Title: Ultrasound Palpation Sensor for Tissue Thickness and Elasticity Measurement - Assessment of Transverse Carpal Ligament

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Abstract: Palpation is a traditional diagnostic procedure for health care professionals to use their fingers to touch and feel the body soft tissues. It is a common clinical approach, though it is rather subjective and qualitative and the palpation results may vary among different people. Tissue ultrasound palpation sensor (TUPS) provides a feasible solution that makes the palpation of soft tissues not subjective feeling any more. It is comprised of an ultrasound transducer together with a load cell to form the finger-sized probe. The probe is used to push against the soft tissue surface to measure the thickness and elasticity of the soft tissues. TUPS has been successfully applied to the assessment of various human tissues. Recently, we have improved TUPS, which can now be linked to personal computer (PC) via universal serial bus (USB) and provide a better user-interface. The information of ultrasound signal and indentation force is displayed on PC in real time during measurement. In this paper, we introduce the recent application of TUPS for the assessment of the transverse carpal ligament. The tissues at the carpal tunnel regions of five normal male subjects were tested using TUPS. The results showed that the average thickness of the tissues covering the carpal tunnel ligament and the tunnel region was 7.98 ± 1.05mm and 9.59 ± 1.12mm, respectively. Under a compression force of 20N applied by a cylindrical ultrasound indenter with a diameter of 9mm, the stiffness of the soft tissue layer and the tunnel region was 6.72 ± 2.10 N/mm and 15.63 ± 8.42 N/mm, respectively. It is expected that TUPS can be a potential tool for non-invasive assessment of carpal tunnel syndrome.

Keywords: Palpation, ultrasound, ultrasound palpation, tissue elasticity, soft tissue, carpal tunnel syndrome

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1. Introduction

Carpal tunnel syndrome (CTS) is a common musculoskeletal disease caused by the compression to the median nerve in the carpal tunnel. As median nerve is a mixed motor and sensory peripheral nerve, compression of it may cause losing of sensation, clumsy and even powerless of the palm sides from the medial half of thumb to the lateral half of the ring finger [1]. There are several clinical tests for CTS to check the sensory and mobility of the hand including provocative tests, Phalen’s test and Tinel percussion test [1]. However, the appearance of these clinical symptoms of CTS means that the patient may suffer from CTS severely. If we can develop a non-invasive approach to test the thickness and elasticity at the carpal tunnel region, it might be feasible to be a tool for diagnosing of CTS in the early states. So, health care professionals may apply non-surgical management such as anti-inflammatory drugs injection or hand support for neutral position to release the carpal tunnel pressure [1, 2]. It is because surgical management, which cuts the transverse carpal ligament (TCL), can help to release the carpal tunnel pressure (CTP) but it may cause the weakness of finger flexion [3].

TCL forms the volar aspect of the carpal tunnel at the wrist. The increase of CTP may be due to fluid retention, infection and excessive use of the fingers, which may cause swelling of the tendons or their synovial sheaths [4]. In pervious studies, TCL was found to be thickened with the increase of the CTP and it is relevance to CTS [5, 6]. In this study, the newly developed version of ultrasound palpation sensor (TUPS) was used to examine the thickness and the stiffness of the carpal tunnel in-vivo.

TUPS has been used to determine different kinds of soft tissues including residual limb tissues [7], burn and surgical scars [8], fibrotic tissues induced by radiotherapy [9]
and plantar foot tissues with diabetes [10]. The studies mentioned above considered the soft tissue and measured the tissue thickness and elasticity in whole. In this CTS study, an improved version of TUPS and its software were used in order to consider the soft tissues in the wrist region into two layers which are the soft tissues superior and interior to the TCL. Therefore, we are able to estimate the effects of the TCL to CTP.

2. Methods

TUPS is an ultrasound indentation with a finger size probe which consists of 5MHz ultrasound transducer with a diameter of 9mm and an in series load cell [11]. Fig 1 shows the block diagram of the TUPS system. Ultrasound was emitted from the ultrasound transducer to measure the thickness and deformation of different layers in the wrist region during indentation using the information of the time-of-flight and the sound speed. The average sound speed in soft tissues of human body was assumed to be 1540 m/s [12]. The load cell was used to record the indentation force. In this study, 20N force was applied to the subject’s palm via the measuring probe within 3 second.

