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Puhakka PH

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Pia H PUHAKKA 1,2, Nikae C R te MOLLER 3, Petri TANSKA 1, Simo SAARAKKALA 1,4,5, Virpi TIITU 6, Rami K KORHONEN 1, Harold BROMMER 3, Tuomas VIRÉN 7, Jukka S JURVELIN 1, and Juha TÖYRÄS 1,2

1 Department of Applied Physics, University of Eastern Finland, Kuopio; 2 Department of Clinical Neuropysiology, Kuopio University Hospital, Kuopio, Finland; 3 Department of Equine Sciences, Utrecht University, Utrecht, the Netherlands; 4 Department of Medical Technology, Institute of Biomedicine, University of Oulu, Oulu; 5 Department of Diagnostic Radiology, Oulu University Hospital, Oulu; 6 School of Medicine, Institute of Biomedicine, Anatomy, University of Eastern Finland, Kuopio; 7 Cancer Center, Kuopio University Hospital, Kuopio, Finland. Correspondence: pia.puhakka@uef.fi

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Background and purpose — Arthroscopic estimation of articular cartilage thickness is important for scoring of lesion severity, and measurement of cartilage speed of sound (SOS)—a sensitive index of changes in cartilage composition. We investigated the accuracy of optical coherence tomography (OCT) in measurements of cartilage thickness and determined SOS by combining OCT thickness and ultrasound (US) time-of-flight (TOF) measurements.

Material and methods — Cartilage thickness measurements from OCT and microscopy images of 94 equine osteochondral samples were compared. Then, SOS in cartilage was determined using simultaneous OCT thickness and US TOF measurements. SOS was then compared with the compositional, structural, and mechanical properties of cartilage.

Results — Measurements of non-calcified cartilage thickness using OCT and microscopy were significantly correlated (ρ = 0.92; p < 0.001). With calcified cartilage included, the correlation was ρ = 0.85 (p < 0.001). The mean cartilage SOS (1,636 m/s) was in agreement with the literature. However, SOS and the other properties of cartilage lacked any statistically significant correlation.

Interpretation — OCT can give an accurate measurement of articular cartilage thickness. Although SOS measurements lacked accuracy in thin equine cartilage, the concept of SOS measurement using OCT appears promising.

Optical coherence tomography (OCT) is a high-resolution, non-destructive imaging method. The resolution can be less than 5 µm (Drexler et al. 1999). OCT helps, for example, in measurement of the thickness of structures in the eyes (van Velthoven et al. 2007), blood vessels (Jang et al. 2005), and teeth (Wilder-Smith et al. 2009). The potential of OCT in osteoarthritis research and clinical diagnostics is also noteworthy (Drexler et al. 2001, Li et al. 2005, Viren et al. 2012).

Arthroscopy is a common diagnostic technique in both human and equine medicine. Arthroscopic measurement of articular cartilage thickness using OCT could help in detecting cartilage thinning (Cernohorsky et al. 2012) or scoring of cartilage lesion severity (te Moller et al. 2013, Niemelä et al. 2014). Studies in rabbits and goats showed high correlations between OCT and microscopy measurements of articular cartilage thickness (Han et al. 2003, Rogowska et al. 2003, Cernohorsky et al. 2015). However, no similar study has been conducted with thicker cartilage, such as in humans or horses.

The speed of ultrasound (US) is lower in osteoarthritic cartilage than in healthy cartilage (Myers et al. 1995). This may be due to changes in water and proteoglycan content in the tissue and to collagen degeneration (Joiner et al. 2001, Suh et al. 2001, Töyräs et al. 2003). Consequently, reduced speed of sound (SOS) in articular cartilage could serve as an indicator of early and local cartilage degeneration. A technique that enables accurate arthroscopic measurement of local variations in SOS could have high diagnostic value.

