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| EXECUTIVE SPONSOR or EXECUTIVE CLINICAL SPONSOR | Dr Greg Keogh  
Surgery stream Director |
| AUTHOR                 | SESLHD Wound committee     |
| POSITION RESPONSIBLE FOR THE DOCUMENT | Carol Stott  
Carol.Stott@health.nsw.gov.au |
| KEY TERMS              | Compression, venous, leg ulcers, cellulitis |
| SUMMARY                | This document outlines the appropriate use of compression therapy for the treatment of venous leg ulcers and lower limb cellulitis. |
1. POLICY STATEMENT
Compression therapy must provide safe and effective treatment for patients with venous leg ulcers and lower limb cellulitis. Where uncertainty about the appropriate use of compression exists, the clinician must seek a review of the patient by a wound care expert, a medical officer or a vascular specialist. This policy is not applicable for the management of Lymphoedema.

Compression whether it be bandages, compression wraps, stockings or intermittent pneumatic compression is a therapeutic treatment and should not be discontinued until all the ramifications of this have been discussed with the patient and carers. Alternative methods of compression intervention should be investigated should the patient not want to wear or is not able to tolerate compression bandages.

The application of compression bandages must not put the patient at falls risk, therefore, when contemplating the type of compression to be used consider how safe footwear can be achieved.

2. BACKGROUND
Compression therapy will increase the healing rate of most venous leg ulcers and will reduce the likelihood of a recurrence\(^1\). Compression is the primary intervention in the prevention and management of venous hypertension, venous oedema and venous leg ulcers\(^2\). Where the leg ulcer has a mixed aetiology of arterial and venous disease, the lower leg arterial occlusion must be addressed prior to application of compression. Compression therapy may also be beneficial in the treatment of lower limb cellulitis. Compression therapy includes compression bandages, compression wraps, compression garments and intermittent pneumatic compression (IPC) systems. The aim of compression therapy is to improve calf muscle pump function, improve venous return, reduce venous hypertension, control venous oedema and facilitate the healing of venous leg ulcers\(^2\).

Definitions
**Ankle Brachial Pressure Index (ABPI):** Ratio of ankle arterial systolic blood pressure to brachial pressure\(^6\).
**Compression Bandages:** Can be cotton and/or synthetic, with or without elastic or latex and are described as short or high/long stretch bandages. A three layer tubular system can deliver compression equal to a short stretch bandage system\(^5\) [Appendix K].
**Compression Garments:** Include manufactured graduated compression hosiery.
**Compression Levels:** Vary depending on type of pressure required, [Appendix I].
**Compression Scales:** Guide compression prescription, [Appendix I].
**Compression Therapy:** Graduated compression can be achieved through compression stockings or compression bandages. This also includes the use of intermittent pneumatic compression pumps.
**Light / Mild Compression Therapy:** Is not an effective treatment for venous leg ulcers. However, higher pressure is better than lower pressure and some pressure is better than no pressure\(^3\).
**Graduated Compression:** Is achieved by applying a bandage at a steady and even pressure from toes to below knee and can be also achieved with a compression stocking.
High / Long stretch bandages: Provide both a high resting pressure and a high working pressure.

**Intermittent pneumatic compression (IPC) systems:** Intermittent pneumatic compression (IPC) is a mechanical method of delivering sequential compression to swollen limbs.

**Lanarkshire Oximetry Index (LOI):** A protocol for pulse oximetry toe / finger O² saturation to check the suitability of compression therapy.

**Short stretch bandages:** Provide a low resting pressure and a high working pressure.

**Toe Brachial Pressure Index (TBPI):** A procedure to determine arterial perfusion in the feet and toes by measuring the systolic pressure in the arm and the great toe.

**Wound Care Expert:** A person with advanced training in wound management and recognised within the facility e.g. CNC Wound Care, CNC Stomal Therapy and Wound Care and Nurse Educators.

### 3. RESPONSIBILITIES

#### 3.1. Employees will:
Ensure that they work within their scope of practice and attend relevant education related to this procedure.

#### 3.2. Line Managers will:
Ensure all clinical staff are given the opportunity to attend District wound management education and that all nursing staff work within this procedure and have appropriate resource and stock items to implement the recommendations within this procedure.

### 4. PROCEDURE

#### 4.1. Prior to application of compression significant arterial disease should be excluded. Arterial disease can be determined by physical examination and the following tests:

- Ankle brachial pressure index (ABPI) every six months
- Toe brachial pressure index (TBPI) every six months
- Lanarkshire Oximetry Index (LOI) every six months
- Arterial/venous duplex every 12 months
- Arteriogram.

