Is there a correlation between biophotonical, biochemical, histological, and visual changes in the cartilage of osteoarthritic knee-joints?

Stephanie Tatjana Stumpfe 1
Julia Karin Pester 1,2
Susanne Steinert 1
Ivan Marintschev 1
Holger Plettenberg 1,3
Matthias Aurich 1
Gunther Olaf Hofmann 1,2

1 Department of Traumatology, Hand and Reconstructive Surgery, Jena University Hospital, Jena, Germany
2 Department of Traumatology and Orthopaedic Surgery, Trauma Centre Halle (Saale), Jena, Germany
3 Arthrospec GmbH, Jena, Germany

Corresponding author:
Gunther O. Hofmann
Department of Traumatology
Hand and Reconstructive Surgery
Jena University Hospital
Erlanger Allee, 101
07747 Jena, Germany
E-mail: gunther.hofmann@med.uni-jena.de

Summary

The aim of this study was to detect characteristic structural changes in the cartilage composition of osteoarthritis (OA), hereby improving the arthroscopic identification of cartilage pathology by the use of a non-destructive technique - NIRS (Near-Infrared Spectroscopy). 682 cartilage samples out of 25 knees with OA were classified visually, using the ICRS system, biophotonically, histologically (n = 66), using the Score of Mankin and the Score of Otte, and biochemically (n = 616), determining the content of glycosaminoglycan (GAG) and hydroxyproline (HP). Significant correlations were found between biophotonical, histological, biochemical and visual characteristics of cartilage lesions. NIRS values corresponded to the content of GAG, HP and to the Score of Mankin and Otte. The data show that changes in the composition and structure of articular cartilage influence the optical properties and can be measured objectively by NIRS. The ease of use during arthroscopy, the quick response and the non-destructive nature of NIRS make it a promising addition to the assessment of disease intervention in OA.

KEY WORDS: cartilage, GAG, hydroxyproline, Mankin score, NIRS, osteoarthritis.

Introduction

OA is the most common joint disorder worldwide, being prevalent in the elderly, but also affecting younger people 1. However, the pathogenesis of OA is a multifactorial process and involves the whole joint and soft tissue structures in and around the joint 2.

OA causes several changes in cartilage and leads to loss of tissue integrity and function. The breakdown of the extracellular matrix (ECM) is most prominent and is indicated through the loss of the major matrix molecules, including proteoglycan and collagen 3. This matrix transformation occurs although the chondrocytes are thought to have anabolic activity, such as matrix synthesis and proliferation. The cells seem to counteract the matrix degradation during the degenerative events of OA 4,5. The biomechanical, biochemical, and histological properties of articular cartilage change as a consequence of cartilage degeneration in OA 6,7. Structural changes result in a break-up of the collagen-proteoglycan matrix with consequent changes in the water-binding properties and a shift in the composition of main cartilage components 8,9.

There is increasing research interest in the physiological and pathological function of vital articular cartilage depending on the structure, the composition and the integrity of the tissue, carried out by correlating results of biophysical, biomechanical, biochemical, and histological methods. These methods proved as reliable and were validated in the past. However, nearly all of the methods in cartilage research are tissue-destructive techniques effecting the cartilage samples and are not applicable in joint preserving surgery. NIRS is a non-destructive optical technique in the range of 780 nm up to 2500 nm and widely employed in many fields of medicine 10,11. With this biophotonic measurement procedure it is possible to detect the overtone and combined vibration of for example C-H, O-H and N-H groupings. Therefore it is possible to get the link with this biophotonic measurement method to the chemical and biomechanical properties of analysed cartilage. Recent studies have shown that NIRS is a proven sensitive and accurate diagnostic in-vivo method for quantitative determination of structural and mechanical changes in cartilage 12,13. Since infrared light interacts with cartilage components affected by OA, such as water, water-binding properties, CH-OH-and SH-groups, early stage OA are detectable with this method 12,13. Still, the rationale of this is not clearly understood.