We investigated the accuracy of OCT in the measurement of equine cartilage thickness, and used a novel arthroscopic technique utilizing OCT thickness measurement and US time-of-flight (TOF) measurement to determine the SOS in cartilage. We hypothesized that combining OCT and US would help to give accurate SOS determination. We assessed the applicability and accuracy of the technique with experimental measurements using phantoms and equine cartilage, and by numerically estimating the measurement uncertainty. We further compared SOS in equine cartilage with the structural, compositional, and biomechanical properties of the tissue.
phate-buffered saline (PBS) during the imaging process, with < 20 µm). The sample and the catheter were placed in phos-
Medical), providing cross-sectional images (axial resolution
with a thin catheter (0.9 mm diameter; C7 Dragonfl y; St. Jude
PCI Optimization System; St. Jude Medical, St. Paul, MN)
We used an OCT device (wavelength 1,305 ± 55 nm; Ilumien
Cartilage thickness measurement using OCT
al. 2010).

Material and methods
Sample preparation
18 metacarpophalangeal joints of 13 horses (age > 2 years)
were obtained from a slaughterhouse and frozen until sample
preparation. 4–7 osteochondral blocks including healthy carti-
lage and a variety of lesions were prepared from each joint
(94 blocks in total; 34 from the proximal phalanx (P1), 34
from the dorsoproximal area of the condyles of the third meta-
carpal bone (MC3), and the sagittal ridge of the third metacarpal bone
(SR).

Figure 1. Osteochondral samples were prepared from equine meta-
carpophalangeal joints at 5 an-
tomical locations: medial and lat-
eral proximal phalanx (P1), medial
and lateral dorsoproximal areas of
the condyles of the third metacar-
pal bone (MC3), and the sagittal
ridge of the third metacarpal bone
(SR).

Figure 2. The measurement setup. Adjacent optical coherence
tomography (OCT) and ultrasound (US) catheters were
inserted through an instrument
channel. The phantom mea-
surements were performed in
distilled water and the cartilage measurements in a bath of
phosphate-buffered saline.

Figure 2. The measurement
setup. Adjacent optical coherence
tomography (OCT) and ultrasound (US) catheters were
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surements were performed in
distilled water and the cartilage measurements in a bath of
phosphate-buffered saline.

Cartilage thickness measurement using OCT
We used an OCT device (wavelength 1,305 ± 55 nm; Ilumien
PCI Optimization System; St. Jude Medical, St. Paul, MN)
with a thin catheter (0.9 mm diameter; C7 Dragonfly; St. Jude
Medical), providing cross-sectional images (axial resolution
< 20 µm). The sample and the catheter were placed in phos-
phate-buffered saline (PBS) during the imaging process, with
the catheter held manually above the ink marking or lesion,
which were both visible in the OCT image. The thin low-
scattering layer observed in OCT images just above the sub-
choondral bone is assumed to correspond to calcified cartilage
(Cernohorsky et al. 2012). The thickness of the non-calcified
cartilage used in mechanical measurements was measured
from the site of interest using the OCT system software. After
matching the measurement points in the OCT and microscopy
images, the thicknesses of non-calcified cartilage, calcified
cartilage, and full cartilage were measured again from the raw
images, taking into account the refractive index of cartilage
(1.358; Wang et al. 2010). Pixel size was determined based on
the known diameter of the OCT catheter and considering the
refractive index of the water inside the catheter (1.322).

Speed of sound (SOS) measurements
First, the SOS in each phantom material was determined using
a reference method; a custom-made acoustic microscope (Fc
= 50 MHz, −6 dB bandwidth = 30–73 MHz, focal length = 25
mm, Fs = 550 MHz) was used to measure the TOF in the phan-
tom and a caliper (resolution = 10 µm; DIGI-MET 1226 417;
Helios-Preisser GmbH, Gammertingen, Germany) to measure
the thickness.

