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Treatment time of ultrasound therapy interferes with the organization of collagen fibers in rat tendons

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ABSTRACT | Background: The application time of therapeutic ultrasound is an infrequently studied dosimetric variable that affects tissue repair. **Objectives:** The aim of this study was to evaluate the effects of different treatment times of therapeutic ultrasound (US) on the organization of collagen fibers in the tendons of rats. **Method:** Forty Wistar rats were selected (300±45 g), and the rats were divided into five groups (n=8 for each group): Control, without tenotomy or any treatment; tenotomy group, with tenotomy and without treatment; US groups (US1, US2, and US3), subjected to tenotomy and treated with US for one, two, or three minutes per area of the transducer, respectively. The animals were sacrificed on the 12th post-operative day, and the tendons were surgically removed for analyses of the collagen fiber organization by means of birefringence analysis. **Results:** The collagen fibers exhibited better aggregation and organization in the US3 group compared with the tenotomy group ($p<0.05$). **Conclusions:** The findings suggest that US applied for three minutes per treated area improves the organization of collagen fibers during rat tendon repair.

Keywords: therapeutic ultrasound; Achilles tendon; physical therapy; collagen; wound healing.

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● Introduction

The healing process of tendon injuries can take months, although the use of electrotherapeutic resources may help accelerate recovery and prevent functional complications that might otherwise delay the rehabilitation process¹. The high incidence of these injuries justifies more studies to improve tendon repair by reducing recovery time and the time to return to functional activities². Researchers have studied non-pharmacological treatment modalities to accelerate tissue repair, including therapeutic ultrasound (US) and low-intensity laser therapy^{1,3-6}.

US therapy has been used to treat musculoskeletal injuries, particularly in tissues with a high percentage of collagen fibers^{4,7-12}. Harvey et al.¹³ described the following physiological responses involved in soft tissue repair when the tissue is submitted to US: acceleration of inflammatory responses by promoting the release of histamines, macrophages, and monocytes; increases in cellular metabolism and collagen synthesis; and decreases in edema and pain¹⁴⁻¹⁷.

Studies regarding the efficacy of US therapy demonstrate that there are no precise guidelines for its parameters, particularly with respect to the dose-response of the treatments¹⁸⁻²¹. Most professionals justify their use of US therapy with their clinical experience^{18,22-24}.

The dose-response of therapeutic US is influenced by many variables, including frequencies, intensities, irradiation times, application modes, type and coupling techniques, and early post-injury interventions²⁵⁻²⁸. A systematic review on the effectiveness of US therapy in musculoskeletal injuries found no evidence of its effectiveness. Warden and McMeeken¹⁹ concluded that US is often used by physical therapists in sport rehabilitation, but there is no evidence to support the dosimetry used. Robertson¹⁸ noted that a therapeutic window ranging from 0.16 to 0.5 W/cm² in pulsed mode, suggested by previous studies, has proven to be illusory; the variables analyzed in the few studies available, such as frequencies, duration of treatments, types and depths of the injury, the effective radiation

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area, and the size of the treated areas, make comparisons impossible. These considerations are corroborated by the findings of Speed²⁹, who supported the exclusion of US therapy due to a lack of evidence.

The US application time influences the amount of energy applied to the tissue, which is calculated by the following formula: Energy (Joules) = Power (Watts) * Time (Seconds)³⁰. The treatment time is an infrequently studied variable in US therapy. The relationship between the time of application and the treated area has been described by Oakley³¹. Oakley³¹ proposed that each area of $1\frac{1}{2}$ times the size of the transducer should be treated for one to two minutes. Furthermore, for each area adjacent to the transducer, an additional one to $1\frac{1}{2}$ minutes of treatment was advised, although these relationships were not grounded or justified. Hoogland³² recommended a maximum treatment time of 15 minutes and at least 1 minute for each treated area. Conversely, Olsson et al.²⁴ concluded that there was no definitive recommended treatment time for US application.

Experimental studies on tendon repair using ultrasound follow the recommendation of 1-2 minutes per transducer treated^{6,33,34}. The rationale for this treatment is based on previous studies using the same application time or following the recommendations of Oakley³¹ and Hoogland³², who proposed this time empirically. There are no studies in cell culture, animal tendons or human tendons concerning different US application times. Thus, there is no evidence to justify the choice of this dosimetric parameter for tendon healing. The effects of irradiation time as a function of frequency and depth of penetration has been evaluated in human muscle tissue by Draper et al.²⁵. There is a good basis for US application in individuals with little adipose tissue overlying the muscle. The effect of treatment time on tendon healing needs to be studied to improve the efficiency of ultrasound application.

Therefore, the aim of this study was to evaluate the effects of different US therapy treatment times on the organization of collagen fibers in rat tendons.

● Method

This study was approved by the ethical research board of the Faculty of Medicine of the Universidade de São Paulo (USP), São Paulo, SP, Brazil (nº 065/11 04/13/2011). The surgical procedures followed the ethical guidelines for animal experiments of the Council for the International Organization of Medical Sciences, the standards of the Brazilian Society of

Laboratory Animal Sciences, and the current national legislation on procedures for the scientific use of animals in research (Federal Law 11,794, October 9th, 2008).