#### 4.2. When compression has been ordered without any of the above, the Lanarkshire Oximetry Index can be undertaken to ensure the arterial circulation is not compromised by the application of compression therapy. This should be undertaken by a clinician trained in this method. If results are within normal limits, compression can be applied as per policy / written order. If results are outside the normal limits contact a wound care expert to discuss the results.

#### 4.3. Prior to application of compression bandages:

##### 4.3.1. In the hospital setting:
- A wound care expert, medical officer, or vascular specialist should document an order in the clinical notes. This should include the type and level of compression.

##### 4.3.2. In the community setting:
- The medical officer or vascular specialist should provide the community nurse with a letter which includes the level of compression, the date of arterial test and the results of this (example Appendix A).
4.4. Registered and enrolled nurses are not permitted to apply compression until they have gained specific education (determined at a local level) in the application and use. The correct degree of compression must be applied and the correct application technique must be used see:

- Appendix B  Complications following the Application of Compression
- Appendix C  Specific Compression Bandage Systems
- Appendix D  Four Layer Bandage Systems
- Appendix E  Two Layer Compression System
- Appendix F  Short Stretch (Inelastic) Compression Bandage
- Appendix G  High Stretch (Elastic) Compression Bandage
- Appendix H  Intermittent Pneumatic Compression (IPC)

4.5. Decisions about the compression system should consider the following issues:

- The shape and size of the leg, unusually shaped legs may require custom made compression garments
- Patient tolerance and preference
- Clinical experience in application
- Environment e.g. temperature
- Ease of application and removal
- Access to compression systems
- Presence of comorbidities
- Level of the individual’s activity

4.6. The patient’s psychological and social factors must be considered in the selection of appropriate compression garments, as they may have difficulty accepting compression therapy due to its effect on work, showering / bathing, choice of clothing and footwear. Climate and cultural factors should also be considered. All product options should be discussed and the compression level / scale Appendix I and type chosen should encourage concordance.

4.7. The patient / carer should be educated on the importance of concordance and of possible complications and problems arising as a consequence of the compression. Appendix B

4.8. Education should include signs and symptoms of arterial compromise, pain management and the management of loose, slipping and wet bandages. Advice should be given about appropriate footwear and manufacturers guidelines regarding laundering and replacement of bandages or stockings. Garments should be discarded and replaced according to the manufacturer’s recommendations. For further information please refer to Appendix J.

4.9. Some wound management products are not suitable for use under compression, e.g. thick dressing products and hydrocolloids. Discuss product selection with a wound care expert if unsure.

4.10. Compression bandages should be applied as per the manufacturer’s instructions and in a manner which will achieve graduated compression.

4.11. Compression therapy of $\geq 40\text{mmHg}$ at the ankle should only be used where the arterial investigations have indicated that there is no significant arterial disease e.g. ABPI or LOI is 0.8-1.3, TBPI $>0.7$. 
4.12. Caution should be exercised if compression has been prescribed for a patient with an ABPI or LOI of less than 0.8 or a TBPI of less than 0.7. Always consult a wound care expert before applying compression therapy in these clients.

4.13. The ankle should be measured prior to the application of compression bandages.

4.14. For ankle sizes less than 18cms apply extra padding to the ankle / lower leg area until the 18cms is reached at the ankle and the calf is proportionally larger than the ankle.

4.15. For patients with an ankle circumference of around 18cm, regular measurement of the ankle is recommended as these patients are at risk of complications caused by the compression.

4.16. For ankles greater than 25cms alteration maybe needed in compression therapy application Appendixes C - H.

4.17. For ankles greater than 30cms consider the use of IPC Appendix H.

4.18. For limbs that are not conical shaped or misshaped consider the use of IPC Appendix H.

4.19. A natural padding bandage is required under all compression bandages to protect the skin.

4.20. Following application of compression the patient should be observed for pain, pressure damage, loss of calf muscle and skin problems Appendix B.

Note: in clients unable to verbalise pain increased pain may present as delirium.

4.21. Directly following application of compression the patient’s limb should be observed for changes in colour / perfusion. If there is a change in perfusion remove compression therapy.

4.22. In the community, compression therapy should not be applied unless the client or carer can remove it if problems arise such as severe pain or changes in colour / perfusion.

4.23. Pain scores should be measured before and after application of compression therapy with reference made to increases in scores or changed sensation as appropriately applied compression should reduce pain. If pain persists remove compression and ensure arterial status has been adequately assessed.

4.24. A compression garment must extend from just proximal to the toes to two fingers widths below the knee. The foot should be positioned at 90 degrees to the leg during application to avoid the bandage wrinkling during standing or walking. A figure of eight technique can be used to anchor the bandage to the foot. Adequate padding is essential to protect bony prominences and to achieve a conical limb shape for patients who have altered leg contour e.g. ‘champagne bottle legs’.