The topic of this current study was to prove whether NIRS is able to identify structural and biochemical
changes in human cartilage of OA and to evaluate the possible correlation between cartilage breakdown and loss of histological structure with biophotonical values. We hypothesized that NIRS is sufficient for the prediction of these processes of intra-cartilaginous degeneration.

Methods

Patients

Twenty-five knee-joints in 24 patients (13 women, 11 men) with severe osteoarthritis (OA) due to clinical (KOOS)\textsuperscript{15,16} and radiological (Kellgren-Lawrence)\textsuperscript{17} scores received total knee replacement (TKR) with patella resurfacing and were enclosed in this prospective, single-centre, open-label study between March, 2009 and December, 2009. Mean age of the patients was 68.71 years (women 71.62 years, men 64.18 years), ranging from 52 to 80 years. All implantations were indicated by clinical (pain, loss of function and mobility) and radiological signs of OA. Informed consent was obtained from all patients after the nature of all examinations had been fully explained, following written approval from the local Ethics Committee for Clinical Trials (1714-01/06). All procedures were in accordance with the Helsinki Declaration concerning human trials.

Surgery

All patients received TKR under general anaesthesia, performed by two experienced surgeons. The cartilage in the femorotibial and femoropatellar compartments was independently examined by them using the classification approved by the International Cartilage Repair Society (ICRS)\textsuperscript{18}. Joint surfaces were resected under continuous irrigation with Ringer’s solution for cooling, covered with PBS-soaked gauze and immediately transferred to the laboratory for further investigations. Only cartilage samples of ICRS grades 1 - 3 were processed further in this study.

Sample preparation

Full-thickness osteochondral cylinders (n = 682) including subchondral bone with a diameter of 4 mm were drilled out of three different surface areas (femoral, tibial and patellar) using a gouge bit under continuous irrigation with Ringer’s solution for cooling. The samples represented various stages of cartilage damage (7 ICRS grade 0; 360 grade 1; 190 grade 2; and 125 grade 3). Articular compartments with ICRS grade 4 were excluded from further investigation. The drilled-out specimens were initially examined biophotonically and then stored in acetone (Roth, Karlsruhe, Germany) for the further histological preparation or frozen for the biochemical examination.

Biophotonical testing

All samples (n = 682) were examined by a prototype NIRS device for diffuse reflectance measurement (arthrospec ONE, arthrospec GmbH, Jena, Germany). The system consists of a photodiode array spectrometer (950 nm – 1700 nm, 256 pixel), a stabilized broadband light source, a fibre-optic reflectance probe and associated hard and software for data acquisition (Fig. 1). For measurement, the face of the probe was placed on the cartilage surface of each specimen and five subsequent reflectance spectra were recorded (100 ms recording time per spectrum). To search for fundamental relations between laboratory data (presented as single value vectors) and optical data (matrix of spectral data) we used a multivariate regression model (partial least square regression) to calculate linear correlations. To avoid overfitting, we restricted the number of latent factors to five. The goodness of fit is described by the correlation coefficient between the best model and the laboratory data. A schematic diagram of the test setup is given in Figure 2.

Due to the exclusion of articular compartments with ICRS grade 4, there were different numbers of samples of each knee available which were examined by NIRS and processed during the following procedures. After the biophotonical testing, the specimens were divided into two groups: the first group concluded specimen of each patient with each ICRS grade for histological evaluation (n = 66). The second group concluded remaining samples for biochemical analysis (n = 616). Criterion for the following selection of samples was the intention to harvest material from each patient with each ICRS grade for histological evaluation. All other specimens served for biochemical analysis.

Biochemical Testing

A total of 616 cartilage samples were shredded and incubated in papain working solution (125 μg/ml) for 16 h at 60°C after evaluation of the dry weight (Ohaus Europe GmbH, Nänikon, Switzerland).

