Effect of light-emitting diode (LED) therapy on the development of osteoarthritis (OA) in a rabbit model

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1. Introduction

Osteoarthritis (OA) is one of the most common diseases of aging, characterized as articular cartilage and underlying subchondral degeneration, with more than 20 million people affected in the United States [1]. OA causes joint pain and a limitation of joint range of motion (ROM), with the end stage resulting in decreased quality of life (QOL). For patients with advanced OA, however, surgical options are available. Osteoarthritic cartilage may be repaired by chondroplasty, drilling and microfracture or it may be replaced through autologous or allogenic osteochondral transplantations in an attempt to remodel the joints to their original morphometry [2–8]. Artificial joint replacement and arthrodesis are salvage procedures for severe osteoarthritic joints [9], and autologous chondrocyte implantation (ACI) is used as a cartilage reconstruction trial [10].

Early OA is usually treated by controlling pain and improving ROM. Conservative procedures begin with weight loss, ROM exercises, muscle stretching and strengthening, followed by medication and intra-articular injections of hyaluronic acid, analgesics and traditional NSAIDs. Long-term use of these drugs may increase gastric mucosal damage and cardiovascular disease [11]. Injection of hyaluronan has been shown to improve joint lubrication and decrease pain [12], however the effectiveness diminishes with time, and repetition can cause joint infection. A long-lasting, noninvasive procedure to provide anti-inflammation and preserve articular cartilage in early OA is therefore clinically desirable.

Light amplification by stimulated emission of radiation (LASER) treatment has been reported to improve wound healing, including increase of cell proliferation, collagen synthesis and growth factor production. The LASER does, however, have limitations in wavelength capacities and beam widths. An optimal wavelength of LASER for each tissue may be difficult for larger areas, and heat productions from LASER light could damage tissue [13].

Light-emitting diodes (LEDs) are complex semiconductors that convert electrical current into incoherent narrow-spectrum light.
LEDs are available at wavelengths ranging from ultraviolet (UV) to visible to near infrared (NIR) bandwidths (247 to 1300 nm). The National Aeronautics and Space Administration (NASA) has reported on the acceleration of plant growth and wound healing following exposure to LEDs. This has led to investigations into nonthermal therapy, especially in dermatology, leading to its approval as a nonsignificant risk procedure for humans by the Food and Drug Administration (FDA) [14].

We hypothesize that LED treatments can penetrate into joints, stimulate the inside of the joint cavity and articular surface, and have the potential to repair OA. The objective of this study was to evaluate the effect of LED light therapy on osteoarthritic knee joints using the anterior cruciate ligament transection (ACLT) model of OA in vivo.

2. Materials and methods

2.1. Animals

A total of 14 skeletally mature, female New Zealand White rabbits (9–15 months old weighing 3.5 to 4.5 kg) with closed epiphyses were used for both control and experimental evaluation (n = 7 for each group). All procedures conformed to the guidelines of the University’s Animal Subjects Committee and the American Association for Accreditation of Laboratory Animal Care.

2.2. Osteoarthritic knee joint model

The study animals were given presurgical general anesthesia injections of 35 mg/kg ketamine and 5 mg/kg xylazine intramuscularly. An anesthesia machine was attached to a facemask delivering 1–2% isoflurane for maintenance of anesthesia during surgery. The central part of the right leg was shaved and draped in sterile fashion using betadine. A 3 cm anterior midline incision was made through the skin of the knee, subcutaneous tissues were dissected, the joint capsule incised with a medial parapatellar retinacular approach, and the articular surfaces of the knee joint exposed. The ACL was then completely transected with a blade (Fig. 1). An intraoperative Lachman test was performed to verify that anterior instability had been created before closing the capsule with 2-0 Vicryl (Ethicon Inc. Somerville NJ) and the skin with 4-0 Vicryl. Previous studies have shown that this method produces a reliable and reproducible degradation of articular cartilage after 9 weeks [15,16]. All rabbits were allowed normal cage activity after the procedure in a temperature-controlled environment with a 12 h light-dark cycle. Intramuscular buprenorphine was administered for at least 72 h for postoperative pain control.

2.3. LED therapy device and treatment

Five weeks following ACLT, LED exposure commenced at intervals of 10 min per day, 5 days per week for 5 weeks, under awake and alert conditions without sedation, in the experimental group (n = 7). The light therapy was accomplished on the experimental knees with a very light custom-designed brace fitting comfortably over the knee and was held in place with two Velcro straps (3 M, St. Paul MN). The device applied two sets of LEDs, with wavelengths of 630 nm (red) and 870 nm (infrared: IR). LEDs mounted to the underside of the brace covered 44 cm², and could treat the entire area of the rabbit knee joint (Fig. 2). It alternated at high frequency between two patterns of off and on diodes. The amounts of energy delivered to the skin were ~2 J/cm² (red) and 2.5 J/cm² (IR) (Light Sciences Oncology, Inc., Snoqualmie WA).

