frequency of	Can be cut to size if necessary. As exudate dictates refer to exudate and debridement management		
dressing changes	guidance (appendix 1&2)		
	Re-assessment of wound to determine if antimicrobial dressing to		
	continue should be undertaken at least two weekly.		
Prescribing	Can make wound bed very moist and may lead to maceration if		
guidance	exudate not managed adequately		
	• A short lived stinging sensation may be experienced when applying the		
	honey, if pain in wound continues/cannot be tolerated discontinue		
	 use and irrigate with saline solution Dressing hardens when cold, can be softened in warm environment, 		
	needs to be softened prior to use		
	Activon contains a high level of glucose, although no incidents of		
	increased blood sugar levels due to use of honey in wounds has		
	been reported, it is advisable to monitor blood sugar level in patients with diabetes		
	Seek specialist advice in diabetic foot conditions and arterial		
	insufficiency		

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L-Mesitran Ointment (Aspen Medical Europe)

Antimicrobial ointment

Description: Hydroactive antibacterial ointment that has anti-inflammatory properties, debrides and reduces malodour, stimulates wound healing, is safe and cost effective.

Contains: 48% medical grade Honey, Medical Grade Hypoallergenic Lanolin, Sunflower oil, Cod liver oil, Calendula officinalis, Aloe Barbadensis, Vitamin C and E, Zinc Oxide.

Sizes	
20g tube	
50g tube	

Indications for use	 Superficial and acute wounds (cuts, abrasions, donor sites, etc). Superficial and partial-thickness burns (first and second degree). Chronic wounds (pressure ulcers, and venous, arterial and diabetic ulcers). Fungating wounds (to help deodorise and debride). Colonised acute wounds and post-operative surgical wounds. 	
Contraindications/cautions	None known to date. Do not use on patients who are sensitive to the product or any of its components.	
How to apply/remove	Apply: Remove lid and clean the mouth of the tube and the lid with an alcohol wipe. Apply a thin layer of L-Mesitran directly on to the wound area and surrounding tissue. Cover and secure with the appropriate secondary dressing, depending on the amount of the wound exudate.	
	Removal : Remove old L-Mesitran by gently cleansing the wound (refer to NHSGGC wound cleansing guidance).	
Frequency of dressing changes	Depends on the amount of exudate and the needs of the wound. Re-apply product once dissolved, usually every 24-48 hours.	
Prescribing guidance	Single patient use; after opening use within three months. Wound may appear to increase in size as devitalised tissue is removed through autolytic debridement. This is normal and prepares wound bed for healing.	

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L-Mesitran Soft (Aspen Medical Europe)

Antimicrobial gel

Description: Hydroactive antibacterial ointment that has anti-inflammatory properties, debrides and reduces malodour, stimulates wound healing, is safe and cost effective.

Contains: 40% medical grade Honey, Medical Grade Hypoallergenic Lanolin, propylene glycol, PEG 4000 and Vitamin C and E.

Sizes	
15g tube	

Indications for use	Superficial and acute wounds (cuts, abrasions, donor sites, etc). Superficial and partial-thickness burns (first and second degree). Chronic wounds (pressure ulcers, and venous, arterial and diabetic ulcers). Fungating wounds (to help deodorise and debride). Colonised acute wounds and post-operative surgical wounds.	
Contraindications/cautions	None known to date. Do not use on patients who are sensitive to the product or any of its components.	
How to apply/remove	Apply: Remove lid and clean the mouth of the tube and the lid with an alcohol wipe. Apply a thin layer of L-Mesitran Soft directly on to the wound area and surrounding tissue. Cover and secure with the appropriate secondary dressing, depending on the amount of the wound exudate.	
	Removal : Remove old L-Mesitran Soft by gently cleansing the wound.	
Frequency of dressing changes	Depends on the amount of exudate and the needs of the wound. Re-apply product once dissolved, usually every 24-48 hours.	
Prescribing guidance	Single patient use; after opening use within three months. Wound may appear to increase in size as devitalised tissue is removed through autolytic debridement. This is normal and prepares wound bed for healing.	

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Urgostart (Urgo)

THIS PRODUCT MUST ONLY BE INITIATED BY A WOUND SPECIALIST

Modulating Matrix

Description: Foam dressing with an adherent TLC-NOSF layer combine with an absorbent Polyurethane foam pad and an outer vapour permeable film.

Sizes (pad s	size in brackets)
6x6cm	
10 x 10cm	
15 x 20cm	

Indications for use	Exuding chronic wounds
Contraindications	Cancerous wounds
	 Fistular wounds with deep abscess formation.
	Known sensitivity to the dressing
	Signs of critical colonisation
How to apply/remove	Clean the wound using the standard protocol.
	• If an antiseptic has previously been used, rinse the wound carefully with normal saline.
	Carefully dry the skin around the wound.
	• UrgoStart can be cut using sterile scissors to fit the dressing size to the wound if necessary.
	• The soft-adherent layer adheres to surgical gloves (latex), it is
	recommended to use the protective tabs to aid application.
	• Apply the soft-adherent side of the dressing in contact with the wound.
	• Secure the dressing in place with a suitable bandage or tape.
	• To remove lift one corner and peel back gently.
Frequency of dressing changes	• The dressing should be changed every 2 to 4 days, and left in place for up to 7 days depending on the level of exudate and the clinical condition of the wound.
	• The recommended treatment duration is a minimum of 4 to 5 weeks.
Prescribing guidance	 In case of an atypical ulcer presenting induration or overgranulation, UrgoStart should only be used after checking the absence of wound-related malignancy in order not to delay the diagnosis.
	 The action of the product on the healing process may possibly cause stinging or painful sensations at the start of treatment with UrgoStart. This rarely warrants suspension of treatment
	• Urgostart is not recommended as a first line treatment in acute wounds, and in the treatment of Epidermolysis bullosa (even for longstanding lesions)
	Check that the sterility protector is intact before use.