GAG content

GAG content of these prepared samples was evaluated using the dimethyl-methylene blue (DMMB) assay. GAGs are a covalently attached functional group to the core proteins of proteoglycans. Aggrecan has a high affinity for water due to its high negative fixed-charge density and causes the interaction with the H\textsubscript{2}O-based DMMB reagent (Sigma-Aldrich, Steinheim, Germany). Therefore, the maximum absorption is changed and can be evaluated with a special reader (Sunrise, Tecan, Switzerland) at a wavelength of λ = 525 nm. The samples were diluted three times (1:200, 1:400, 1:800) with dilution buffer (0.05 M Na-acetate and 0.05% Tween 20) and the results were averaged to get reliable information about the GAG content of
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For each sample, for each passage a standard curve was made. The results are reported as the mean ± SD of the analysis of three different diluted cultures.

**HP content**

All chemicals were purchased from Sigma Aldrich (Steinheim, Deutschland) or Thermo Fisher (Waltam, USA) unless otherwise noted. Determination of HP was performed by a colorimetric assay after papain digestion and freeze-drying. Aliquots from each digest were hydrolysed in 4 N NaOH for 30 min at 125°C. The hydrolysates (ph = 6.0) were incubated for 20 min at room temperature with chloramine-T and afterwards with aldehyde-perchloric acid for 20 min at 70°C for colorimetric reaction. The absorbance at 550 nm was determined with a spectrophotometer (Sunrise, Tecan, Switzerland). The content of HP in each sample was calculated after generation of a standard curve using known concentrations of gelatin. The calculated HP content was used to determine the total amount of collagen by multiplying the results by a factor of 8.2. The results are reported as the mean ± SD of the analysis of three separate cultures.

**Histological examination**

**Sample embedding**

The embedding of the specimens was carried out using Technovit 9100 (Heraeus Kulzer, Wehrheim/Ts., Germany) according to the manufacturers’ protocol. This polymerisation system is based on methylmethacrylate (MMA) that hardens at low temperatures and has been developed especially for the embedding of mineralised tissues. The embedded samples were cut into 4 µm sections using a Leica microtome (Polycut S Reichert-Jung, Bensheim, Germany). Each sample was stained histologically for grading according to Mankin and Otte.

![Figure 1. Design of biophotonic test setup. Full-thickness osteochondral cylinders from femoral surface area (a) were analysed by a prototype NIRS device (c-e). For measurement, the face of the probe was placed on the cartilage surface of each specimen (a, b). The system consists of a photodiode array spectrometer (e), a stabilized broadband light source, a fibre-optic reflectance probe (c, d) and associated hard and software for data acquisition.](image)
Microscopic staging

Cartilage samples were staged microscopically according to the modified Mankin score\textsuperscript{19}, which is estimated to be the actual gold standard scale concerning the evaluation of cartilage degeneration. The sections were stained with Hematoxylin-Eosin and Safranin-O and investigated microscopically. The four different aspects of the score were determined and summed up leading to a scale that ranges between 0 and 14. The Mankin score 0-2 represents normal cartilage, 3-5 - superficial fibrillation, 6-7 - moderate cartilage destruction, 8-10 - severe damage of cartilage, and over 10 - complete loss of cartilage. In addition, cartilage samples were classified microscopically according to the Otte score\textsuperscript{20}. Chondral lesions were subdivided into four grades: Grade 0 represents intact normal cartilage, grade 1 - superficial fibrillation, grade 2 - chondral lesions (fissures and cracks) without extending down to subchondral bone, grade 3 - chondral lesions extending down to subchondral bone, and grade 4 - focal or general full-thickness cartilage loss.