Animals

Forty male Wistar rats were used in this study (weight: 300 ± 45 g, age: 90 days). The animals were kept in a controlled environment at a temperature of 25°C and a light/dark cycle of 12 hours and were provided with a balanced diet for rodents and water *ad libitum*. The rats were randomly divided into five groups of eight rats. The number of animals needed to achieve statistical significance was based on previous studies^{35,36}.

The animals were divided into the following groups: control group, without surgery and US therapy; tenotomy group, with tenotomy of the Achilles tendon and without US therapy; US1 group, with tenotomy of the Achilles tendon and US treatment for one minute per transducer area, totaling two minutes of treatment; US2 group, with tenotomy of the Achilles tendon and US treatment for two minutes per area of the transducer, totaling four minutes of treatment; and the US3 group, with tenotomy of the Achilles tendons and US treatment for three minutes per area of the transducer, totaling six minutes of treatment.

Experimental model

The model for the total tenotomy of the middle portion of the Achilles tendon (transverse incision of the dissected tendon) was based upon experiments performed by Cunha et al.¹⁴, Reddy et al.³⁷, Koeke et al.³⁸, and Carrinho et al.³⁹.

Surgical technique

The animals were weighed and anesthetized with veterinarian tiletamine hydrochloride and zolazepam hydrochloride (Zoletil 50 - VIRBAC®) at a dose of 25 mg/kg of body mass via intraperitoneal injections⁴⁰. The posterior areas of the right tibia, corresponding to the location of the Achilles tendon, were epilated. Afterwards, asepsis was achieved with alcohol 70%, and the skin and the panniculus carnosus were cut longitudinally over the site corresponding to the middle third of the Achilles tendon with a scalpel blade, exposing the tendon to obtain a cross-section²² (Figure 1). After sectioning the tendon, the skin incision was completely sutured in the central area using 4.0 nylon thread (Figure 2). Post-operatively, the animals received a single intramuscular dose



Figure 1. Longitudinal incision and exposure of the Achilles tendon on the posterior of the rat's hind limb.



Figure 2. Surgical incision sutures with 4.0 nylon surgical thread.

of Ceftriaxone in the contralateral hind limb for antibiotic prophylaxis and were returned to sterilized cages without immobilization^{14,37-39,41}. The right Achilles tendon was submitted to US in the US1, US2 and US3 groups.

Treatment protocols

The Sonacel BIOSET® US emitter was used. The design of its transducer, which has an area of 0.5 cm², was modified to fit the posterior portion of the rats' hind limbs and the size of the lesions. The equipment was calibrated with an acoustic scale (GT210 Glutymax®) and an oscilloscope (Intermetro®).

The animals were submitted to the first application of US therapy 24 hours after the surgery. The ultrasonic irradiation employed the following parameters: 1 MHz frequency, pulsed mode with 20% of the duty cycle (2 ms emission/8 ms interval), 100 Hz repetition frequency, 0.5 W/cm² of intensity (spatial average time average - SATA), and 0.5 cm² ERA and was performed once per day. The treated area was 1 cm² in size. The animals were sacrificed after the 10th treatment session on the 12th post-operative day. The irradiations occurred consecutively at one day intervals after the fifth treatment day^{3,4,18,38,42}.

During the US applications, the transducer was placed perpendicular to the treated area using movement techniques. Water-soluble gel was used as a coupling agent to better conduct the waves and facilitate movement over the rat's skin^{27,43}. The animals were also stabilized with a standard retainer (Figure 3)^{14,38}.



Figure 3. Constraint and irradiation of the Achilles tendon on the posterior of the rat's hind limb

Preparation and analysis of the histological slides

The animals were sacrificed using a CO₂ chamber, and the damaged tendon areas were surgically removed. For the quantitative assessment of tendon repairs, the removed tendons were placed in a 10% formalin solution for 24 hours and were subsequently dehydrated with four alcohol baths of one hour each at dilutions of 50%, 70%, 80%, and 100%. Subsequently, the samples were washed in xylene for one hour and embedded in paraffin at 58°C. The blocks were longitudinally cut into serial sections using a microtome (LAICA®) with a standard thickness of 7 µm. The sections were mounted on glass slides without staining and identified by a blinded evaluator.