Fig 2 shows the user interface of the custom-developed program for the TUPS system. It can show the ultrasound and force signals during indentation in real time. The ultrasound signal can also be shown in M-mode to present the overall displacement profile of the ultrasound echoes reflected or scattered from the tissues at different depths. As we introduced in earlier papers [9, 11], signal peak or cross-correlation tracking approaches could be used to track the shift of a selected echo, which corresponded to a tissue interface, such as tunnel-bone interface. However, these methods could not be well used for the ultrasound signals collected in this study. Since the applied load was
very large and tissues were deformed significantly, the tracking using signal peak or cross-correlation for a segment of signals did not work very well for the interfaces of TCL due to the obvious change of the interface echoes (de-correlation). In this study, we used the M-mode image to trace the shift of echoes quasi-automatically by selecting a number of critical points in the M-mode image for each interface echo under different loading levels. As an example, two groups of manually selected points were shown in the M-mode image of Fig 2. The software then automatically links the points together to form a deformation profile under different loads using a linear interpolation. Normally, at least 10 points were manually selected for tracking of each interested interface echo. The obtained deformation-time data were then further analyzed together with load-time data to obtain the stiffness information.

Five normal male subjects were recruited in this study with an average age of 29.8 ± 5.1 years old. They have no neuro-musculoskeletal disorders in their upper limbs. The location for the indentation of the palm is on the skin overlying the TCL. A line was firstly drawn to connect the palpable pisiform and scaphoid and a point 10mm distal from the mid-point of this line was marked as the centre of indentation. During the testing, the hand was supinated on a testing table with the palm side facing upward. 10 trails were recorded for each subject.

3. Results

The ultrasound echoes reflected from the carpal bone surfaces were obviously identified (Fig 2 and Fig 3). The original thickness of the tissues between the skin and the TCL and between the TCL and bone surface were 7.98 ± 1.05mm and 9.59 ± 1.12mm,
respectively, for the 5 subjects. We applied about 5N to compress the soft tissue between
the skin surface and TCL, it is because the stiffness of that layer was assumed to be softer.
In the M-mode display of ultrasound signals as shown in Fig 2, we can observe that the
level of deformation is different between the layer of skin-TCL and TCL-Carpal bone.
Figs 3a, 3b and 3c show the ultrasound echo trains and different tissue interfaces obtained
with the applied load of 0 N, 5 N and 20 N, respectively. In Table 1, the total
dehformations in skin-TCL layer and TCL-carpal bone layer are 3.18 ± 0.75mm and 1.16
± 0.42mm, which are 38.1% and 13.7% respectively when the indentation force is up to
20N. When the indentation is less than 5N, the deformation of skin-TCL layer is 2.2mm
± 0.8mm as the deformation of TCL-carpal bone layer is 0.65mm ± 0.25mm only, which
is 4 times larger. When the indentation force is between 5N to 20N, the deformation of
skin-TCL layer is 0.98 ± 0.51mm as the deformation of TCL-carpal bone layer is 0.50 ±
0.20 mm, which is 2 times larger. Therefore, the stiffness of TCL-carpal bone layer is
much larger than the skin-TCL layer and it is deformation dependent.

The stiffness of soft tissues and carpal tunnel layer under different indentation load
were shown in Table 2. We found that the difference of stiffness between two layers
keeps reducing when the indentation force increases. The overall stiffness of skin-TCL
layer and TCL-carpal bone layer are 6.72 ± 2.10 N/mm and 15.63 ± 8.42 N/mm,
respectively.

4. Discussion

In this study, we developed the TUPS system together with the program to perform
the indentation on the wrist region. The new developments of TUPS system and its
program not only provide the real time information of ultrasound and force signal to the
operator during indentation, but also provide a feasible solution to distinguish different
tissue layers rather than overall information of tissue thickness and elasticity. By the help
of M-mode display of ultrasound in the user-interface of the program, we tried to
distinguish TCL in the wrist region so as to study the deformation and the elasticity of
skin-TCL (soft tissues) layer and TCL-carpal bone (Carpal Tunnel) layer. The results
from Table 2 show that the stiffness of skin-TCL layer is 2.60 times less than TCL-carpal
bone layer when the indentation force is below 5N. The results proved the assumption
that the stiffness of the skin-TCL layer should be less than that of TCL-carpal bone layer.

However, when the indentation force is larger than 15N, the stiffness of skin-TCL
and TCL-carpal bone layer become similar (1.15 times larger), so the deformation of the
whole thickness is contributed relatively closer by both two layers. Therefore, we may try
to study the stiffness and deformation of TCL-carpal bone layer during the indentation
when the force is between 5N to 20N. The stiffness of TCL-carpal bone layer in this
study is $21.76 \pm 10.00$N/mm.

