Thonic bandage: A new application technique as a first step towards controlled compression? Reprocomp study

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Introduction

Compression bandages are one of the corner stones of veno-lymphatic conditions management and all international publications agree on the fact that bandages have to be applied by trained practitioners.1

This is clearly understandable because the level of compression reached or, in other words, the dosage of the treatment, is the result of a complex equation (Laplace’s Law) involving many factors: \( P = \frac{T \times N \times 4630}{C \times W} \) where \( P \) = sub-bandage pressure (mmHg), \( T \) = bandage tension, \( N \) = number of layers, \( C \) = limb circumference (cm) and \( W \) = bandage width (cm).1

It is important to notice that in this equation, 2 of the factors are operator dependant: the tension with which the bandage is applied and the number of layers (overlap). This is why international compression experts agree to say that the compression is not in the bandage but in the hands that apply it which is in total contradiction with the concept of controlled compression. These 2 factors will be the focus of this study.

A German study2 showed that, in practice, primary healthcare professionals struggled to reach the recommended sub-bandage pressures between 50 and 60 mmHg. The range of pressures reached went from 6 to 143 mmHg and only 10% of them achieved the target.

Traditionally, the applicator creates the tension by stretching the bandage maintaining the roll at a distance from the limb. The resulting torque effect could explain part of these major discrepancies.

Thonic bandage’s patented system allows to mechanically limit the stretchability of the elastic component to 30%, which means it can be wound up under maximum tension before application. In this case, Thonic bandage can be applied by unwinding it in contact with the limb, which allows the application of a more regular and controlled tension.

Objectives of the study

The first objective of this study is to check whether stretching Thonic bandage before application allows reaching more reproducible pressures and to narrow the range of pressures achieved by an applicator. The second objective is to compare the sub-bandage pressures reached by different groups: people who have never applied a bandage before, patients, primary care professionals, and secondary care professionals.

Materials and Methods

The sub-bandage pressures achieved by 4 different bandaging systems (2 short-stretch + 2 long stretch) will be measured in this study.

Both short-stretch bandaging systems will be applied traditionally and previously stretched at maximum strength. Both long stretch-bandaging systems will be applied as recommended.

Sub-bandage pressures will be measured with a Pico-Press system. In order to avoid any bias due to the 2 non-operator dependant elements of Laplace’s law, the measurements will be made on a plastic leg and the sensor will remain in exactly the same position for all measurements. For this reason, the compression levels reached by the different systems will not be of any clinical value. All bandages will also have the same width.

Each applicator will repeat 3 times and randomly the following applications: i) thonic bandage: applied traditionally; ii) thonic bandage: applied previously stretched; iii) short stretch bandage: applied traditionally; iv) short stretch bandage: applied previously stretched; v) long stretch bandage A: applied traditionally; vi) long stretch bandage B: applied traditionally.

Both long stretch bandages allow the delivery of compression force 1 (20 to 45 cN/cm²) and force 2 (46 to 100 cN/cm²) with 2 different techniques: increase of stretch for one and increase of overlap for the other.

The average pressure reached in 3 applications and the range (maximum pressure - minimum pressure) will be calculated. Obviously, the lower the range, the most controlled the compression will be.

The participants will be split between 4 categories: i) group 1: people who have never applied a bandage before; ii) group 2: patients; iii) group 3: primary care professionals (nurses, physios, …); iv) group 4: secondary care professionals (nurses, physios, …).

References

3. IFTH measurements, norm NF G 30 104-1.