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• Single-use, individual and sterile dressing: re-using a single-use product
may lead to risks of infection.

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Promogran Prisma 3M + KCI

THIS PRODUCT MUST ONLY BE INITIATED BY A WOUND SPECIALIST

Wound Balancing Matrix

Description: Promogran Prisma is a topical wound treatment containing collagen, oxidisedregenerated cellulose (ORC) and 1% silver-ORC. It is a protease-modulating matrix that has the potential to alter the wound environment by reducing the protease activity in the wound and thus stimulate healing.

Sizes
28cm2
123cm2

Indications for use	All chronic wounds free from necrotic tissue healing by secondary	
	intention.	
Contraindications	Known hypersensitivity to any components in the dressing	
How to apply/remove	For optimal effect, apply directly to the whole wound bed. For a wound with low or no exudate apply PROMOGRAN PRISMA and hydrate with saline or Ringer's solution. This will initiate the gel forming process. The biodegradable gel is naturally absorbed over time. It can be cut to size.	
Frequency of dressing changes	The dressing should be changed every 72 hours or more frequently if the exudate level is high. If the gel has not biodegraded the dressing should be left in place until the next dressing change, minimising disturbance to the wound.	
Prescribing guidance	This product must only be initiated by a wound care specialist	

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Atrauman (Hartmann) N-A Ultra (Systagenix)

Low adherence dressings

Description:

N-A Ultra Primary wound contact layer consisting of a knitted viscose rayon sheet with a silicone coating.

Atrauman (Petrolatum free) Non-adherent, polyester mesh wound contact layer (1mm pore size and impregnation of neutral triglycerides prevent penetration of granulation tissue into dressing).

Atrauman sizes	N-A Ultra sizes	
5 x 5cm	9.5 x 9.5cm	
7.5 x 10cm (052295)	9.5 x 19cm	
10 x 20 cm (052301)		
20 x 30cm		
Indications for use	 Provides a contact layer directly onto the wound surface. Basic wound dressing for non-complex wounds: Minor burns Abrasions Superficial wounds as a contact layer under compression bandage on leg ulcers A cost effective alternative to silicone contact layer products when dressings are changed more than once a week 	
Contraindications/cautions	 dressings are changed more than once a week. Can be used under compression; however risk of adherence to wound bed if minimal exudate present. Use with caution on chronic low exuding wounds with viscous exudate which may result in pooling and restricted drainage through dressing pores. 	
	N-A Ultra Do not use if allergic to silicone.	
How to apply/remove	 Apply: Place flat onto the wound surface with 2.5cm border May be applied in multiple layers "fluffed" up to reduce risk of adherence and frequency of dressing changes. Can be cut or folded to size. Removal: Raise corner and peel back off wound. Should lift off wound with no adherence. 	
Frequency of dressing changes	 Dependent on the nature of the wound, can be left in place for up to 7 days; however may require more frequent changes if there is a risk of desiccation or unexpected increase in exudate with need to review dressing regimen and/or more frequent changes. If secondary dressing allows strike through e.g. bandages or dry dressings there is a risk of bacterial ingress with requirement for review of dressing regimen or more frequent changes. Refer to exudate and debridement management guidance (appendix 1 & 2) 	
Prescribing guidance	Usually used for low exuding wounds	

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Atrauman is a cost effective alternative to silicone contact layer dressings, if greater than once a week dressing changes are required.
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CarboFLEX (Convatec)

Odour absorbant dressings

Description: Primary contact non-adherent wound dressing in 5 layers: wound facing absorbent layer containing hydrocolloid and alginate; water resistant second layer; third layer containing activated charcoal; non-woven absorbent fourth layer; water resistant backing layer.

	over absorbent fourth layer, water resistant backing layer.
Sizes	
10 x 10cm	
8 x 15cm oval	
15 x 20cm	
Indications for use	 Discharging, malodorous, sloughy, and moderate to heavily exuding wounds
	 Hydrocolloid and alginate layer will gel where moisture present and sequester exudate, proteases and bacteria into dressing facilitating debridement
	Water resistant layer reduces rate of charcoal becoming wet and ineffective, whilst outer layer reduces risk of strikethrough
	• The underlying cause of wound odour should be identified and wound bed treated appropriately for example debridement
	 CarboFlex dressing may be used as a primary dressing for shallow wounds or with deeper wounds as a secondary dressing over a wound filler.
Contraindications	 Not suitable for dry wounds, as requires moisture to activate gelling process
	 Any known sensitivity to the dressing or its components
How to apply/remove	Select dressing size large enough to overlap the wound edge by 3cm.
Secondary Dressing	Bandage or tape.
Frequency of dressing changes and removal	As exudate and slough dictates – refer to exudate and debridement management guidance (appendices 1&2)
changes and removal	
Prescribing guidance	 Useful in palliative and fungating wounds, as conforms to shape of wound
	Cannot be cut to size
	Suitable for surface and shallow wounds
	 If large cavity or tracking wound, can be used additionally with packing dressing

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CliniSorb (CliniMed)

Odour absorbent dressings Description: A non-adherent activated charcoal cloth enclosed in viscose rayon with outer polyamide coating.