4.25. Changes of limb shape due to reduced oedema should be monitored by measuring circumference at defined sites (ankle and calf). This can be achieved by asking the patient to place the foot flat on the floor and recording the distance from the floor to the site of measurement.

4.26. Excess bandage should be ‘taped off’ or ‘cut off’ as winding around the limb or turning it over can impair circulation. If one bandage does not adequately cover the leg a second bandage should be used. Finishing the bandaging too low or applying increased stretch to reach the knee may result in adverse client outcomes.

4.27. Application of compression bandages can cause injury to the clinician or carer. The patient should be positioned to ensure easy access to the leg. Appropriate posture throughout the procedure must be maintained. A position must be assumed which will
minimise twisting, reaching and bending. Avoid squatting and kneeling for long periods whilst applying garments. Take breaks as necessary between garment layers and between legs. Avoid rushing the procedure as this may result in inappropriately applied garments and increase the risk of injury.

4.28. Reduced compression
4.28.1. A reduced compression pressure may be used in patients:
- Initially upon commencing compression therapy, the level of compression should be increased as the patient’s tolerance improves
- Where a mild degree of arterial impairment exists (consultation with wound care expert should occur prior to commencement of compression for these clients)
- If ordered by treating specialist and tolerated by the client
- During periods of infection where pain may be increased
- Where tolerance of optimal compression levels is unable to be obtained

4.28.2. A three layer tubular bandaging system may be considered if the patient is unlikely to tolerate full compression. The outer layers can be removed by the patient if required. This system can be used if the applicator does not have the expertise to apply other compression systems\(^5\) Appendix J for appropriate sizing and application).

4.28.3. Moderate compression can be unsafe or painful for patients with arterial insufficiency, neuropathy or cardiac failure. Mild or light compression may be required\(^4\).

4.29. Patient unable to tolerate compression bandage
4.29.1. Alternatives to compression bandages should be considered:
- IPC Appendix H
- Compression wraps e.g. Farrow wraps™ (BSN), Ready wraps™ (Cosmac), CircAid™ (Reis Orthopaedics)
- These are a short-stretch compression system designed for patients with fluctuating oedema, rebound oedema, problems getting compression stockings on or off for patients who are unable to tolerate bandaging. The overlapping bands provide support and rigidity to control oedema. They may be able to be removed and reapplied by a patient. These wraps can also be used on a patient with open wounds. The wrap needs to be specifically measured to fit the patient to ensure graduated compression will be achieved.

4.30. Patients unsuitable for compression bandage due to safety concerns, educate patient on leg elevation and refer to medical officer for review.

4.31. Compression post healing of venous leg ulcer
4.31.1. Compression needs to be continued for life unless surgical intervention is an option and is successful.

4.31.2. Once the venous leg ulcer has been closed for one month consider:
- If the patient could be reviewed by a vascular specialist for possible vascular surgery to prevent recurrence
If the patient can be fitted with appropriate compression stockings, if they are able to get the stockings on and off
If the patient is unable to get the stockings on and off consider alternatives e.g. compression wraps or referral for home assistance package.

4.32. **After hours compression on patient with cellulitis**

4.32.1. After undergoing a physical assessment by a medical officer, compression can be initiated:
- When all leg pulses are present i.e. femoral, popliteal, posterior tibia, dorsalis pedis
- The patient has no known co-morbidities
- The patient or their carer are capable of taking off bandages if they are too painful
- The patient has a referral to an appropriate service for testing to occur as soon as possible.

5. **DOCUMENTATION**

SESLHD wound assessment and management plan SEI060.118 or the Electronic equivalent e.g. in Ambulatory and Primary Health Care (APHC) use Wound Assessment Treatment Evaluation Plan (WATEP).

6. **AUDIT**

Nil

7. **REFERENCES**

3. Australia and New Zealand Clinical Practice Guidelines for Prevention and Management of Venous Leg Ulcers (2011)

8. **REVISION AND APPROVAL HISTORY**

<table>
<thead>
<tr>
<th>Date</th>
<th>Revision No.</th>
<th>Author and Approval</th>
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| February 2015 | 0            | Area wound committee  
              |               | Endorsed by Executive Sponsor |
| April 2017   | 1            | Minor amendment to Appendix A |
| April 2018   | 2            | Minor amendment to Appendix E. Approved by Executive Sponsor |
| May 2018     | 2            | Processed by Executive Services prior to publishing. |
Appendix A: Example of Doctor Letter

<table>
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<tr>
<th>Surname:</th>
<th>Mrn:</th>
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<tbody>
<tr>
<td>South Eastern Sydney Local Health District</td>
<td></td>
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</table>

**Authority to Apply Compression Therapy**

I Dr (please print)…………………………………………….. give permission for registered nurses to apply compression therapy for the above client/ patient (Please select compression below).