Statistical Analysis

We used PASW Statistics (PASW Statistics 17, SPSS Inc., Chicago, USA) for statistical analysis and Microsoft Excel (Microsoft Office Excel 2003, Microsoft Corp., Redmond, USA) for data visualization. The data are presented as means, whereas we used normalised data to evaluate the relative change between the consecutive measurements. For the establishment of the biophotonical correlation analysis all samples deemed as independent variable because bivariate correlations between several samples and six methods were performed. All other statistical analyses were made assuming that the variables were dependent. Normal distribution was tested with the Kolmogorow-Smirnow test. In case of normal distribution, we employed the t-test; otherwise, we used the Wilcoxon test. When parameters showed no normal or linear distribution, Spearman’s rank correlation coefficient $\rho$ was used to characterise links between parameters. Significance was defined as $P \leq 0.05$. Samples of the small group of ICRS grade 0 were excluded for statistical evaluation.

Results

Macroscopic visual evaluation

The samples represented various stages of cartilage damage, and cartilage abrasion was evident in most specimens. Only a small number of classified samples ($n = 7$) showed no OA signs (ICRS grade = 0), 360 samples showed mild OA signs (ICRS grade = 0.1), and 360 samples showed moderate OA signs (ICRS grade =...
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1), 190 samples showed grade 2 associated alterations, and 125 samples showed advanced OA signs (ICRS grade = 3).

Biochemical evaluation

The group includes a total of 616 samples, consisting of six samples of ICRS grade 0, 338 samples of ICRS grade 1, 167 samples of ICRS grade 2, and 105 samples of ICRS grade 3. The results are summarized in Table 1 and visualized in Figure 3. The average content of GAG was 35.62 µg/mg (SD: 14.15 µg/mg) and of HP 96.34 µg/mg (SD: 49.78 µg/mg). If considered separately according to the ICRS grades, the content of GAG and HP showed no significant differences in the mean values. This is reflected by the highly variable values of the GAG and HP content as a function of an inconsistent intra-individual and inter-individual appearance. Values obtained from the biochemical analysis showed a significant (P < 0.001) correlation between increasing ICRS grade and the content of GAG (p = 0.521 for grade 1~3 up to p = 0.826 for grade 1~2; Fig. 3a) and HP (p = 0.645 for grade 1~3 up to p = 0.852 for grade 1~2; Fig. 3b). Furthermore, values showed minor regional distinctions with significant correlations between biochemical evaluation and the ICRS grade according to the femoral, medial and patellar compartment (Tab. 1). Values showed no correlation between the content of GAG and HP.

Histological evaluation

A total of 66 specimens were chosen for the histological examination representing the different stages of cartilage damage according to the ICRS classification (one sample of ICRS grade 0, 22 samples of ICRS grade 1, 23 samples of ICRS grade 2, and 20 samples of ICRS grade 3).

Values assessed from the histological analysis showed strong significant (P < 0.001) correlations between increasing ICRS grade and Mankin (p = 0.725) or Otte score (p = 0.736). Histological classification increased significantly with the macroscopic grade of cartilage damage classified by ICRS. Furthermore, values assessed from the histological analysis measured by the Score of Mankin were associated significantly with the grade of cartilage lesion according to the Score of Otte (p = 0.814). The results are summarized in Table 2 and visualized in Figure 4. The histological evaluation showed clear structural disorganization; fibrillation and cartilage defects increased significantly (P < 0.001) from ICRS grade 0 to 3 and approved the ICRS classification. Areas macroscopically classified as ICRS grade 1 showed fissures and cracks in the superficial and intermediate layer, hypercellularity and a decrease in safranin-O staining and were adjusted with 5.5 ± 0.71 points on the Mankin score and 1.19 ± 0.15 on the Otte score. ICRS grade 2 with cell-clustering of chondrocytes, deep cracks penetrating

Table 1. Biochemical results of the evaluation of GAG and HP content according to regional affiliations of the samples and the grade of cartilage lesion classified by ICRS.