Since a relative large load was required to deform the carpal tunnel, the profile of
the ultrasound echo from a certain interface could significantly change, particularly that
from TCL. Automatic tracking methods using signal peak or cross-correlation that we
earlier developed could not work well in this study. We developed a quasi-automatic
measurement method to trace the shift of echoes by manually selecting critical points in
the M-mode display of ultrasound images. The manual selection of the points involved
subjective judgements. Therefore, inter- and intra-operator variations of the measurement
should be further documented in future studies. In addition, these manual operation also
limited the option to obtain the tissue deformation data in real-time during the test. We plan to develop more robust automatic tracking algorithms to precisely tracking the echo movement of tissue interface using the ultrasound signals. The ultrasound signals were digitized in 100 MHz, which corresponds to a displacement measurement resolution of approximately 8 µm if the echo did change its profile during the compression and could be automatically tracked, such as the echo reflected from the bone surface. However, manual selection for the points in this study could significantly affect the measurement accuracy. Further study will be required to systematically investigate the measurement accuracy under various conditions of the echo.

Another limitation of the reported method is that the measured stiffness value could be affected by the boundary condition of the carpal tunnel region. When a certain load was applied at the carpal tunnel region, the deformations of different tissue components might not only be controlled by the tissue elasticity, but also affected by the surrounding boundary conditions. Therefore, the stiffness value measured from different subjects could be affected by the dimension of the carpal tunnel region. The boundary conditions should be taken into account in the extraction of the stiffness from the load-deformation data in the future studies.

In spite of the above limitations, this study demonstrate the feasibility of the novel approach of using TUPS to measure the tissue thickness and elasticity of TCL which is very relevant to CTS. In future study, a higher frequency ultrasound (such as 20 MHz focused ultrasound) should be used to detect the TCL layer. It is because the resolution of the 5 MHz ultrasound may not be enough to distinguish the TCL easily. Afterward,
subjects with CTS should be recruited to perform the indentation to compare the difference of tissue thickness and stiffness between normal subjects and CTS patients.

Acknowledgements

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108.
Table 1 Deformation (mm) of the skin-TCL layer and TCL-carpal bone layer under different indentation load (5N and 20N).

<table>
<thead>
<tr>
<th>Force</th>
<th>Skin-TCL layer</th>
<th>TCL-carpal bone layer</th>
</tr>
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<tbody>
<tr>
<td>0 - 5N</td>
<td>2.20 ± 0.80</td>
<td>0.65 ± 0.25</td>
</tr>
<tr>
<td>5 - 20N</td>
<td>0.98 ± 0.51</td>
<td>0.50 ± 0.20</td>
</tr>
<tr>
<td>Overall</td>
<td>3.18 ± 0.75</td>
<td>1.16 ± 0.42</td>
</tr>
</tbody>
</table>

Table 2 Stiffness (N/mm) of the skin-TCL layer and TCL-carpal bone layer under different indentation load (5N and 20N).

<table>
<thead>
<tr>
<th>Force</th>
<th>Skin-TCL layer</th>
<th>TCL-carpal bone layer</th>
</tr>
</thead>
<tbody>
<tr>
<td>0 - 5N</td>
<td>3.51 ± 1.69</td>
<td>9.11 ± 5.83</td>
</tr>
<tr>
<td>5 - 20N</td>
<td>18.98 ± 7.73</td>
<td>21.76 ± 10.00</td>
</tr>
<tr>
<td>Overall</td>
<td>6.72 ± 2.10</td>
<td>15.63 ± 8.42</td>
</tr>
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</table>

Figure Captions

Fig. 1 Schematic of the TUPS system including the finger size probe together with other electronic parts as a control system of the load cell and ultrasound transducer. The control unit can be directly linked with personal computer via USB to have fast transmission of signals as real time measurement.
Fig. 2 User program which was used during indentation. In this interface, TUPS connects to PC via USB to transmit the ultrasound signal (the bottom window) and force signal (the top right window). The top left window was used in the signal processing for showing the displacement curve or global view for the whole tissue (M-mode). The points in the M-mode display were manually selected to represent the shifts of the echoes during the loading process.

Fig. 3 A sample result of ultrasound signals of a carpal tunnel. (a) Echoes obtained with no compression, (b) echoes obtained during indentation when about 5N was applied, (c) echoes obtained during indentation when totally 20N was applied.
Fig. 1

- load cell
- driver/amplifier
- A/D converter
- Ultrasound pulser/receiver
- high speed A/D converter
- USB
- PC system
Ultrasound signals

M-mode

Force

Ultrasound signals

Skin-interface   TCL     Bone-interface

Fig. 2
**Fig. 3**

Indenter-skin interface

Fat-TCL interface

TCL-tunnel interface

Tunnel-bone interface

(a) 0 N

(b) 5 N

(c) 20 N