Please specify which limbs/limb compression to be applied:……………………………………

Signed Dr/ Specialist:…………………………………………….. Date:……………………………………

Print Name:…………………………………………….. Phone number:……………………………………

**Allergies:** ………………………………………………………………………………………………

In the last six months has this patient had the following (indicate by √):

- □ Ankle Brachial Pressure Index Date……………………………………
- □ Toe Brachial Pressure Index Date……………………………………
- □ Lankshire Oximetry Index Assessment Date……………………………………
- □ Vascular studies Date……………………………………

Results……………………………………………………………………………………………………

………………………………………………………………………………………………………………

**Medical Officer/Specialist Vascular Diagnosis:** …………………………………………………

………………………………………………………………………………………………………………

**Vascular Specialist Name:** ……………………………………………………………………..

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<thead>
<tr>
<th>Graduated Compression Therapy: Bandages</th>
<th>Graduated Compression Therapy: Stockings</th>
</tr>
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<tbody>
<tr>
<td><strong>Moderate:</strong> 2 layer 20-40mmHg: padding / short stretch e.g. Comprilan, Coban 2…………………..</td>
<td>□ Very Strong: &gt;60mmHg (Lymphoedema)………..</td>
</tr>
<tr>
<td><strong>Moderate:</strong> 2 layer 20-40mmHg padding / high stretch e.g. Surepress, Setapress…………………..</td>
<td>□ Strong: 40-60mmHg (Class three stocking) ……</td>
</tr>
<tr>
<td><strong>Moderate:</strong> 4 layer high stretch 20-40mmHg e.g. Profore, Veno4………………………………………..</td>
<td>□ Moderate: 20-40mmHg (Class two stocking)…..</td>
</tr>
<tr>
<td><strong>Moderate:</strong> Zinc Bandage / then padding /short stretch or high stretch compression…………………..</td>
<td>□ Mild: 18-24mmHg (Class one stocking)…………</td>
</tr>
<tr>
<td><strong>Light:</strong> Tubular bandage e.g. Tubigrip, TubularForm, Flexigrip …………………………………………..</td>
<td>□ Light: 15mmHg tubular system (e.g. three layer Tubular Form system)…………………………………</td>
</tr>
<tr>
<td><strong>Ex Light:</strong> 5mmHg tubular system (single layer) Note Not used for VLU……………………………………..</td>
<td>□ Ex Light: 5mmHg tubular system (single layer) Note Not used for VLU……………………………………..</td>
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</tbody>
</table>

**Other:** …………………………………………………………………………………………………… |

**Comments:** (Recommended dressing for wound review only)

…………………………………………………………………………………………………………………………

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COMPLIANCE WITH THIS DOCUMENT IS MANDATORY

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APPENDIX B: Complications following the application of compression

**Pain:** The application of compression bandages should not increase pain in the limb. If pain persists remove compression and recheck arterial status of the limb, also reassess for infection.

**Pressure Damage:**
Patients with impaired peripheral perfusion, thin or altered limb shape, foot deformity or dependent oedema are at increased risk of pressure damage. Other risk factors include reduced sensation, reduced pain sensation, long term systematic steroid use and presence of chronic disease associated with reduced mobility, loss of calf muscle and foot / ankle deformity.

- Avoid using sustained compression on these patients, consider inelastic systems or IPC
- Apply extra padding over bony prominences
- Ensure bandaging is not too tight and overlap is even. At risk areas include the ankle, the dorsum of the foot and the calf
- Observe for signs of pressure damage such as erythema, blistering or altered limb shape
- Encourage limb elevation for dependent oedema.

**Loss of Calf Muscle:**
Wastage of calf muscle can occur for patients receiving long term compression. This is usually directly not due to the compression but is often caused by reduced patient activity, underlying co-morbidities and medication.

- Ensure bandage allows good knee and ankle mobility. Ensure flat comfortable shoes are worn
- Encourage exercise and rehabilitation.

**Skin Problems:**
Maceration, excoriation, dryness, itching, allergic or irritant eczema and erosive pustular dermatosis are often associated with compression, topical preparations or chronic inflammation.

- Ensure adequate exudate control with appropriate primary dressings
- Use cotton liner or paste bandage against the skin
- Moisturise the skin with a simple emollient. Use downward movement in direction of the hair growth to avoid folliculitis
- Treat eczema
- Review all products use in treatment of the limb

**Allergy Alert:**
Some bandages / compression garments may contain latex remember to check this and do not use if patient allergic or sensitive to this.

**Bandage slippage:**
Reassess method of bandage application to ensure it has not been applied too loosely. Slippage can also occur if bandages have been applied correctly as the reduction of oedema and subsequent limb size may cause the bandages to slip.