The SOS in each phantom was subsequently measured
using the new technique based on simultaneous OCT and US
imaging. We used the OCT system and a clinically applicable
US system (Clear View Ultra; Boston Scientific Corporation,
Marlborough, MA) providing cross-sectional images. Both
the thin US catheter (diameter = 1.0 mm, Fc = 40 MHz, Fs
= 250 MHz; Atlantis SR Pro; Boston Scientific Corporation)
and the OCT catheter were inserted through an oval-shaped,
custom-made instrument channel to enable imaging of the
same location simultaneously with both modalities (Figure
2). Considering the largest diameter of the instrument chan-
el (3.33 mm), it is possible to insert the instrument into a
joint through normal arthroscopy portals. The phantoms on
the metal plate and the instrument were immersed in degassed
distilled water during the measurements. The instrument was
aligned to direct the US beam perpendicular to the phantom
surface. Each phantom SOS measurement set consisted of 10
consecutive US scans at the same location recorded simulta-
neously with 10 adjacent OCT cross sections. 5 of these sets
of measurements were subsequently performed without repo-
positioning the catheters. The reproducibility of the technique
was evaluated by performing the set of 5 measurements 3
times and repositioning the catheters between each set.

Our technique was tested further by measuring SOS in
equine cartilage in the laboratory. In these measurements,
degassed PBS containing disodium EDTA and benzamidine
HCl was used instead of water. To mimic arthroscopic con-
ditions, the instrument was manually held over the cartilage
surface and—to ensure a perpendicular US incidence angle—
aligned so that the US reflection from the tidemark was maxi-
mal. The measurement was performed 5 times for each osteo-
chondral sample. After the measurements, the samples were stored at −20°C until they were prepared for histology.

**Determination of time of flight (TOF) and sample thickness**

TOF and sample thickness were measured from the exact raw US and OCT data, respectively. In osteochondral samples, US reflects from the cartilage surface and tidemark (Modest et al. 1989). The surface of the phantom or cartilage was defined automatically, and the bottom of the phantom or cartilage tidemark was defined manually for each of the 10 OCT cross sections. A 1-mm-wide analysis window was placed under the US catheter to match the thickness and TOF measurement sites in the lateral direction (Figure 3). The average thickness within the window was calculated.

Hilbert envelopes of bandpass filtered (tenth-order Butterworth filter with upper and lower cutoff frequencies of 80 MHz and 15 MHz, respectively) ultrasound A-scan lines were determined. Phantom or cartilage surface and phantom bottom or cartilage tidemark were manually determined from as wide an area of the cross-sectional image reconstructed from the envelope signals as was feasible (Figure 3). TOF in cartilage was measured from the average time distance between the US catheter and cartilage, and the optimization was conducted in the second step (5–10% strain). A more detailed description of the model is available in earlier studies (Li et al. 1999, Korhonen et al. 2003, Wilson et al. 2004, Julkunen et al. 2007, Kulmala et al. 2012).

**Histological and spectroscopic analysis**

Osteochondral blocks with the marked site of interest exactly in the middle were immersed in formalin for at least 48 h and then processed for histological evaluation (Kulmala et al. 2012). Three safranin-O-stained, 3-µm-thick sections from each sample were prepared and imaged with a light microscope. Cartilage thickness was measured manually after careful comparison of measurement location with OCT image. Three investigators graded the 3 stained sections of each sample using the Mankin score (Mankin et al. 1971). Final grades were obtained by averaging the scores and rounding up the average to the nearest integer. Optical density (OD) measurement of safranin-O distribution was conducted by means of digital densitometry (computer-controlled CCD camera; SenSys; Photometrics Inc., Tucson, AZ) to evaluate the fixed charge distribution in tissue (Panula et al. 1998). Collagen and PG contents were determined by measuring the areas under the absorption spectra at 1,585–1,720 cm⁻¹ and 984–1,140 cm⁻¹ measured from 3 unstained 5-µm-thick sections using a Fourier transform infrared spectroscopy (FTIR; Spotlight 300 FTIR; Perkin Elmer, Waltham, MA) (Boskey and Pleshko Camacho 2007). Polarized light microscopic analysis of 3 unstained 5-µm-thick sections was conducted (PLM; Photometrics Inc., Tucson, AZ) to provide information on the orientation of the collagen fibrils. We determined the average collagen fibril orientation consisting of two 5% strain steps with a strain rate of 100% per second relative to the pre-strain thickness was performed. A relaxation slope of less than 10 Pa/min was used as the equilibrium criterion.