Sizes
10 x 10cm
10 x 20cm
15 x 25cm

Indications for use	 Apply as a primary or secondary dressing.
	 Management of malodorous wounds whilst underlying cause is being addressed (e.g. debridement, management of infection)
Contraindications	None listed
How to apply/remove	Place directly on wound bed or over primary dressing. Can be cut to size.
Frequency of dressing	Can be left in place for up to 7 days, as exudate and slough dictates.
changes	Refer to exudate and debridement management guidance (appendices
	1&2).
Prescribing guidance	Can be cut to size
	 For use in low to moderate exuding wounds
	 Inactivated when wet

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PolyMem_(Non-adhesive) (Aspen Medical) FOR Radiotherapy RTOG damage.

Foam dressings, Polyurethane Foam film dressing without adhesive border Description: Non-adherent thin polyurethane foam dressing with a vapour permeable film backing. Dressing structure contains a wound cleansing agent and glycerol.

Size	
10 x 61cm	

Indications for use	Radiotherapy induced skin reactions
Contraindications	Not suitable for full thickness burns. Do not use in conjunction with solutions containing hypochlorite.
How to apply/remove	Apply directly to wound bed, grid side showing, secure with bandage or tape at edges.
Frequency of dressing changes	As exudate dictates – refer to exudate management guidance (Appendix One)
Prescribing guidance	Seek specialist guidance before use
	 Do not use a foam dressing unless exudate levels and wound conditions indicate appropriate
	 No need to cleanse wound bed as dressing contains cleanser
	 A dramatic increase in fluid may be observed in first few days which should resolve in this time; if not reassess wound.
	DO NOT USE WITH ANY OTHER WOUND CARE PRODUCT, THIS IS
	A PRIMARY DRESSING AND DOES NOT REQUIRE A SECONDARY DRESSING

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Absopad (Medicareple	us int)
Absorbent perforated	plastic film faced dressing
•	ence contact layer dressing consisting of three layers: perforated film
-	ntact layer, absorbent cotton pad and hydrophobic backing
Sizes 10 x 10cm	
20 x 10 cm	
20 % 10 CIII	
Indications for use	Superficial wounds
	Abrasions
	Post op wounds
	Lightly exuding wounds
	Lower depth of these products may be of value in difficult to dress
	areas e.g. toe nail avulsion
Contraindications/	Use with caution on chronic wounds which produce copious or viscous
cautions	exudate. Under these circumstances, the exudate may become trapped
	under the dressing, leading to maceration and inflammation of the
	surrounding skin.
How to apply/remove	Absopad film surface direct to wound bed
	Absopad him surface direct to would bed
Secondary layer	Retention bandage or secure with tape
Frequency of	As exudate dictates – for low or minimal exudate.
dressing changes	
	Patient may prefer to change their own dressing when carrying out
	general social hygiene and to promote independence.
Dress with in a social second	
Prescribing guidance	Consider volume of dressings required for treatment
	Sterile dressings are individually wrapped

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<u>ActivHeal Silicone</u> Foam Adhesive (Advanced Medical Solutions) Foam dressings

Description: A polyurethane foam pad with a waterproof, high moisture vapour transmission rate film backing and adhesive border.

Sizes (pad size	in brackets)	
7.5 x 7.5cm (5 x	(5cm)	
10 x 10cm (6.25	5 x 6.25cm)	
12.5 x 12.5cm (7.5 x 7.5cm)	
15 x 15cm (11 x	(11cm)	
20 x 20 cm (13.	5 x 13.5cm)	
	· · · · · · · · · · · · · · · · · · ·	
Indications for use	Moderate to heavily exuding wounds	
Contraindications	Any known sensitivities	
	Third degree burns	
	Surgical implantation	
	• Do not use with oxidising agents such as hypochlorite solutions or	
	hydrogen peroxide as these can break down the absorbent	
	polyurethane component of the dressing.	
How to apply/remove	Select a dressing large enough so that the pad overlaps the wound	
	edges by 2cm.	
	Centre the dressing on the wound and apply it gently to wound bed.	
Frequency of	Depending on the nature and condition of the wound, may be left in	
dressing changes	place for up to 7 days.	
an eeo mig onangoo	As exudate dictates – refer to exudate management guidance (appendix	
	1)	
Prescribing	Do not use a foam dressing unless exudate levels and wound	
guidance	conditions indicate it is appropriate	

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ActivHeal Silicone Foam Borderless (Advanced Medical Solutions)

Foam dressings

Description: a polyurethane foam dressing with waterproof film backing and perforated wound contact layer (non-adhesive)

	Sizes		
	5 x 5cm	15 x 15cm	
	7.5 x 7.5cm	10 x 20cm	
	10 x 10cm	20 x 20cm	
Indica	tions for use	Suitable for moderate to heavily exuding chronic and acute woundsCan be used under compression	
Contra	aindications	 Third-degree burns. Do not use with oxidising agents such as hypochlorite solutions or hydrogen peroxide as these can break down the absorbent polyurethane component of the dressing. 	
How to	o apply/remove	 Select a dressing large enough to overlap the wound edges by 2cm Dressing can be cut to shape 	
Secon	ndary dressing	Bandage or tape	
-	ency of ing changes	Depending on the nature and condition of the wound, may be left in place for up to 7 days. As exudate dictates – refer to exudate management guidance (appendix 1)	
Presc	ribing guidance	Do not use a foam dressing unless exudate levels and wound condition indicate it is appropriate.	