**Swelling of the Toes or the area around the Knee:**
This may result from the bandage being too tight, too low from the knee, too far back from the toes, lack of exercise or sitting for long periods with legs down. This reduces the effectiveness of the pump action required for venous return and increases oedema in these areas.
Foot wear:
Some clients will not tolerate compression bandaging as they can’t wear their usual footwear. Therefore adjustable footwear to accommodate compression bandaging may need to be sourced.

Ineffective compression:
Reassess the client’s limb shape. Make certain there is enough padding and the bandage materials are appropriate. Ensure the primary dressing is not reducing the sub-bandage pressure. Check that the bandage is it being applied at the correct tension.

Tourniquet effect
This can occur at the top of the limb when compression bandaging is finished off incorrectly. At the completion of applying the compression layer any leftover bandage needs to be cut off or taped off so that the bandage is held in place without causing a tourniquet effect. This can also occur in limbs where a skin lobule over hangs the joint e.g. at base of leg over ankle joint. This skin fold needs to be padded out to be level with surrounding skin.

Tourniquet effect from stockings
Circular / round knit stockings may not be appropriate in patient with a skin lobule that over hangs the joint e.g. at base of leg over ankle joint, customise flat knit stockings may be required to level skin out.
Appendix C: Specific Compression

**Bandage Systems** giving 40mm Hg (regular compression) at the ankle include:
- Four Layer Bandage Systems
- Two Layer Bandage Systems
- Short stretch compression bandages
- High stretch compression bandages.

**Compression therapy** giving 40mm Hg at the ankle include:
- Intermittent Pneumatic Compression
- Compression wraps
- Compression Stockings 25mm Hg and above.

**Reduced compression bandage systems and alternatives**
- Four Layer Bandage Systems Lite
- Two Layer Bandage Systems Lite
- Short stretch compression bandages
- Intermittent Pneumatic Compression reduced intensity
- Three (3) Layer tubular bandaging e.g. TubiForm / Tubigrip
- Compression wraps applied at reduced intensity
- Compression stockings grade 1 or up to 25mm Hg.

Compression bandages should be applied as per the manufacturer’s instructions and in a manner which will achieve graduated compression.

**Compression achieved by static stiffness**
- Short stretch compression bandages however, may need multiple bandages to achieve desired effect.
Appendix D: Four Layer Bandage Systems e.g. Profore™ (Smith and Nephew) or Veno 4™ (Hartmann)

A four layer bandage system achieves 40mmHg at the ankle through application of a number of layers of low compression that together exert a cumulative effect. To achieve 40mmHg at the ankle all four layers must be applied correctly and to the correct ankle size.

The four layers include:
1. The padding bandage
2. Crepe or similar retention bandage
3. Light weight long stretch (elastic) bandage
4. Elasticised rubber bandage.

Apply in the following sequence for ankle size 18-25cms:
- Wound contact layer. Apply directly to the wound. If wound has a moderate to high exudate an alternative dressing might be required
- Padding bandage (layer 1). Apply from toes to knee with slight tension (to avoid puckering) using a spiral technique with 50% overlap. Ensure shin and ankle is adequately padded
- Light retention bandage (crepe or similar) (layer 2). Apply from toes to knee using spiral technique with 50% overlap
- Light compression bandage (layer 3). Apply from toes to knee using figure of eight technique with 50% extension of bandage. Use central yellow line as a guide to overlap. Secure with tape
- Flexible cohesive bandage (layer 4). Apply from toes to knee using a spiral technique with 50% extension and 50% overlap. This bandage will adhere to itself. The use of tubifast over the flexible cohesive bandage is acceptable if the client finds the cohesiveness uncomfortable.

A reduced compression can be achieved by omitting the light compression bandage (layer 3) or the cohesive bandage (layer 4). Application of only three (3) layers will approximately halve the level of compression.

For ankle size greater than 25cms the four layer bandage system will need to be modified as per table.

<table>
<thead>
<tr>
<th>18-25 cm Ankle Circumference</th>
<th>25-30 cm Ankle Circumference</th>
<th>&gt; 30 cm Ankle Circumference</th>
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</thead>
<tbody>
<tr>
<td>Padding bandage</td>
<td>Padding bandage</td>
<td>Padding bandage</td>
</tr>
<tr>
<td>Crepe or similar retention bandage</td>
<td>Crepe or similar retention bandage</td>
<td>Lightweight long stretch bandage</td>
</tr>
<tr>
<td>Light weight long stretch bandage</td>
<td>Long stretch bandage</td>
<td>Long stretch bandage</td>
</tr>
<tr>
<td>Elasticised cohesive bandage</td>
<td>Elasticised cohesive bandage</td>
<td>Elasticised cohesive bandage</td>
</tr>
</tbody>
</table>
Appendix E: Two Layer Compression System e.g. Coban™ (3M) or UrgoK2 (Link Medical)

Two Layer Compression System - UrgoK2
This system comprises two layers which cohere to form one thin conforming compression bandage which can be left in place for up to seven days. The system chosen is latex free and dependent on the ankle size (either ankle size 18-25cm or 25-32cm) and the amount of compression required e.g. regular or lite option.

