Tissue fluid pressure and flow in the subcutaneous tissue in lymphedema – hints for manual and pneumatic compression therapy

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ABSTRACT

Physiotherapy of lymphedema requires knowledge of: a) how high external pressures should be applied manually or set in compression devices in order to generate tissue pressures high enough to move the fluid to the non-swollen regions and b) how to measure the tissue fluid flow.

We measured tissue fluid pressure and flow under the skin in the subcutaneous tissue of lymphedematous limbs stage II to IV at rest and during manual and pneumatic compression under various pressures and sleeve inflation timing.

In obstructive lymphedema of lower limbs tissue fluid pressures in the subcutaneous tissue was 2.5±3.0 mmHg (range -1 to +10 mmHg) and did not differ from those measured in normal subjects. During manual massage the applied force generated pressures ranging from 60 to 120 mmHg. Pneumatic compression generated tissue fluid pressures depending on sleeve inflation pressures, however, they were on the average 20% lower than in the inflated chambers. The high pressure gradient across skin and subcutis could be explained by skin rigidity (fibrosis), low hydraulic conductivity of subcutis and dissipation of the applied force in subcutis to the proximal non-compressed regions. Strain gauge put around the limb provided data on girth changes during compression and allowed to calculate the approximate volume of the proximally displaced fluid. It showed that tissue fluid flow occurred during manual compression only during pressing of tissues to stop immediately after its cessation. In contrast, pneumatic sequential compression produced unidirectional flow toward groin without backflow. The total proximally displaced volume from ankle to groin was up to 100ml/cycle.

The obtained data should be useful for physiotherapy allowing to set the manual or pneumatic compression parameters at levels corresponding to the physiological conditions.
INTRODUCTION

Treatment of lymphedema of limbs is a combined modality comprising administration of medicines, compression procedures based on manual and pneumatic massage, wearing elastic stockings or bandages preventing accumulation of fluid and growth of expanded skin and subcutis, also enhancing the efficiency of muscular pump, and in selected cases lymphovenous shunts, liposuction and debulking of abundant tissues.

The mechanism of action of administered anti-lymphedema medicines as diosmins and related drugs as well as indications for surgical procedures have been well defined and the results of treatment can be quantified. In contrast, compression procedures bring about diverse effects because of differences in manual massage protocols, pneumatic devices and poor knowledge what pressures and compression timing should be used. There is no information in the literature on mobile tissue fluid pressure and flow in human skin and subcutaneous tissue under normal conditions and in various types of lymphedema. We previously measured lymph pressures and flow in human calf collectors, however, the obtained data did not give insight into the hydraulic conditions in the interstitium. In lymphedema most lymphatic collectors do not contract as they have a fibrotic wall and are partially obliterated. The bulk of the capillary filtrate accumulates in tissue spaces.

Compression therapy partly supports, partly replaces the force necessary for propelling tissue fluid toward the root of the limb. Unfortunately, this treatment modality has so far not been based on the knowledge of hydraulic conditions in the interstitium of skin and subcutaneous tissue and blood perivascular spaces. The technical parameters of pneumatic compression devices are based on blood rheological data derived from textbooks. These are the capillary and venous blood pressure, venous blood flow and capacitance and capillary filtration. Unfortunately, there is little analogy between the blood and lymphatic system rheology at the tissue level. Pressures applied by pneumatic compression are set arbitrarily, because of lack of knowledge of skin and subcutis compliance (fibrosis), exact sites of lymphatic obstruction and accumulation of excess of tissue fluid as well as tissue fluid pressures and flow. The recommended compression pressures for pneumatic massage are around 50mmHg, timing of sleeve chamber inflation is usually 5 seconds and it is followed by a rapid deflation. Total compression time by an 8-chamber sleeve usually ranges between 20 to 40 seconds. It is not known whether this timing is sufficient to overcome tissue hydraulic resistance and move the stagnant tissue fluid.

What needs to be elaborated for clinical practice is how much external pressures should be applied either manually or by compression devices in order generate tissue pressures high enough to move the fluid to the non-swollen regions and how to measure the tissue fluid flow.

In this study, we measured tissue fluid pressure and flow under the skin in the subcutaneous tissue of lymphedematous limbs stage II to IV at rest and during manual and pneumatic compression under various applied pressures. The obtained data should be useful for physiotherapy and allow to set the compression parameters at levels corresponding to the physiological conditions.