Using Abaqus (v6.10-1; Dassault Systèmes Simulia Corp., Waltham, MA) and Matlab (2012a, The MathWorks Inc., Natick, MA), an axisymmetric fibril-reinforced poroelastic finite element model was fitted to the experimental stress-relaxation data (Julkunen et al. 2007, Mäkelä et al. 2012). Cartilage was modeled using axisymmetric 4-node continuum pore pressure elements (CAX4P). Elastic fibrillar and biphasic porohyperelastic non-fibrillar matrices represented the collagen network and the PGs with a porous structure filled with fluid, respectively. The behavior of the collagen network was expressed with the fibril network modulus (E_f) and that of the non-fibrillar matrix with the non-fibrillar matrix modulus (E_{nf}), permeability (k), and Poisson’s ratio. The Poisson’s ratio (0.42) and fluid fraction (80%) were fixed (Li et al. 1999, Korhonen et al. 2003), while the other aforementioned material parameters were obtained through optimization, minimizing the mean square error between the experimental and simulated reaction forces. The first 5% achieved a perfect contact between the indenter and cartilage, and the optimization was conducted in the second step (5–10% strain). A more detailed description of the model is available in earlier studies (Li et al. 1999, Korhonen et al. 2003, Wilson et al. 2004, Julkunen et al. 2007, Kulmala et al. 2012).

**Mechanical measurements**

The mechanical properties of the osteochondral samples were determined using a custom-made material testing system (resolution for deformation and force: 0.1 µm and 5 mN, respectively) (Töyräs et al. 1999) equipped with a cylindrical plane-ended indenter (530 µm diameter). The indenter was driven into contact with the sample surface; this was seen from the reading of the load cell. Thereafter, the stress-relaxation test
in relation to the cartilage surface, and the parallelism index (PI) describing the anisotropy of the collagen matrix.

**Numerical assessment of SOS measurement error**

The theoretical performance of the SOS measurement technique was numerically assessed by calculating the propagation of uncertainty. First, the effect of varying sample thickness and resolution of TOF measurement on SOS determination was studied. Perfect accuracy in thickness measurement was assumed while the thickness was varied from 0.3 mm to 3 mm, and TOF resolution corresponding to the measurement error was varied between 0 µs and 0.05 µs. Then we examined the effects of varying sample thickness and the resolution of thickness measurement on SOS determination. Perfect accuracy in TOF was assumed while the sample thickness and resolution of thickness measurement were varied from 0.3 mm to 3 mm and from 0 µm to 50 µm, respectively. In the analyses, true SOS was assumed to be 1,600 m/s.

**Statistics**

All 94 osteochondral samples were included in the analysis of OCT thickness measurement. 36 samples were excluded from the analysis of SOS measurement because the US reflection from either the cartilage surface or the tidemark was too weak for TOF measurement; 9 other samples were excluded because the cartilage was too damaged for mechanical measurement.

Spearman’s rank correlation coefficient was calculated to investigate the correlation between cartilage thickness measured from OCT and microscopic images of the samples. The same method was used to study the correlation between the SOS values in phantoms, measured using the reference and the novel techniques. Possible bias in the measurements was investigated using Bland-Altman analysis. The reproducibility of the SOS measurements was verified by calculating the root mean square coefficient of variation (CVrms(%) for the phantom and cartilage measurements. The statistical significance of differences in cartilage thickness and SOS values between the anatomical locations was tested using the Kruskal-Wallis test. Linear dependencies between the SOS and the cartilage properties determined were evaluated using linear mixed-model analysis, which accounts for the correlations or clusters in the data set. The present study used the joint from which the sample was prepared as a random variable. The models assume that the residuals and the random effect are normally distributed and that the residuals are homoscedastic. The normality was assessed from quantile-quantile plots and homoscedasticity by investigating the variation of residuals as a function of predicted values. The results of these analyses supported the initial assumptions. The significance level in all the analyses was set to p < 0.05. The analyses were performed using SPSS and Matlab.

**Results**

The SOS values in the phantoms measured using the combination of US and OCT were consistent with the values measured using the reference technique ($\rho > 0.99, p < 0.001$) and no systematic error was found based on Bland-Altman analysis (Figure 4). CVrms(%) in SOS measured for the phantoms was 3.4%.