- First Layer KTECH: white, short-stretch bandage, providing compression, protection and absorbency. Composition: wadding: viscose, polyester; knitted layer: polyamide, elastane
- Second layer: KPRESS: pink/beige, cohesive long-stretch bandage, providing additional compression necessary to achieve the therapeutic pressure and securing the bandages in place. Composition: cotton, polyester, polyamide, elastane; synthetic latex free cohesive material.

Application Method
Before applying the bandages:
- Examine the shape of the leg and identify any areas at risk of excessive pressure (i.e. bony prominences)
- Protect and reshape leg with wadding if necessary. If a wound is present, apply an appropriate dressing before applying any bandages
- Apply the compression system first thing in the morning or after the patient’s legs have been elevated for an hour to minimise any orthostatic oedema.

Ankle circumference 18-25cm kit – 50% overlap –
1. Place foot at a 90% angle – ‘toes to nose’. Start applying KTECH Lite at the base of the toes using two turns to anchor the bandage, ensuring wadding side is in contact with the skin and the pressure indicator is at the top edge, towards the patient. Secure the heel by using a figure of eight, ensuring full coverage of the heel (picture 1). Do not apply with pressure indicator at full stretch on the foot.
2. Spiral KTECH Lite up the leg from malleolus, stretching the bandage so that the pressure indicator (printed on the bandage) forms a circle, achieving the therapeutic pressure. A correct overlap is applied when the pressure indicator is just covered (50% overlap). Finish 2cm below popliteal space and cut off any excess bandage. Secure with tape.
3. Apply KPRESS (or KPRESS Latex Free) over KTECH Lite using the same application technique as KTECH Lite. For patient comfort, allow a small border of KTEC Lite at the toes and knee. Once applied, press down gently on bandage to ensure full cohesion.

Ankle circumference 25-32cm kit – 2/3 overlap
- Apply in the same way as the 18-25cm kit, stretching the bandage so that the pressure indicator forms a circle
- Cover the pressure indicator (printed in the middle of the bandage) to achieve the correct overlap (2/3 overlap).

Two Layer Compression System – Coban 2™ 3M
This system comprises two layers which cohere to form one thin conforming compression bandage which can be left in place for up to seven days.
Apply from just proximal to the toes to two fingers widths below the knee. The foot should be positioned at 90 degrees to the leg during application to avoid the bandage wrinkling during standing or walking. A figure of eight technique can be used to anchor the bandage to the foot.

Apply in the following sequence:
- First Layer (comfort layer): Is composed of foam laminated to a latex-free cohesive bandage and is wrapped upwards around the foot and leg with a minimal overlap
- Second Layer (Compression Layer): Is wrapped over the first layer with a 50% overlap using full stretch to provide effective sustained compression.

Two Layer Compression System – Coban 2 Lite™ (3M) 25% less resting pressure compared to standard - Combine 2.

Apply from just proximal to the toes to two fingers widths below the knee. The foot should be positioned at 90 degrees to the leg during application to avoid the bandage wrinkling during standing or walking. A figure of eight technique can be used to anchor the bandage to the foot.

Apply in the following sequence:
- First Layer (comfort layer): Is composed of foam laminated to a latex-free cohesive bandage and is wrapped upwards around the foot and leg with a minimal overlap
- Second Layer (Compression Layer): Is wrapped over the first layer with a 50% overlap using full stretch to provide effective sustained compression.

Appendix F: Short Stretch (inelastic) Compression Bandage e.g. Comprilan™ (BSN)
Short stretch bandages do not contain significant amounts of elastomer; rather they rely heavily on heavily twisted cotton yarns for their elastic properties.

Short stretch bandages exert low resting pressures – i.e. a low pressure is exerted whilst the patient is resting and high working pressures – i.e. a high pressure is exerted whilst the patient is walking and the calf muscle is pushing against the inelastic bandage.

Short stretch bandages might need to be reapplied frequently in patients with oedema as they do not have the ability to alter tension and will therefore become loose.

Short stretch bandages may be applied singularly, as directed and then if needed a second short stretch bandage may be applied if additional pressure is required.