MATERIAL AND METHODS

Patients.

Study was carried out on 25 patients with lymphedema of one lower limb, stage II to IV, duration of 2 to 15 years. Edema developed month to years after small foot abrasion, insect bite or soft tissue trauma. In 50% of cases edema was complicated by 1 to 3 attacks of dermato-lymphangio-adenitis. In 3 patients edema developed without any detectable reason. Cases with acute inflammation were excluded from the study. Five male volunteers with healthy legs served as controls. Staging was based on the level of edema starting from foot to groin, degree of skin keratosis and fibrosis and soft tissue tonometry. Evaluation of lymphatic pathways was done on lymphoscintigraphic pictures. Color Doppler investigation was also carried out to exclude cases with the postthrombophlebitic changes. Calf and thigh circumference were measured in supine position 15cm below and 15 cm above the lower edge of patella. The study was approved by the Warsaw Medical University and the Indian Universities ethics committees.

Tissue fluid pressure measurement

The wick-in-needle technique was used. An 8 gauge injection needle with a polyethylene tubing (OD 1.34
mm) containing glass-wool wick protruding from the tubing tip at 5mm was introduced under the skin. The outer part of tubing was led outside the compression sleeve, connected to the pressure transducer (Honeywell, Elblinger, Poland) and recording was done using a 3 channel device, pressure range -20 to 150 mmHg (Telsoft, Warsaw, Poland) and LabView software (National Instruments, Austin, TX, USA). The position of transducer was zeroed and pressure recording was started simultaneously with manual compression or sequential inflation of sleeve chambers. The data were collected using Microsoft Excel program and were presented graphically on a pressure/time scale.

**Continuous limb girth (volume) measurement**

Strain gauge plethysmography was used to measure girth changes in the calf and thigh. The plethysmograph (Hokanson, Bellevue, WA, type EC6) recording vein mode was applied. The girths of mid-calf and mid-thigh were measured and a 2 cm shorter mercury strain gauges were put around limb at chamber levels 3 to 8. Elongation was read off on the recorder graph scale in mm as a change in limb circumference. Inflation of chambers located distally to the strain gauge propelled mobile tissue fluid in proximal direction. Once fluid reached the limb region with strain gauge, its volume increased and so did the girth. Increase in girth was recalculated into volume by multiplying cross area of limb at the studied level by the length of the compressing chamber. Subtracting the volume value before compression from that during compression provided data on the transferred volume.

**Manual massage**

Limb massaging started distally from the level of pressure sensor and strain gauge, slowly approached them, then it was continued over the site of recording in proximal direction to the groin. Tissue fluid pressure and girth changes were recorded continuously.

**Pneumatic compression appliance**

We used such a device designed for us by Biocompression (Moonachie, NJ). It was built of 8 segments 9 cm long each, sequentially inflated, inflation pressures were regulated from 50 to 125 mmHg, gradient pressures was proximally decreasing by 20%, inflation time of each chamber was 50 sec, there was no deflation of distal chambers, the deflation time at the end of the cycle was 50 sec. The long inflation time was based on our observations that manual squeezing of tissue fluid from the wound during debulking surgery took a minimum of 50 to 100 sec. No deflation prevented venous return to the distal parts of the limb and venous stasis with increased filtration rate. Sites of pressure and flow recordings have been shown on Figure 1.

**RESULTS**

**Subcutaneous tissue fluid pressures at rest**

Tissue fluid pressure measured under the skin in lymphedematous calfs ranged between – 1.5 and 10 mmHg (mean 2.5±3.0) and in controls between -1.8 and

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**Figure 1.** Schematic presentation of lower limb in a pneumatic sleeve with 8 chambers 9 cm wide each. Tissue fluid pressure was measured at 6 points indicated by large dots. The lines encircling calf and thigh show the site of strain gauge location for continuous measuring of girth changes.

**Figure 2.** Tissue fluid pressure measured under the calf skin in a horizontal position in 15 patients with lymphedema stage II to IV and 5 healthy subjects. No significant differences between lymphedema and normal limbs. There were no differences depending on the stage of the disease. Data are means +SD.
Tissue fluid pressure and flow in lymphedema

3.0mmHg (mean 0.8±1.2) (Figure 2). There were no statistically significant differences between the groups. These readings corroborate our previous clinical and experimental findings.10, 11 The mechanism of low pressure can be explained by high skin compliance in the early stages of edema formation leading to expansion of the subcutaneous space absorbing excess of capillary filtrate not drained away by lymphatics.