<table>
<thead>
<tr>
<th>Localization</th>
<th>ICRS</th>
<th>General</th>
<th>Femoral</th>
<th>Tibial</th>
<th>Patellar</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>GAG</td>
<td>µg/mg</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>µg/mg cartilage</td>
<td>0</td>
<td>32.4 (18.2) [6]</td>
<td>24.5 (17.1) [3]</td>
<td>48.1 (--o) [3]</td>
<td>-- [0]</td>
</tr>
<tr>
<td></td>
<td>1</td>
<td>36.2 (11.4) [334]</td>
<td>37.6 (14.2) [254]</td>
<td>34.9 (14.8) [58]</td>
<td>34.1 (8.1) [25]</td>
</tr>
<tr>
<td></td>
<td>2</td>
<td>34.4 (10.9) [168]</td>
<td>37.5 (11.1) [96]</td>
<td>33.2 (12.9) [40]</td>
<td>33.9 (12.9) [32]</td>
</tr>
<tr>
<td></td>
<td>3</td>
<td>32.4 (14.2) [105]</td>
<td>39.5 (15.6) [42]</td>
<td>28.3 (10.6) [42]</td>
<td>26.1 (9.3) [21]</td>
</tr>
<tr>
<td></td>
<td>HP</td>
<td>µg/mg</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>µg/mg cartilage</td>
<td>0</td>
<td>115.6 (68.3) [6]</td>
<td>139.5 (77.1) [3]</td>
<td>67.9 (--o) [3]</td>
<td>-- [0]</td>
</tr>
<tr>
<td></td>
<td>1</td>
<td>95.8 (44.9) [334]</td>
<td>95.7 (47.3) [254]</td>
<td>102.5 (56.1) [58]</td>
<td>99.9 (25.7) [25]</td>
</tr>
<tr>
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<td>2</td>
<td>80.9 (42.4) [168]</td>
<td>82.9 (46.7) [96]</td>
<td>79.4 (35.3) [40]</td>
<td>94.2 (38.5) [32]</td>
</tr>
<tr>
<td></td>
<td>3</td>
<td>77.8 (45.6) [105]</td>
<td>76.2 (46.1) [42]</td>
<td>96.4 (58.8) [42]</td>
<td>72.1 (46.5)[21]</td>
</tr>
</tbody>
</table>

Data are expressed as means, standard deviation and total of samples. * Significant change at a level of P < 0.05; ** significant change at a level of P < 0.001

Figure 3. Dependence and correlation analysis between biochemical changes measured by the content of GAG (a) and HP (b), and the macroscopic grade of cartilage lesion classified by ICRS via bivariate correlations.
to the middle zone and a progressive decrease in safranin-O staining showed 7.51 ± 0.43 points on the Mankin score and 1.94 ± 0.15 on the Otte score. Cartilage changes associated with areas evaluated as ICRS grade 3 appeared with increased clefts and disruption of the matrix with cell-clustering, hypercellularity and an increase of the cartilage thickness. The Mankin score of this advanced stage of degeneration was 9.53 ± 0.3 points and the Otte score was 2.67 ± 0.14. Furthermore, values showed minor regional distinctions with significant correlations and differences between histological evaluation (Score of Mankin and Otte) and the ICRS grade according to the femoral, medial and patellar compartment (Tab. 2).

**Discussion**

The purpose of the current investigations was to prove whether NIRS is an objective and valid predictor identifying structural and biochemical changes in human cartilage of OA. To establish a diagnostic probe for the non-destructive evaluation of cartilaginous changes in composition and structure articular cartilage from human knee-joints with different degrees of OA was examined using biochemical, histological and biophysical techniques. Our data showed that biophotonical, histological, biochemical and visual characteristics of cartilage lesions correlated significant. Especially, NIR light absorption corresponded to the macroscopical grade of cartilage lesions and to biochemical changes of cartilage structure and composition detected by the content of GAG and HP. According to previous studies, our results show that there is a strong correlation between the histological grading of the OA and the ICRS classification, and that the histological evaluation is one of the most reliable methods to determine the qualitative and quantitative extent of changes in cartilage. However, this method is limited to *ex vivo* examinations. We are conscious that the arthroscopic grading of cartilage lesions due to the ICRS score as applied in this study is limited and has a poor inter-observer reliability, especially between intact cartilage (grade 0) and lesions that consist of softening of the cartilage.