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Celludress (Medicarep	lus International)
· ·	
Absorbent dressings	
Description: Absorbent	cellulose pad with fluid repellent backing for moderate to heavy exudate.
Sizes	
10 x 10cm	
10 x 15cm	
10 x 20cm	
15 x 20cm	
20 x 25cm	
20 x 30cm	
Indications for use	Basic wound pad
	 Use as primary or secondary dressing for moderate to heavily
	exuding wounds
	Under compression therapy for increased fluid handling capability
Contraindications	None listed
How to apply/remove	Apply blue backing uppermost, facing away from the wound
	Apply blue backing uppermost, facing away from the wound
Secondary dressing	Bandage or tape
Frequency of	A a avudata distatas (ass appandiv 182)
dressing changes	As exudate dictates (see appendix 1&2)
dicooning changes	 If strike through occurs review frequency of change requirement or consider e.g. Kliniderm Superabsorbent
	 If exudate increases review treatment regimen to establish
If exudate increases review treatment regimen to establish underlying cause	
Prescribing guidance	Consider alternative to secondary foam or silicone dressing
	Consider volume to be prescribed for treatment period to avoid waste

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Comfifast (Synergy Health plc)

Tubular bandages and garments (elasticated)

Description: Conformable elasticated viscose stockinet tubular bandage. 92% viscose, 5% elastane, 3% polyamide

Sizes:

colour	sizes width and length	Application	
code	available		
red line	3.5 cm x 1m	small limb (8-15cm)	
green line	5.0cm x 1m, 3m, 5m	small/medium limb (10 – 25cm)	
blue line	7.5 cm x 1m, 3m, 5m	large limb (20 – 45 cm)	
yellow line	10.75cm x 1m, 3m, 5m	extra-large limb, head, children trunk (35-65cm)	
beige line	17.5 cm x 1 m	adult trunk (50 – 120cm)	
Indications for	 holds wound dr Can be used for reduce staining Is not intended 	 Can be used following application of dermatology products to reduce staining to clothing Is not intended as compression therapy 	
Contraindicatio utions		 Ensure Comfifast is correct size is applied by competent practitioner to prevent tourniquet effect, slippage or damage to skin integrity 	
How to	Apply: Measure ar	ea for correct size choice	
apply/remove	Removal: Roll off I	oval: Roll off like a stocking.	
Frequency of dressing chang	When dressing cha	Vhen dressing changes or treatment required	
Prescribing guidance	For irregular sh considered	ned and reused when appropriate shaped limbs Comfifast Multi Stretch may be zes are colour coded to inform correct prescription	

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Hydrofilm (Hartmann)

Vapour permeable film dressing (Semi-permeable Adhesive Dressing) Description: Conformable adhesive waterproof film dressing with high moisture vapour transmission rate.

Sizes	
6 x 7cm	12 x 25cm
10 x 12.5cm	15 x 20cm
10 x 15cm	20 x 30cm
10 x 25cm	

Indications for use	 Superficial wounds with minimal exudate Abrasions Provide showerproof bacterial barrier wound contact layer on post op incisions
Contraindications/cautions	 Heavily exuding wounds Fragile skin if risk of skin tears Known sensitivities to dressing components
How to apply/remove	 To apply: Remove film backing paper Apply direct to wound surface with approximately 2.5 cm border Peel off frame and smooth edges. The frame prevent stretching the dressing to apply and reduces risk of discomfort and skin tears For removal: Gently lift corner and pull backwards towards centre of wound and stretch off For further advice refer to manufacturer's instructions
Frequency of dressing changes	 May be left in situ up to seven days. Transparency of dressing will allow assessment of wound bed without removal, to inform frequency of changes
Prescribing guidance	Consider volume of dressings required based on number of wound care interventions required

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Hydrofilm Plus (Hartmann)

Vapour permeable adhesive film dressing with absorbent pad Description: Conformable adhesive waterproof film dressing with high moisture vapour transmission rate and adsorbent island pad

Sizes	
7.2 x 5cm	10 x 20cm
9 x 10cm	10 x 25cm
9 x 15cm	10 x 30cm

Indications for use Contraindications/ cautions	 Low exuding wounds Provide showerproof bacterial barrier wound contact layer Minor traumatic wounds such as grazes, abrasions and lacerations Post-operative surgical wounds Superficial burns Can be used as a secondary dressing Hydrofilm should not be used as a primary dressing on clinically infected, bleeding or heavily secreting wounds Known sensitivities
How to apply/remove	Apply:
	Remove film backing
	 Apply to wound ensuring absorbent pad is covering wound bed or incision line Peel off frame following application to prevent stretching skin and risking epidermal blistering and smooth edges Removal: When absorbent lift corner and pull backwards towards centre of wound For further advice refer to manufacturer's instructions
Frequency of dressing	When absorbent pad is 80% discoloured change dressing or earlier
changes	if wound assessment dictates
	 If exudate level results in greater than 2-3 times per week changes, or exudate becomes more discoloured or viscous reassess treatment regimen
Prescribing guidance	Film allows inspection of wound and surrounding skin when used
	as a primary dressing
	Low absorbency capacity

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Kliniderm Foam Silicone Border (Aria Medical)

Soft polymer dressings with absorbent pad with ADHESIVE BORDER Description: Absorbent foam dressing with a soft silicone wound contact layer (non-adherent) and adhesive border plus a waterproof vapour-permeable polyurethane (film) backing.