2.4. Harvest tissues

Ten weeks following ACLT, and 3 days after the 5-week treatment, the treated animals (n = 7) were sacrificed with intravenous injections of 97.8 mg/kg sodium pentobarbital via a lateral ear vein following anesthesia. In the control group without any treatment (n = 7) the animals were sacrificed at 9 weeks following ACLT in a similar manner.

2.5. Gross morphology

The rabbit femurs were harvested and initially put in a phosphate buffered solution (PBS) with protease inhibitors: 2 mM disodium-ethylenediaminetetraacetic acid (Na₂-EDTA), 1 mM phenylmethanesulphonyl fluoride (PMSF), 5 mM benzamidine hydrochloride (Benz-HCl), and 10 mM N-ethyl maleimide (NEM) for 1 h. The femoral condyles were then stained with india ink (Higgins Waterproof Drawing Ink, Sanford Co of Bellwood IL). The ink was applied to the surfaces of the femoral condyles with a small paintbrush and a quick rinse of sterile water. Images of the condyles were taken before and after staining with a digital camera (Nikon DX 5 megapixel) for analysis. These images were assessed for residual ink remaining on the cartilage surface by three blinded individuals. A modified Outerbridge classification score was used to grade each condyle. The score criteria consisted of: Grade 1 (intact surface) – surface appears normal and does not retain ink;
Grade 2 (minimal fibrillation) – site appears normal before staining but retains ink as elongated specks or light gray patches; Grade 3 (overt fibrillation) – the cartilage is velvety in appearance and retains ink as intense black patches; or Grade 4 (erosion) – loss of cartilage exposing the underlying bone. Using these grades, an average for each group was calculated [15,17,18].

2.6. Reverse transcriptase-polymerase chain reaction (RT-PCR) technique

Anabolic and catabolic pathways in the osteoarthritic joints were also examined. Articular cartilage of the lateral femoral condyle and synovial tissue around the knee joints was shaved and the mRNA harvested using RNeasy Mini Kit (Qiagen Inc., Valencia CA).

The mRNA expressions of various markers were evaluated by RT-PCR using glyceraldehyde-3-phosphate dehydrogenase (GAPDH) as an internal control gene, type II collagen and aggrecan as anabolic markers of articular cartilage, interleukin-1 beta (IL-1β) and tumor necrosis factor-alpha (TNF-α) as inflammatory markers, and matrix metalloproteinase-3 (MMP-3) and MMP-13 as catabolic markers of cartilage. Based on the published sequences, PCR primers were constructed (see primer sequences: Table 1).

National Institute of Health (NIH) image analysis software (version 1.61, Natl Inst Health, Bethesda MD) was used to quantitatively scan RT-PCR profiles following agarose gel electrophoresis and ethidium bromide visualization. The relative mean density was measured over a fixed gray scale range after correction for background. All values were normalized to GAPDH, data expressed as mean mRNA expression ± standard deviation (SD), and evaluated with a two-tailed Student’s t-test assuming equal variances (statistical significance $P < 0.05$).

3. Results

The attached LED knee device alternated between a wavelength of red (630 nm) and IR (870 nm). The peak irradiance of red is 20 mW/cm² and IR is 25 mW/cm². This device applied energy in pulsed modes with a duty cycle of 16%. The average irradiance was then 3.2 mW/cm² for red and 4 mW/cm² for IR. With a daily irradiation time of 10 min, the daily energy dose applied of red is 2 J/cm² and IR 2.5 J/cm². Because the treated area is 44 cm², the total daily dose of red is 88 J and of IR is 110 J. Rabbits of both groups showed no remarkable knee joint swelling or infective symptoms. Compared to the control group, there were no remarkable side effects seen on the knee joints in the experimental group.

3.1. Gross morphology

The interobserver differences in morphological grading were not significant. Macroscopically, the control group presented with four Grade II, two Grade III and one Grade IV condyles (average was 2.6) (Fig. 3). The femurs in Grade II showed slight fibrillation in both medial and lateral condyles. Grade III femurs showed slight fibrillation in the lateral condyles and overt fibrillation in the medial condyles. These degenerative patterns were more pronounced in Grade IV condyles where the overt fibrillation in the medial condyle and articular cartilage defects were present in the lateral condyles (Fig. 4).