**Table 2. Histological results of the evaluation of the Mankin score and Otte score according to regional affiliations of the samples and the grade of cartilage lesion classified by ICRS.**

<table>
<thead>
<tr>
<th>Localization</th>
<th>ICRS</th>
<th>General</th>
<th>Femoral</th>
<th>Tibial</th>
<th>Patellar</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mankin score</td>
<td>0</td>
<td>5.5 (--) [1]</td>
<td>5.5 (--) [1]</td>
<td>-[0]</td>
<td>-[0]</td>
</tr>
<tr>
<td></td>
<td>1</td>
<td>5.5 (0.7) [22]**</td>
<td>5.4 (0.5) [21]**</td>
<td>6.0 (--) [1]**</td>
<td>-[0]</td>
</tr>
<tr>
<td></td>
<td>2</td>
<td>7.5 (0.4) [23]**</td>
<td>7.4 (0.6) [16]**</td>
<td>9.0 (0.2) [5]**</td>
<td>4.5 (1.4) [2]**</td>
</tr>
<tr>
<td></td>
<td>3</td>
<td>9.5 (0.3) [20]**</td>
<td>9.9 (0.3) [10]**</td>
<td>9.0 (0.3) [9]</td>
<td>10.0 (--) [1]**</td>
</tr>
<tr>
<td>Otte score</td>
<td>0</td>
<td>1 (--) [1]</td>
<td>1 (--) [1]</td>
<td>-[0]</td>
<td>-[0]</td>
</tr>
<tr>
<td></td>
<td>1</td>
<td>1.2 (0.2) [22]**</td>
<td>1.1 (0.7) [21]**</td>
<td>1 (--) [1]**</td>
<td>-[0]</td>
</tr>
<tr>
<td></td>
<td>2</td>
<td>1.9 (0.2) [23]**</td>
<td>1.9 (0.7) [16]**</td>
<td>2.4 (0.5) [5]**</td>
<td>1.5 (0.7) [2]**</td>
</tr>
<tr>
<td></td>
<td>3</td>
<td>2.7 (0.1) [20]**</td>
<td>2.9 (0.6) [10]**</td>
<td>2.4 (0.5) [9]</td>
<td>3 (--) [1]**</td>
</tr>
</tbody>
</table>

Data are expressed as means, standard deviation and total of samples. * Significant change at a level of P < 0.05; ** significant change at a level of P < 0.001
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(grade 1), and differentiation between superficial and deep lesions. All specimens, therefore, underwent a detailed histological scoring according to Mankin and Otte. As has already been shown in previous studies, our results demonstrated that there is no histologically normal tissue in the OA joint. The representative selection of histological samples (n = 66) shows no regions that are classified microscopically (n = 0) and macroscopically (n = 1) as normal cartilage.

It is a challenging task to establish a simple correlation between defined microscopical changes of cartilage structure and changes in biochemical composition. The biochemical composition of human cartilage (content of collagen and GAG) changes as a function of age, but also in OA. According to age, the collagen content of human cartilage is discussed controversially. Williamson et al. found that the collagen concentration in bovine cartilage increases during life as a function of age. In contrast, the GAG content remains unchanged or decreases slightly. Other investigators found that proteoglycans are smaller and less functional with age. Nevertheless, it is important to note that while changes in the composition and structure of the cartilage matrix take place inevitably, the development of OA with aging is not a uniform process. This could be one explanation for our results showing no significant differences in matrix composition according to ICRS, and reflected the high variability of the GAG and HP content as a function of an inconsistent intra-individual and inter-individual appearance. Furthermore, the mean age of our study patients is 68.71 years and structural changes observed during aging are not discriminated from disease-associated changes of articular cartilage.