Subcutaneous tissue fluid pressures and flow during manual massage

The manual force for massage is set arbitrarily and never seems to be too high. However, our measurements showed that hands generated tissue fluid pressures ranging as high as 140 mmHg (Figure 3). This was observed in stages II and III. In stage IV high skin rigidity limited the force transfer to subcutis and lower pressures were observed. To investigate whether the manual force is transferred to proximal regions of the limb, we measured pressures at different distances from the massaging hand. Compression 3 cm from the sensor did not show any transfer pressure through this distance. This could be explained by skin rigidity and low hydraulic conductivity of subcutis.

Tissue fluid flow during manual massage was seen only during compression of tissues, to stop immediately after removal of the pressing hand (Figure 4). The calculated flow per compression values were around 5 ml. The skin indentation by the massaging hand usually disappeared within 10-15 sec. This indicated that the displaced fluid returned to its site of origin.

Subcutaneous tissue pressures and flow during pneumatic massage

The pressures generated in tissue fluid during the first inflation of a sleeve chamber were in all cases lower than those in the chamber itself. (Figure 5, 6A, 6B). This high gradient was most likely caused by skin rigidity (fibrosis) and dissipation of the applied force to the proximal non-compressed regions. Interestingly, there was also little tissue fluid pressure transmission in subcutis from the compressed to the non-compressed proximal segments for a distance of 9 cm (width of the chamber). In advanced stages IV limited pressure transmission in proximal direction could be seen. An interesting finding was building up pressure in the distal parts of the limb during sequential inflations of proximal chambers. This could be due to flow obstruction at the inguinal level. We also noticed that tissue fluid pressures reached lower levels in the popliteal and upper thigh than in other limb regions. These two regions are usually less swollen and
Figure 5. Tissue fluid pressure recording in a normal calf subcutaneous tissue at the level of compression sleeve chambers 3, 4 and 5. Inflation pressures 50, 80 and 120 mmHg. Inflation time of each chamber 55 sec, no deflation. Number 1 to 8 denote consecutive sleeve chambers. Pressures recorded under chambers 3 (yellow line), 4 (red line) and 5 (blue line). Note that pressures were always lower than those in chambers. Inflation of chambers 1, 2 and 3 to 50 mmHg did not generate pressures at level 3. Inflation of chamber 4 produced pressure of 30 mmHg at level 3 stepwise rising during inflation of consecutive chambers to 45 mmHg. Inflation of chamber 4 also produced tissue fluid pressure rise to 40 mmHg and of chamber 5 to 12 mmHg, to increase during inflation of consecutive chambers to 45 and 38, respectively. Similar pressure curves were observed during inflation of chambers to 80 and 120 mmHg. Blue line (level 5) represents pressure at the medial aspect of calf just below the knee usually with less edema.

Figure 6a. Tissue fluid pressure in the calf lymphedematous subcutis stage III during pneumatic compression of 50 mmHg (inflation time of each chamber 55 sec, no deflation). Pressures recorded at level 3 (blue line), 4 (red line) and 5 (yellow line). Note that tissue fluid pressures during first inflation of chambers at level 3, 4 and 5 were lower than those in the chambers themselves. Inflation of chambers 1 and 2 to 50 mmHg did not generate pressures at level 3. Inflation of chamber 3 produced at level 3 pressure of 25 mmHg (blue), stepwise rising during inflation of consecutive chambers to 55 mmHg. Inflation of chamber 4 produced tissue fluid pressure rise at level 3 and 4 to 45 mmHg and at level 7 to 30 mmHg to increase during inflation of consecutive chambers to 55 and 35 mmHg. The yellow line represents pressure at level 8 close to the groin usually with less edema and low flow resistance.

Figure 6b. Tissue fluid pressure in the thigh lymphedematous subcutis stage IV during pneumatic compression of 50 mmHg. Pressures recorded at level 6 (blue line), 7 (red line) and 8 (yellow line). Note that as in the calf, tissue fluid pressures during first inflation of all chambers were lower than those in the chambers. Inflation of chambers 1 to 5 to 50 mmHg did not generate pressures at level 6. Inflation of chamber 6 produced at level 6 pressure of 35 mmHg (blue), stepwise rising during inflation of consecutive chambers to 55 mmHg. Inflation of chamber 7 produced tissue fluid pressure rise at level 6 to 50 mmHg and at level 7 to 30 mmHg to increase during inflation of consecutive chambers to 55 and 35 mmHg. The yellow line represents pressure at level 8 close to the groin usually with less edema and low flow resistance.