Apply in the following sequence:
- Padding bandage
- Short stretch bandage: Apply from toes to knee using a spiral technique with 75 to 100% extension and 50% overlap*
- Tubifast if needed

* Apply second Short stretch bandage if required: Apply from toes to knee using a spiral technique with 100% extension and 50% overlap in the opposite direction to the first Short stretch bandage.
Appendix G: High Stretch (elastic) Compression Bandage e.g. Surepress™ (ConvaTec)
High stretch bandages contain elastomers and their length can increase significantly when stretched.

High stretch bandages exert a high resting pressure – i.e. a high pressure is exerted whilst the patient is resting and high working pressures – i.e. a high pressure is exerted when the patient is walking and the calf muscle is pushing against the bandage.

Apply in the following sequence:
- Padding bandage
- Long stretch bandage: Apply from toes to knee using spiral technique with 50-75% extension and 50% overlap.
  - Note Surepress is a guided compression bandage, the rectangles on the bandage show the correct amount of tension (pull) required to achieve the correct compression based on the ankle size.
- In-elastic Retention tubular bandage e.g. Tubifast™ Molnlycke, if needed.

Appendix H: Intermittent Pneumatic Compression e.g. Flowtron Hydroven (Arjohuntleigh)
- Intermittent Pneumatic Compression is generally tolerated well by most people. This is an external compression device consisting of an inflatable boot and machine. Consider if external providers will cover the cost of this for community clients e.g. Veterans Affairs will pay for this for their clients.
Appendix I: Compression Level / Scales

Compression Scale 1

<table>
<thead>
<tr>
<th>COMPRESSION LEVEL</th>
<th>INDICATIONS FOR USE</th>
</tr>
</thead>
<tbody>
<tr>
<td>RENTENTION BANDAGES</td>
<td>● Retention of wound product e.g. Crepe or similar bandages <strong>DO NOT</strong> provide sufficient compression to result in an increase in venous return.</td>
</tr>
</tbody>
</table>
| LIGHT COMPRESSION 14-17 mm Hg at ankle | ● Relief of leg discomfort associated with tired aching legs or mild varicose veins  
  ● People who spend long periods of time standing  
  ● Relief from leg discomfort during pregnancy |
| MILD COMPRESSION <20 mm Hg at ankle | ● Varicose veins  
  ● Prevention or treatment of mixed ulcers  
  ● Management of mild moderate oedema  
  ● Post surgery for leg veins or muscles weakened by surgery or lack of exercise |
| MODERATE COMPRESSION 20-40 mm Hg at ankle | ● Varicose veins  
  ● Prevention of venous ulcer recurrence  
  ● Management of moderate oedema |
| STRONG COMPRESSION 40-60 mm Hg | ● Venous ulcer treatment  
  ● Post thrombotic venous insufficiency  
  ● Severe chronic venous insufficiency  
  ● Severe varicose veins |
| VERY STRONG COMPRESSION >60 mm Hg | ● Lymphoedema |

Compression Scale 2: Compression Hosiery

<table>
<thead>
<tr>
<th>Pressure (mmHg)</th>
<th>British Class</th>
<th>European Class</th>
</tr>
</thead>
<tbody>
<tr>
<td>14-17</td>
<td>I</td>
<td>I</td>
</tr>
<tr>
<td>18-21</td>
<td></td>
<td>I</td>
</tr>
<tr>
<td>18-24</td>
<td>II</td>
<td></td>
</tr>
<tr>
<td>25-32</td>
<td>II</td>
<td>II</td>
</tr>
<tr>
<td>25-35</td>
<td>III</td>
<td>III</td>
</tr>
<tr>
<td>36-46</td>
<td></td>
<td>III</td>
</tr>
</tbody>
</table>

15-20 mmHg
● Mild ankle, foot and leg swelling  
● Leg fatigue  
● Pregnancy  
● Mild varicose veins  
● Spider veins  
● Tired, aching legs  
● Travel (usually greater than four hours).
20-30 mmHg
- Moderate to severe varicose veins
- Post sclerotherapy / vein stripping surgery
- Moderate venous disease
- Helps prevent recurrence of venous leg ulcers
- Prevention of post thrombotic syndrome
- Treatment of deep venous thrombosis.

30-40 mmHg
- Venous ulcer management and prevention
- Severe leg swelling e.g. post fracture or trauma
- Chronic venous insufficiency
- Severe varicose veins.

Notes
- Anti-embolic stockings are indicated for prevention of deep venous thrombosis whilst lying in bed. They do not provide sustained adequate compression whilst ambulating.
- A parallel support bandage will not achieve adequate compression at the ankle to enhance venous return and may cause a reverse pressure gradient e.g. TubiForm
- The level of compression is dependent on the type of garment chosen and application technique e.g. a long stretch bandage applied with 40% stretch will apply less compression pressure than one applied with 60% stretch.
- Specialised compression stockings are available in a range of compression pressure levels.
Appendix J: Compression stocking to prevent venous leg ulcers returning

What are venous leg ulcers?
Venous leg ulcers are caused by your veins not working properly to bring the blood from your legs back to your heart. This condition leads to increased swelling in your lower legs, which causes ulcers to form. To help your veins return your blood back to your heart and reduce this swelling, you must wear a compression stocking to prevent the ulcer returning.