In OCT images, calcified cartilage was discernible from non-calcified cartilage and bone in all but 3 samples (Figure 5). The mean full cartilage thickness and the mean non-calcified cartilage thickness measured using OCT were 1.00 mm (SD 0.26) and 0.83 mm (SD 0.25), respectively. Samples from P1 (mean 1.03 mm (SD 0.27)) had significantly thicker non-calcified cartilage than samples from SR (0.74 mm (SD 0.16), $p = 0.001$) or MC3 (0.71 mm (SD 0.13), $p < 0.001$).
The maximum full cartilage thickness and the maximum non-calcified cartilage thickness measurable with OCT were 1.80 mm and 1.44 mm, respectively. For the thinnest cartilage, the thicknesses were 0.53 mm and 0.37 mm, respectively.

There was a significant correlation between full cartilage thickness measured from OCT images and that measured from microscopic images ($\rho = 0.85; p < 0.001$), while the corresponding correlations for non-calcified and calcified cartilage thicknesses were $\rho = 0.92 (p < 0.001)$ and $\rho = 0.22 (p = 0.04)$, respectively (Figure 6).

The CVrms(%) for the repeated measurements of thickness, TOF, and SOS of the equine cartilage was 7.3%, 7.6%, and 7.8%, respectively. The mean SOS in the cartilage samples was 1,636 m/s. The median Mankin scores of all the samples and of those included in the SOS measurement were 4 and 3, respectively (Table 1). SOS was not significantly dependent on anatomical location ($p = 0.3$). No statistically significant linear dependence was detected between SOS and the cartilage properties determined (Table 2).

The numerical assessment of error in SOS measurements revealed that especially in thin equine cartilage, the resolutions of US and OCT systems have a prominent effect on the accuracy of SOS measurement (Figure 7).

**Table 1.** Mean speed of sound (SOS), mean non-calcified cartilage thickness, and median Mankin score of the samples included in the speed of sound measurements, grouped by anatomical location

<table>
<thead>
<tr>
<th>Anatomical location</th>
<th>n</th>
<th>SOS, m/s (SD)</th>
<th>Thickness, mm (SD)</th>
<th>Mankin score median (IQR)</th>
</tr>
</thead>
<tbody>
<tr>
<td>MC3</td>
<td>26</td>
<td>1,604 (195)</td>
<td>0.66 (0.16)</td>
<td>2 (1)</td>
</tr>
<tr>
<td>P1</td>
<td>7</td>
<td>1,720 (169)</td>
<td>0.74 (0.26)</td>
<td>7 (3)</td>
</tr>
<tr>
<td>SR</td>
<td>16</td>
<td>1,651 (210)</td>
<td>0.68 (0.14)</td>
<td>3 (2)</td>
</tr>
<tr>
<td>All</td>
<td>49</td>
<td>1,636 (197)</td>
<td>0.68 (0.17)</td>
<td>3 (3)</td>
</tr>
</tbody>
</table>
| **Note:** MC3: dorsoproximal areas of the condyles of the third metacarpal bone; P1: proximal phalanx; SR: sagittal ridge of the third metacarpal bone; IQR: interquartile range.

**Table 2.** p-values for linear association between speed of sound (SOS) and the fixed effects (structural, compositional, and biomechanical properties) using the linear mixed model

<table>
<thead>
<tr>
<th>Mankin score</th>
<th>Optical density</th>
<th>Collagen orientation</th>
<th>Parallelism index</th>
<th>Proteoglycan content</th>
<th>Collagen content</th>
<th>$E_f$</th>
<th>$E_m$</th>
<th>$k$</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.5</td>
<td>0.8</td>
<td>0.4</td>
<td>0.9</td>
<td>0.8</td>
<td>0.4</td>
<td>0.5</td>
<td>0.4</td>
<td></td>
</tr>
</tbody>
</table>

$E_f$: fibril network modulus; $E_m$: non-fibrillar matrix modulus; $k$: permeability.
et al. 2003). However, no statistically significant dependence and Mankin 1997, Joiner et al. 2001, Suh et al. 2001, Töyräs control its mechanical and acoustic properties (Buckwalter al. 2005). The structure and composition of articular cartilage with the values given in the literature (1,696 m/s) (Brommer et materials. The mean SOS in the equine cartilage samples agreed OCT and US would enable SOS measurement in layered mate-

trials. The relative difference between OCT and microscopic materials. The relative difference between OCT and microscopic measurement was found between SOS and the structural or mechanical properties determined.