Figure 7. Summarized data of tissue fluid (TF) pressures in calf and thigh subcutaneous tissue in patients with lymphedema (n=15) and healthy subjects (n=5) during the first inflation of pneumatic sleeve to 50, 80 and 120 mmHg. L- lymphedema, N- normal, horizontal line denotes pressure in the sleeve, mean values ± SD. Note that TF pressure was in all cases lower than in the sleeve both in the lymphedema cases and healthy subjects (for explanation see text). There were also evident differences between lymphedema and control cases.
slowly accumulate fluid translocated during sequential massage. Summarized data of 15 patients have been presented on Figure 7.

**Circumference (volume) changes during pneumatic massage**

Continuous recording of circumference changes during sequential compression gave indirect insight into the volumes of fluid translocated from the compressed segments to the proximal ones. Following inflation of a chamber, increase of limb circumference occurred proximally to this chamber. Sequential inflations of chambers from 1 to 8 resulted in stepwise increase of circumference in consecutive segments of the limb (Figure 6). The increase in circumference at each level was recalculated into increase in volume. Summarized data of 15 patients are presented on Figure 7. The calculated proximally transferred volume was evident in the calf but much more in the thigh containing large volumes of fluid and fat.

Continuous recording of circumference changes during sequential compression gave insight into the volumes of fluid translocated from one pressed segment to another. Following inflation of a chamber there was increase of circumference in limb segment above this chamber. When the segment with strain gauge was compressed, the girth decreased. Further sequential inflation of proximal chambers brought about girth decrease (Figure 8). This was observed in cases with low resistance to tissue fluid flow. In case of flow obstruction, girth rose to high values. The increase in girth was recalculated into increase in volume. The total volume accumulating fluid volume in the proximal parts of the limb could also be calculated. Summarized data have been presented on Figure 9.

**Correlation between pressure and girth changes during sequential compression**

Correlating pressure and flow data provides information of whether there is a high resistance to tissue fluid flow. High fluid pressures and low flow would suggest major flow obstruction, usually at the groin level, whereas, low pressures and low flows would point to low efficacy of compression due to high skin rigidity. High flows and low pressures were usually seen in early stages of lymphedema without fibrotic changes of skin and subcutaneous tissue. There was no correlation between tissue fluid pressure generated by external compression and flow in proximal direction in tissue fibrosis and scars after surgery and radiotherapy. There could be high pressure but practically no flow. This resistance could only be overcome by strong external compression generating very high tissue fluid pressures.
DISCUSSION AND CONCLUSIONS

Taken together, in obstructive lymphedema of lower limbs tissue fluid pressures in the subcutaneous tissue were low and did not differ from those measured in normal subjects. During manual massage the applied force is set arbitrarily and it generated pressures ranging from 100 to 150 mmHg. Compression at a distance of 3 cm from location of the pressure sensor didn’t not show any rise in pressure what could be explained by skin rigidity and low hydraulic conductivity of subcutis. Tissue fluid flow appeared during manual compression only when the hand was pressing, but stopped immediately after cessation of massage. The calculated flow values during manual massage were low.

Pressures generated in tissue fluid by pneumatic compression chambers were lower than those measured in the inflated chamber. The gradient depended most likely on skin rigidity (fibrosis) and dissipation of the applied force in the subcutaneous tissue to the proximal non-compressed regions. There was little tissue fluid pressure transmission in subcutis to the noncompressed proximal segments. Interestingly, building up pressure in the distal parts of the limb during inflation of proximal chambers was observed. This was presumably the consequence of flow obstruction at the inguinal level. Strain gauge put around the limb provided data on circumference changes during compression and allowed to calculate the approximate volume of displaced fluid. This volume ranged in our studies from 10 to 30 ml per inflated chamber, to reach above 100 per the whole sleeve in the groin region.

Interestingly, there was direct correlation between tissue fluid pressure generated by external compression and flow in proximal direction. This was presumably caused by fibrosis of the subcutaneous tissue and scars after radiotherapy creating mechanical resistance to flow. It could only be overcome by external compression generating very high tissue fluid pressures.

REFERENCES


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