Sizes	
7.5cm x 7.5cm	
10cm x 10cm	
12.5cm x 12.5cm	
15cm x 15cm	
10cm x 20cm	
15cm x 20cm	
Indications for use	Suitable for exuding chronic and acute wounds
Contraindications	Do not use if allergic to silicone or any other components of the dressing
How to apply/remove	 Ensure wound margins are dry Apply directly to wound bed ensuring the dressing overlaps the wound margins by at least 2cm. Remove dressing by gently lifting one corner and slowly peel back the dressing.
Frequency of Dressing changes	May be left in place for up to 7 days depending on wound exudate. Refer to exudate and debridement management guidance (appendices 1 & 2).
Prescribing guidance	Do not use a foam dressing unless exudate levels and wound condition indicate it is appropriate.
	Sloughy wounds may initially appear to increase in size due to autolytic debridement promoted by the moist conditions produced beneath the dressing.
	Do not use with oxidising solutions such as hypochlorite or hydrogen peroxide

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Kliniderm Foam Silicone Non- Border (Aria Medical)

Soft polymer dressings with absorbent pad Description: A soft conformable absorbent polyurethane foam with a silicone wound contact layer and a moisture permeable backing

Sizes	
5cm x 5cm	
10cm x 10cm	
15cm x 15cm	
20cm x 20cm	

Indications for use	Suitable for exuding chronic and acute wounds
Contraindications	Do not use if allergic to silicone or any other components of the dressing
How to apply/remove	 Ensure wound margins are dry Select a suitable size so that the dressing overlaps the wound margins by at least 2cm Gently apply directly onto the wound site Secure with simple secondary dressing Remove dressing by gently lifting one corner and slowly peel back the dressing.
Frequency of Dressing changes	May be left in place for up to 7 days depending on wound exudate. Refer to exudate and debridement management guidance (appendices 1 & 2).
Prescribing guidance	 Do not use a foam dressing unless exudate levels and wound condition indicate it is appropriate. Sloughy wounds may initially appear to increase in size due to autolytic debridement promoted by the moist conditions produced beneath the dressing. Do not use with oxidising solutions such as hypochlorite or hydrogen peroxide

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Knit Band (Clinisupplies)

Lightweight conforming bandages

Description: Lightweight knitted polyamide and cellulose contour retention bandage

Sizes:	
5cm x 4m	
7cm x 4m	
10cm x 4m	
15cm x 4m	

Indications for use	Dressing retention
Contraindications/ cautions	 Bandage should be applied by practitioner to prevent tourniquet effect, slippage or damage to skin integrity Allow for swelling following application of product between changes to ensure there is no constriction
How to apply/remove	Apply:
	Bandaging is a basic procedure but if carried out incorrectly it has the potential to cause considerable harm, for example by restricting movement or blood flow.
	Bandages can be used to fix or retain a primary dressing product. In some instances, the bandage is simply wrapped around the affected area and secured with tape
	Removal: Unwind bandage.
	To avoid trauma, particularly if bandage is in direct contact with skin, do not remove using scissors, loosen and unwind dressing.
Frequency of dressing changes	When wound dressing change dictates
Prescribing guidance	Volume of bandages should be in line with number of dressing changes

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KSoft (Urgo)

Sub compression wadding bandage

Description: Soft absorbent padding layer

Sizes 10 cm x 3.5m 10 cm x 4.5m	
Indications for use	 Normally used as sub compression wadding layer for shaping and protecting bony prominences under compression bandages May also be used for padding, protecting bony prominences and extra absorbency on limbs under retention bandages
Contraindications/ cautions	None listed
How to apply/remove	Apply: as directed dependent on purpose
Secondary Dressing	Retention bandage
Frequency of dressing changes and removal	As wound dressing or exudate dictates
Prescribing guidance	Consider purpose of product, absorbent pads may also be used if extra absorbency required

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Premierpore (Shermond)

Absorbent dressings Description: An absorbent perforated dressing with adhesive border.

Sizes (pad size in brackets)
5 x 7cm (3 x 4cm)
10 x 10cm (6 x 5cm)
10 x 15cm (5 x 10cm)
10 x 20cm (5 x 15cm)
10 x 25cm (5 x 20cm)
10 x 30cm (5 x 25cm)
10 x 35cm (5 x 30cm)

Indications for use	 Post-operative incision sites
	 Lightly exuding wounds
Contraindications	Any known sensitivity to adhesives
How to apply/remove	Place directly over wound ensuring the absorbent pad covers the wound and/or suture line
	Removal: Lift one corner and peel back gently.
Frequency of dressing changes	 Post-operative dressings should be removed 48 hours post op or as per surgeons instructions
	 Remove and inspect wound if a large amount of exudate is visible on the outer dressing
	Refer to exudate and debridement management guidance (appendix 1 & 2)
Prescribing guidance	Care must be taken on removal to prevent skin stripping
	• Do not use as primary dressing on wounds with moderate to heavy levels of exudate; this will result in strike through, increased risk of bacterial contamination and increased frequency of dressing changes

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Telfa pad (Aria Medical)

Absorbent perforated plastic film faced dressing non adhesive Description: low adherence contact layer dressing consisting of three layers: perforated film polyester film wound contact layer, absorbent cotton pad and hydrophobic backing

Sizes:

Telfa sterile	
5 cm x 7.5 cm	
10 cm x 7.5 cm	
7.5 cm x 15 cm	
7.5 cm x 20 cm	
Telfa non sterile	
20 cm x 7.5 cm	
25 cm x 20 cm	
Indications for use	Superficial wounds
	Abrasions
	Post op wounds
	Lightly exuding wounds
	• Lower depth of these products may be of value in difficult to dress areas
	e.g. toe nail avulsion
Contraindications/	Use with caution on chronic wounds which produce copious or viscous
cautions	exudate. Under these circumstances, the exudate may become trapped
	under the dressing, leading to maceration and inflammation of the
	surrounding skin.
How to	Telfa can be applied any side down
apply/remove	
Secondary layer	Retention bandage or secure with tape
Frequency of	As exudate dictates – for low or minimal exudate.
dressing changes	
Prescribing	Consider volume of dressings required for treatment
guidance	Sterile dressings are individually wrapped
	Telfa is also available in non-sterile form, if required in bulk

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Biatain Silicone (Coloplast)

THIS PRODUCT MUST ONLY BE INITIATED BY A WOUND SPECIALIST AND SHOULD ONLY BE USED ON WOUNDS OF LESS THAN 2CM DEEP AND SHOULD NOT BE USED WITH ANOTHER WOUND PRODUCT

Foam

Description: Biatain silicone is a soft conformable polyurethane dressing with semi-permeable and waterproof top film, a lock-away pad and soft silicone adhesive layer. The pad absorbs vertically and retains exudate and bacteria.

It has a unique and patented 3D Fit technology, conforming to the wound bed up to 2cm in depth to fill the dead space and reduce exudate pooling for optimal healing. There is no requirement for a wound filler unless the wound is greater than 2cm in depth.

Sizes:				
7.5x7.5cm	12.5x12.5cm		Multi-shape 14x19.5cm	
10x10cm	15x15cm		Sacral 15x19cm	
10x20cm	17.5x17.5c	m		
10x30cm	Heel 18x18cm			
Indications for use	including acute wounds such as donor sites; postoperative and traumatic wounds; and chronic wounds such as leg ulcers, pressure ulcers and non-infected diabetic foot ulcers. May be used to prevent postoperative blistering. Can be used under compression therapy		nor sites; postoperative and nds such as leg ulcers, abetic foot ulcers. May be ring. Can be used under	
		 allergic reaction to the dressing or its components. Do not use with oxidising solutions such as hypochlorite and hydrogen peroxide solutions. Ensure that any other evaporating solution is completely dried off before dressing application. 		
How to apply/remo	ove	dry an Removinot tou the dre	d free from emollients.	-
			applied, run your fingers arou e contact between skin and si	0
			val : gently lift the border of th intil fully removed.	e dressing and slowly pull
Frequency of dress changes	sing		n Silicone can be left in place nount of wound fluid and type	for up to 7 days, depending on of wound. The dressing

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	should be changed when there is 1cm between the exudate and the edge of the foam pad.
Prescribing guidance	This product can only be initiated by a wound specialist

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Kliniderm (H and R Healthcare)

Silicone Wound Contact Layer

Description: One sided soft silicone wound contact layer with porous structure that allows exudates to pass into an outer absorbent dressing. The one sided layer prevents the secondary dressing from stocking to the wound contact layer. Can also be used as a protective layer in negative pressure wound therapy.

Sizes	
5cm x 7.5cm	17cm x 25cm
7.5cm x 10cm	20cm x 30cm
10cm x 18cm	

Indications for use	Can be used in acute and chronic wounds	
Contraindications	Not to be used on third degree burns or with patients with known hypersensitivity to any of the components of the dressing. Not for surgical implantation.	
How to apply/remove	 Ensure peri wound skin is dry Ensure 1cm overlap over wound margins Remove protective film and apply directly to the wound and smooth down the edges Cover with appropriate secondary dressing 	
Frequency of dressing changes	As wound exudate dictates	
Prescribing guidance	N/A Ultra or Autraman should be considered before using Kliniderm contact	

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Cutimed Sorbion Sachet S (Essity)

THIS DRESSING MUST ONLY BE INITIATED BY A WOUND CARE SPECIALIST

Superabsorbent dressing with Hydration Response Technology. This is a Primary dressing not a secondary dressing.

Description:

Superabsorbent dressing with Hydration Response Technology. It has a hypoallergenic polypropylene outer sheath and an inner core consisting of superabsorbent polymers and cellulose. Does not contain glue or adhesives. Can be used under compression.

Sizes	
7.5cm x 7.5cm	15cm x 15cm
12cm x 5cm	20cm x 20cm
10cm x 10cm	20cm x 30 cm
20 cm x 10 cm	

Indications for use	For wound cleansing and debridement with moderate or high exudates levels.	
Contraindications/ cautions	Do not apply to dry wounds or wounds with low exudates levels Only apply to tunnel cavities with caution as product expands as it absorbs fluid.	
How to apply/remove	 Apply: Directly to wound bed that has high exudate Ensure at least 1.5cm margin of dressing beyond wound edge Do not use greasy barrier products under dressing as this compromises abseorbency Do not use restrictive circumferential tape to hold in place as the dressing swells and may cause ischaemia if resticted Removal: Should come away from wound bed easily if used appropriately on exudating wound. It may adhere if used on wound that has low exudate 	
Secondary Dressing	Do not use circumferential tape or film as this may restrict and cause ischaemia. Simple non restrictive retention bandage Frame with tape or film	
Frequency of dressing changes and removal	As wound exudates regimen dictates	
Prescribing guidance	A wound care specialist must initiate this dressing Do not use on wounds with no or low exudate	

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Kliniderm Super Absorbent (Aria Medical)

Absorbent dressings

Description: Superabsorbent polymer/cellulose dressing with fluid repellent backing.