Compression stockings need to be very firm at all times. Please observe the following advice:

COMPRESSSION STOCKINGS: Once the ulcer has healed using bandages, you will be required to wear a compression stocking every day to prevent an ulcer from reoccurring (coming back). Your doctor or nurse will advise if you need to wear compression stocking on one leg or both legs.

TO HELP YOU WEAR YOUR COMPRESSION STOCKING EACH DAY PLEASE:

- Shower of an evening immediately prior to going to bed (do not shower in the morning)
- Massage moisturiser (e.g. Sorbolene cream) into skin of legs (after shower)
- Sleep with legs elevated (raise foot end of bed slightly)
- Put compression stockings on before putting feet to floor in the morning (to prevent swelling). Note: keeping stocking by bedside may help with this
- Cover any open wounds (sores or ulcers) before putting compression stocking on
- If due to hot weather the compression stockings become unbearable to wear you may remove them **BUT DO NOT** walk around whilst compression stockings are off, as your legs will immediately swell making re-application difficult. **Rest with your ankles higher than your hips and move feet back and forwards to improve circulation!**

STOCKINGS MUST BE REPLACED EVERY SIX MONTHS and as per Manufacturer’s instructions!!!
Fitting: Compression stockings can be very hard to put on. When you buy your stocking ask if there is something to help you ‘put on’ and ‘take off’ your compression stocking.

When applying compression stocking:

- Protect the stocking from jewellery and fingernails – by wearing cotton or rubber gloves
- When pulling up stocking do not over stretch the stocking. The stocking should start at the toes and stop just below the knee cap. (do not fold or roll the top of the stocking over as this can stop the blood flow in the leg)
- Always wear stocking as the instructions say.

Care of Compression Stocking:

- Do not use Vitamin E or petroleum based moisturisers
- Wash stocking by hand or gentle machine wash daily
- Use a mild laundry detergent
- Do not dry in direct sun – DO NOT use a clothes dryer
- Do not soak garments or use bleach
- Always lie stocking flat to dry (hanging may stretch stocking).

Please remove the compression stocking and contact your GP, Community Health Nurse or hospital emergency department if you are concerned or notice any of the following:

- Increasing pain in toes, foot or leg
- Blue discolouration of the toes
- Numbness, coldness or swelling of the toes, foot or leg
- Staining from a wound coming through the compression stocking.
Appendix K: 3 Layer tubular bandaging system

How do I apply 3 layer compression bandaging with Tubular Form™ tubular compression bandage?

Why would I use 3 layers of Tubular Form™?

- Weller et al. Wound Practice & Research 2010 shows that 3 layers of Tubular Form straight is useful to inform future compression bandages studies that plan to measure venous ulcer healing rates.
- Bale S, Harding KG. Br J Nurse. 2003 shows that tubular bandages are useful for managing patients who cannot tolerate therapeutic forms of compression.

How do I measure for the correct size Tubular Form™?

- Always measure smallest circumference (ankle) to select the correct Tubular Form™ bandage size and refer to table below.
- Each bandage length is determined from measuring along back of the toes around the heel to desired length along the leg (Long, Medium, Short).

How do I apply?

- Apply Long 1st, Medium 2nd and Short 3rd (refer diagrams).

What if the patient is unable to tolerate 3 Layers?

- Simply remove Layer 3 (short) first, then if patient is still unable to tolerate remove Layer 2 (medium).

What if I am unsure of the patient’s vascular condition?

- Always consult physician if unsure of patient’s vascular condition.
- This method of compression therapy is recommended for Venous Insufficiency.

Tubular Form™ Measuring guide (ankle circumference):

<table>
<thead>
<tr>
<th>Size</th>
<th>Measurement</th>
</tr>
</thead>
<tbody>
<tr>
<td>C</td>
<td>15 - 23cm</td>
</tr>
<tr>
<td>D</td>
<td>26 - 35cm</td>
</tr>
<tr>
<td>E</td>
<td>36 - 45cm</td>
</tr>
<tr>
<td>F</td>
<td>46 - 55cm</td>
</tr>
<tr>
<td>G</td>
<td>56 - 65cm</td>
</tr>
</tbody>
</table>

Layer 1: From base of toes to back of knee.
Layer 2: From base of toes to mid calf.
Layer 3: From base of toes to mid point between mid calf and ankle.