Repeatability in SOS measurement is reduced if the surface is fibrillated, as fibrillation impairs the accuracy of locating the cartilage surface, particularly from US images. However, SOS measurements may be unnecessary when there is visible fibrillation, as osteoarthritis that has advanced as far as this can be diagnosed with conventional techniques.

We estimated the error in SOS measurement caused by inaccuracy in the thickness and TOF measurements with different cartilage thicknesses. Theoretically, the resolutions of the current OCT and US systems are sufficient for SOS measurement. However, the accuracy of the present measurement is partially limited by the less than optimal optical and acoustic contrasts between non-calcified and calcified layers. Attenuation of light and US in cartilage further limits the contrast in the tidemark, and therefore the highest cartilage thickness and TOF measurable with this technique. Thus, the technique requires further development before it would be suitable for clinical use in equine or human medicine. Advanced signal and image processing could enhance the contrast, and the use of point measurement or linear scanning instead of rotational scanning could improve measurement accuracy. Furthermore, the US frequency and intensity could be optimized to increase the penetration of US into the tissue. The accuracy of the US and light incidence angle, which should be perpendicular to the cartilage surface, could also be improved through improved probe design.

In conclusion, articular cartilage thicknesses below 1.8 mm were successfully measured with OCT. OCT-based thickness measurements were more accurate when calcified cartilage was excluded. The measurement could be used for arthroscopic detection of cartilage thinning during arthroscopy and estimation of lesion depth. Before becoming suitable for clinical use, the measurement accuracy of the SOS measurement technique presented must be improved. However, the concept of simultaneous OCT and US imaging for SOS measurement appears to be both feasible and promising.

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All authors contributed to the study design, interpretation of results, and critical review of the manuscript. In addition, PP performed experimental work, data analysis, and writing of the manuscript, NiM performed experimental work and data analysis, and PT performed experimental work.

No competing interests declared.

of the measurement of calcified cartilage thickness, and consequently that of the measurement of full cartilage thickness. According to our Bland-Altman analysis, the absolute difference between the 2 modalities increased with thicker cartilage. The relative difference between OCT and microscopic measurement was similar to that in previous studies that investigated rabbit and goat cartilage with thicknesses less than 1 mm (Han et al. 2003, Rogowska et al. 2003, Cerno-

Figure 7. Numerically evaluated effect of resolutions of US (ΔTOF) measurements (panel a) and OCT (Δd) measurements (panel b) on error in SOS values determined for cartilage of varying thickness.

We compared light microscopic images and OCT images before selecting the thickness measurement points. Although the mismatch between the measurement locations was small in all samples, it could still be the main reason for the difference in thickness values between OCT and microscopy. Additionally, fixation and processing of the samples for microscopy may have a small shrinking effect on sample dimensions. Some inaccuracy in OCT measurements may be due to the use of the same value in the literature for the refractive index of both non-calcified and calcified cartilage.

We hypothesized that by using OCT thickness measurement with simultaneous US TOF measurement, arthroscopic determination of cartilage SOS would be possible. The technique that we introduced enabled simultaneous measurement of US TOF in articular cartilage and cartilage thickness at the same location. Our technique may be used in conventional arthroscopy, as the instrument fits through normal arthroscopic portals.

Based on the phantom measurements, the combination of OCT and US would enable SOS measurement in layered materials. The mean SOS in the equine cartilage samples agreed with the values given in the literature (1,696 m/s) (Brommer et al. 2005). The structure and composition of articular cartilage control its mechanical and acoustic properties (Buckwalter and Mankin 1997, Joiner et al. 2001, Suh et al. 2001, Töyräs et al. 2003). However, no statistically significant dependence was found between SOS and the structural or mechanical properties determined.