<table>
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<th>Table 1</th>
<th>Primer sequencing.</th>
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<tr>
<td><strong>GAPDH</strong>: Sense 5’-TCA CCA TCT TCC AGG AGC GA-3’</td>
<td>Anti sense 5’-CAC AAT GCC GAA GTG GTC GT-3’</td>
</tr>
<tr>
<td><strong>Type II collagen</strong>: Sense 5’-AAC TGG TGG AGC AGG AAG AG-3’</td>
<td>Anti sense 5’-CTG CAC CAC GGT ATA GGT GA-3’</td>
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<tr>
<td><strong>Aggrecan</strong>: Sense 5’-GAG GTG GTC ATG AAA GGT GT-3’</td>
<td>Anti sense 5’-AGG TCT GTG TAC CGC AGC AG-3’</td>
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<tr>
<td><strong>MMP-3</strong>: Sense 5’-GCC AAG AGA TGC TGT TGA TG-3’</td>
<td>Anti sense 5’-AGG TCT GTG AAA GCC GTG TG-3’</td>
</tr>
<tr>
<td><strong>MMP-13</strong>: Sense 5’-AAC GAG CAT CAT GTG ATG CG-3’</td>
<td>Anti sense 5’-TGG GCC AGG AGG AAA AGC GTG AG-3’</td>
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<tr>
<td><strong>IL-1β</strong>: Sense 5’-TGC AAC ACC TGG GTG CAT CAC TA-3’</td>
<td>Anti sense 5’-GCC CAC ACC TAT CTC GTC GT-3’</td>
</tr>
<tr>
<td><strong>TNF-α</strong>: Sense 5’-ATG GTG ACC CTC AGA TCA GC-3’</td>
<td>Anti sense 5’-TGA CCT TGT TCG GGT AGG AG-3’</td>
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Fig. 3. Results of macroscopic findings: There were four Grade II, two Grade III and one Grade IV in the control group. There were one Grade I, and five Grade II in the experimental group.
In the experimental group, after 5 weeks of treatment with the light brace, there were two Grade I and five Grade II (average 1.7) knees. For the Grade II condyles, one had minimal fibrillation only in the medial condyle and another had minimal fibrillation only in the lateral condyle of the femur. The remaining three Grade II knees showed minimal fibrillation in the medial and slight fibrillation in the lateral femoral condyles. There were two out of seven rabbits without any fibrillation in the femoral condyles (Grade I), and no severe osteoarthritic joints in the experimental group (Grades III and IV) as seen in Fig. 5.

3.2. RT-PCR

Articular cartilage from the femurs of both experimental and treated knees showed no difference in mRNA expression of aggrecan. Type II collagen expression, however, was observed to be increased in the experimental group compared to control \( (P=0.01) \). The expressions of MMP-3, MMP-13, and IL-1\( \beta \) in the cartilage showed no differences between the groups, however TNF-\( \alpha \) expression was decreased in the experimental group compared to control \( (P < 0.01) \) (Fig. 6).
In the synovial tissue the expressions of MMP-3, MMP-13, and IL-1β showed no differences between groups. There was a decrease, however, in TNF-α expression of the experimental group compared to control ($P < 0.01$) (Fig. 7).

4. Discussion

OA is one of the most common diseases affecting articular cartilage. While there are several surgical options for OA, there are relatively few conservative, noninvasive procedures effective at treating OA clinically. Preserving and producing cartilage noninvasively is desirable, especially for early OA. ROM, muscle stretching and strengthening are the first steps for treating early OA as a noninvasive treatment. These exercises may include stretching and strengthening. Initial reports demonstrated that the right light therapy can reduce pain and swelling and increase ROM in early OA knees. While there is still a need to examine the detail of the parameters of wavelength, dose, intensity, irradiation time and continuous or pulsed modes. De Morias et al. [28] examined the anti-inflammatory effect of low-level laser therapy (LLLT) and LED therapy for arthritic knee joints. They concluded that LLLT could reduce inflammatory signs more effectively than LED therapy, however they indicated that new inquiries should be carried out to evaluate the effectiveness of LEDs with different parameters.

Different wavelengths have different chromophores and can have various effects on tissues. In general, LEDs of longer wavelength can penetrate deeper into tissues. Red light has also been shown to be more successful at affecting deeper targets than smaller wavelengths [29–31]. The red and infrared wavelengths applied through LED therapy were adequate for knee joint tissue (i.e. cartilage and synovial tissue) of the rabbit. The effect of the cellular response to pulsed wave versus continuous wave has not been fully evaluated. Because the pulsed wave was thought to stimulate more collagen production than continuous wave, the pulse wave was applied [14]. This may have contributed to the increased expression of type II collagen. If the light intensity is lower than the physiological threshold value for a given target, it does not produce photostimulatory effects, even when irradiation time is extended [14]. Therapeutic outcomes are dependent upon these parameters, making it necessary to find the optimal LED and dose for the intended treatment. The treatment group responded with our applied intensity levels and wavelengths, while the nontreatment groups demonstrated significantly different outcomes.

This study has examined the effect of LED irradiation on OA knees. While there is still a need to examine the detail of the function and mechanism, as well as the optimal conditions for articular cartilage application, there appears to be a capacity to accelerate the regeneration of articular cartilage with LED therapy. Therefore, this treatment may offer the benefit of an alternative, noninvasive procedure to treat OA joints at outpatient clinics.

5. Conclusions

With LED light therapy, the in vivo cartilage surface was preserved macroscopically and the mRNA expression of type II collagen was upregulated in the treated group. Inflammation levels appeared to decrease in both cartilage and synovial tissues at the mRNA level, and LED light therapy shows a potential for anti-inflammatory function which may preserve and potentially repair articular cartilage. As such, this procedure deserves more study as a noninvasive treatment of OA joints clinically.

Disclosure of interest

The authors declare that they have no conflicts of interest concerning this article.

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References