Sizes:	
10 x 10cm	
10 x 15cm	
20 x 20cm	
20 x 30cm	
20 x 40cm	
Indications for use	Basic wound pad
	• Use as primary or secondary dressing for heavily exuding wounds
	As a contact layer under compression bandage on leg ulcers
	 To provide excess exudate management for oedematous legs
	due to chronic venous insufficiency
	Kliniderm Super Absorber is low profile and can be used when less bulk is required
Contraindications/cautions	
	adherence
	 Do not use with larvae therapy. Outer waterproof layer will
	• Do not use with larvae therapy. Outer waterproof layer will suffocate larvae
	Suilocale laivae
How to apply/remove	Direct to wound bed, or as secondary dressing over primary
	dressing.
	droconig.
Secondary dressing	Bandage or tape
, , , , , , , , , , , , , , , , , , ,	
Frequency of dressing	As exudate dictates – refer to exudate and debridement
changes	management guidance (appendix 1&2)
Prescribing guidance	Cost effective alternative to foam or silicone dressings when used
	as secondary dressings

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Vliwasorb Pro (formerly Flivasorb) (Lohmann & Rauscher)

Soft polymer dressings

Description: Superabsorbent wound dressing with non-adherent wound contact layer and outer clothing protection layer. Contains sodium polyacrylate super absorber particles and cellulose that form a gel on contact with fluid.

Sizes		
12.5 x 12.5cm		
12.5 x 22.5cm		
22 x 22cm		
22 x 32cm		

Indications for use	 Primary dressing for the management of heavily exuding and sloughy wounds 	
	 Secondary dressing for deep heavily exuding wounds 	
	Can be used under compression bandaging	
Contraindications	 Known sensitivity to any components of the dressing 	
	Lightly/non-exuding wounds	
	Cavity wounds	
How to apply/remove	Direct to wound bed	
Secondary dressing	Bandage or tape	
Frequency of	As exudate dictates – refer to exudate and debridement management	
dressing changes	guidance (appendix 1 & 2)	
Prescribing guidance	 Reduces the need for secondary foam or silicone dressing 	
	Dressing must not be cut or torn	

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Zetuvit E non sterile (Hartmann)

Superabsorbent dressing

Description: Absorbent cellulose pad with fluid repellent backing for moderate to heavy exudate.

Sizes: 10 x 10cm 10 x 20cm 20 x 30cm 20 x 40cm		
Indications for use	 Basic wound pad Use as primary or secondary dressing for moderate to heavily 	
	exuding wounds	
	 Under compression therapy for increased fluid handling capability 	
Contraindications	None listed	
How to apply/remove	Apply blue backing uppermost, facing away from the wound	
Secondary dressing	Bandage or tape	
Frequency of	As exudate dictates (see appendix 1&2)	
dressing changes	 If strike through occurs review frequency of change requirement or consider Kliniderm Superabsorbent 	
	 If exudate increases review treatment regimen to establish underlying cause 	
Prescribing guidance	Alternative to secondary foam or silicone dressing	
	 Consider volume to be prescribed for treatment period to avoid waste 	

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Seal-Tight Wound Care Protector (Autonomed)

Category: Wound care accessory Description: Waterproof dressing protector for lower limb

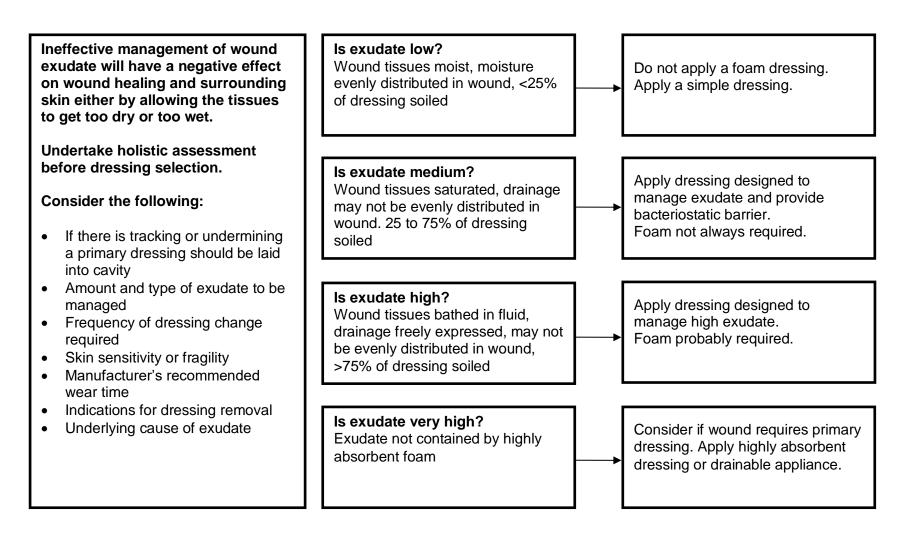
Sizes	
Adult short leg	CV27103
Adult wide short leg	CV27106
Adult foot/ ankle	CV27105