Concerning the changes of the composition of cartilage in OA, some authors showed that the content of proteoglycan of the ECM is reduced in general and that these proteoglycans exist in a non-aggregated and smaller form than in normal articular cartilage. Aurich et al. were able to confirm the pronounced increase in collagen II synthesis and aggrecan turnover for the ankle-joint but not for the knee-joint. Early cartilage lesions in the knee-joint typically show an increase of collagen II cleavage. Another result of the OA process and, additionally, a factor of etiology of the disease is the deterioration in the mechanical properties of the cartilage. As a function of age, the elastic response of articular cartilage to loading forces decreases with decreasing content of GAG. In a number of ex vivo studies in sheep femoral condyles, our group found an increased water content in combination with a decrease in mechanical stiffness, going along with an increasing degree of OA. In contrast to the aforementioned studies in this setting, the biochemical analy-
sis presented no significant differences in GAG and collagen content between intact cartilage (grade 0) and early stages of articular lesions (grade 1). Collagen and GAG content within the chondral defects decreased in tendency only in more severe lesions (grade 2 and grade 3)\textsuperscript{12}. Marticke et al.\textsuperscript{29} found a significant correlation between the mechanical strength (Young’s Modulus) of cartilage and different stages of degeneration quantified by the ICRS grade by investigating human knee-joint material. They also described a strong correlation between these mechanical alterations and biophotonical characteristics in the knee-joint cartilage analysed by the NIRS. In 2008, our group published the first results in evaluation of cartilage defects with NIRS from an \textit{ex vivo} study in sheep femoral condyles\textsuperscript{12}. The NIRS absorption in ICRS grade 1 defects was significantly higher than in intact cartilage (grade 0). ICRS grade 2 specimens tended to a higher NIR absorption than the ICRS grade 1 specimen. A high correlation was found between the water content of the cartilage and NIR absorption, but no correlation was found between NIR absorption and collagen or GAG content\textsuperscript{12}. In contrast, our results demonstrated moderate but significant associations between NIR light absorption and changes of cartilage structure and composition detected by the content of GAG and HP, or the grade of cartilage lesion classified by the Mankin and Otte score. There are several lines of evidence indicating that the biochemical structural properties of cartilage are associated directly with the biophotonical properties\textsuperscript{14,30}.

Our study has a number of limitations which may impair the interpretation of the study findings. Firstly, only cartilage of patients with severe OA needing TKR was available for this investigation. Secondly, our samples are acquired by a plurality of knees not delivering independent samples of each region (femoral, tibial and patellar) and ICRS grade. Thirdly, our investigations were performed \textit{in vivo}. Tissue destructive methods were correlated with one non-destructive \textit{in vivo} method describing changes of cartilage composition. Therefore, it is not remarkable that the correlations to biophotonical properties are only moderately represented compared to well-defined laboratory conditions or to quantitative assays. Nevertheless, we confirmed the correlation between cartilage degradation and changes of cartilage structure and composition with our investigation even if femoral or tibial cartilage was separately surveyed, and we established NIRS for the detection of biochemical and microscopical changes of cartilage composition and lesion. Furthermore the quantitative differentiation of early-stage cartilage defects (grade 1) and intact cartilage (grade 0) is a rather difficult task and is, in part, still unsolved\textsuperscript{7,18}. Moreover, even the differentiation between grade 2 and 3 cartilage defects is difficult concerning the depth of the lesion\textsuperscript{18}. The intraoperative visual and tactile classification is very subjective, based on the individual experience of the investigator and sometimes hardly reproducible\textsuperscript{31}, and has a poor interobserver agreement\textsuperscript{21}. Thus, a more precise description of the location, depth and size of the cartilage lesion adjacent to unchallenged cartilage is mandatory for diagnosis and therapy\textsuperscript{18}. Identi-
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