Indications for use	To keep wound dressings and compression bandages dry when bathing for patients with chronic wounds		
	For further information and support contact podiatric team or vascular nurse specialist.		
Contraindications	 Acute wounds where healing is projected to take place within 4 weeks Patients who are able to or have assistance to self-manage dressing changes Patients who have ability to remove their own dressing, can shower/bathe and apply a temporary cover prior to new dressing application by a health care professional Inability or has no assistance to apply and remove Seal-Tight[™] Routine prescribing for acute limb fractures with casts 		
How to apply/remove	 Pull diaphragm of device open as per manufacturer's instructions Put foot through the opening and pull device up and over any dressings/bandages Check seal prior to bathing by pulling device up a few cm then pushing it down. The device should have a visible flare at the top Remove by expanding the diaphragm again and pulling device down over the foot Avoid using any chemicals to clean the device Hang up and allow to dry between use 		
Frequency of dressing changes	N/A		
Prescribers guidance	Check appropriate sizing to ensure correct choice of product and effective seal is achieved. Product is one per patient limb and should not require repeat prescriptions. Device should last at least 6 months before replacement required		
Acute variation	NA		

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Appendix 1

Exudate management guidance notes



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Appendix 2

Debridement Guidance

<u>Definition:</u> the removal of dead non-viable/devitalised tissue, infected or foreign material from the wound bed and surrounding skin

Non-viable tissue is detrimental to healing in the following ways:

-is a physical barrier to healing -reduces the effectiveness of topical antimicrobials -can mask or mimic signs of infection -can delay wound healing by contributing to prolonged inflammatory response -can be a barrier to comprehensive wound assessment -can increase exudate and odour

Debridement is an important aspect of wound bed preparation and facilitates wound healing. Following structured holistic assessment, decision to debride and selection of method can be made (see Figure 1)

Types of Debridement

Autolytic: the naturally occurring process in which the body's own enzymes and moisture rehydrate, soften and liquefy devitalised tissue. Can be facilitated by dressings which promote debridement through donation of moisture-i.e. hydrogels or hydrofibre (Generalist) Mechanical: using a moistened, soft mono filament pad to physically remove moist, loose slough (Generalist) Larval (Bio-Surgical): Larvae from the green bottle fly ingest and secrete enzymes to breakdown devitalised tissue. Available loose or contained small bags for application to the wound bed (Generalist) Ultrasonic: delivery of ultrasonic sound waves in combination with irrigation to remove devitalised tissue (Specialist) Hydro surgical: delivery of high pressure saline jet to remove devitalised tissue (Specialist) Sharp: using scissors, a scalpel and/or forceps above tissue level to remove devitalised tissue (competent practitioner) Surgical: excision or wide resection of devitalised tissue in a theatre setting (Specialist)

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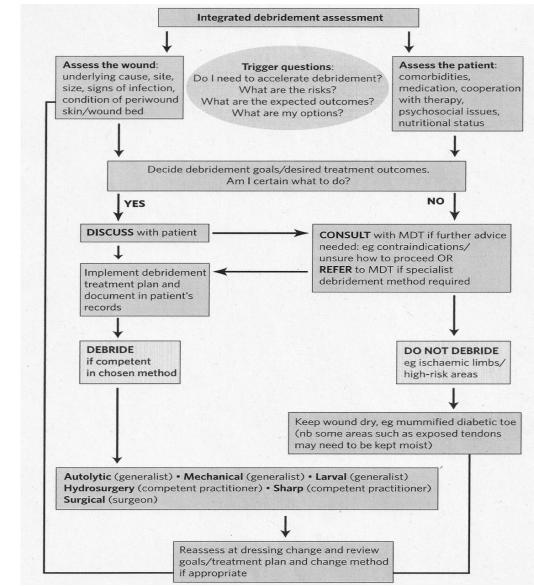


Figure 1

Note:

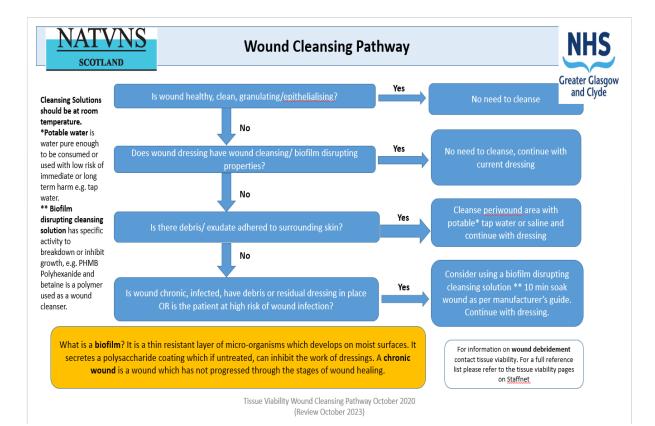
Please seek specialist advice if further support on any aspects of debridement is required.

If patient unable to give consent please discuss with carer.

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Appendix 3

Wound Cleansing Pathway



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Appendix 4

Useful Links to related guidelines and resources

Acute Care Paediatric and Neonatal Wound Management Formulary

Acute Care Wound Resource Folder – Guidelines & Tools

Antimicrobial wound dressings (AWDs) for chronic wounds: Health Technology Assessment <u>13 -</u> Healthcare Improvement Scotland (HIS) guidance on antimicrobial wound dressings

<u>GGC Medicines</u> homepage. A tool to assist in promoting high quality, safe and cost-effective prescribing within the Greater Glasgow and Clyde Health Board. The site includes formulary information (including Stoma Care Joint Formulary, Wound Product Formularies & Urology Formulary; accessed under <u>Other Formularies</u>), BNF, clinical guidelines and resources (including information on GGC medicines app).

Negative Pressure Wound Therapy Systems guideline

NHSGGC Code of Business Conduct for staff

NHSGGC Tissue Viability Service site

Prescribing Larvae, Unlicensed Medicine Protocol

Single use negative pressure wound therapy guideline

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