The uneven penetration of light through an object is a measure of birefringence. This measure assesses the density and organization of the analyzed material and is used to evaluate the organization and structure of the collagen fibers. The purpose of this procedure

was to analyze the organization, the aggregation states, and the alignment of the tendon collagen fibers by measuring their birefringence^{14,20,39}. For the birefringence analyses, the slides for each group were immersed in distilled water (refractive index $\eta=1.333$) for 30 minutes. The optical retard (OR) was measured using a polarized light microscope (LEICA®) with a 10X/0.22 objective, pol. 0.9 condenser, Senarmont $\lambda/4$ compensator, and monochromatic light ($\lambda=546$ nm) obtained by an interference filter (LEICA®) of the LAMAV (Vitreous materials laboratory, Department of material Engineering of UFSCar). These types of analyses have been used to quantitatively measure the degrees of organization of the collagen fibers in several studies^{14,20,38}. The resulting measures, in degrees, were converted into nanometers (nm) by multiplying the values by 3.03. The total birefringence of the collagen fibers was measured after soaking the tendons in distilled water. To perform the measurements along the axis of the tendon, the longitudinal axis of the collagen fibers was oriented 45° from the direction of the propagation of the transmission light. In this position, the collagen fibers have the highest OR. Measurements were taken at five different points on the central areas of

the tendons, which corresponded to the areas of the injuries^{14,20,38,39}. The birefringence data were collected by two previously trained, independent, and blinded assessors.

Statistical analyses

The data distributions were tested using the Kolmogorov-Smirnov test. Subsequently, the data were analyzed using the Kruskal-Wallis test, followed by Dunn’s *post-hoc* test to investigate differences between the groups with a significance level of $\alpha<0.05$.

Results

Kruskall-Wallis tests revealed statistically significant differences among the groups with respect to the OR of the five different areas of each injury region ($p<0.0001$). Dunn’s multiple *post-hoc* tests demonstrated statistically significant differences when comparing the OR data between the following groups: US1xUS2, US1xUS3, US1xC, US2xC, US3xT, and CxT (Figure 4).

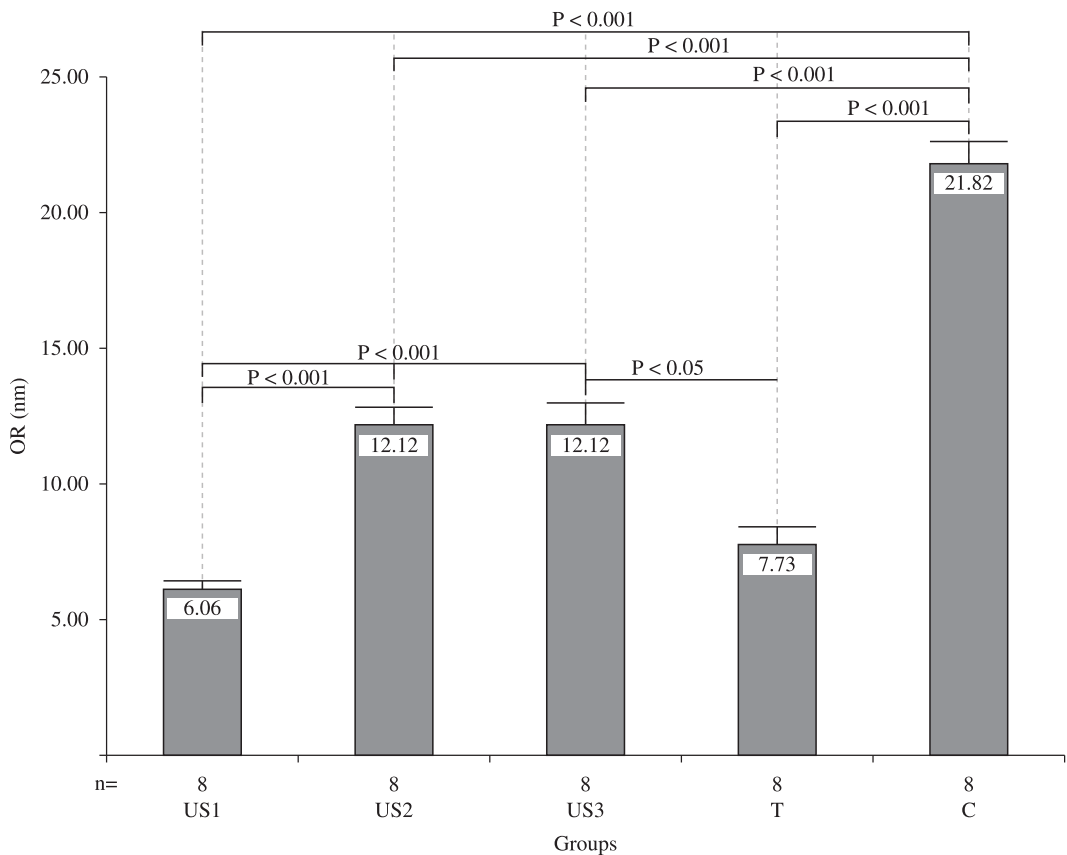


Figure 4. Median and standard errors of the optical delay values for the experimental groups.

Qualitative histological analyses using polarized light microscopy allow the definition of normality standards regarding the organization of the collagen fibers (Figure 5). When comparing the images in Figure 5, it is possible to observe that the animals of the US3 group demonstrated better organization of the collagen fibers when compared to the other treated groups (US1 and US2). It is also possible to observe that the collagen fibers of the animals in the US1 and tenotomy groups were not organized.

● Discussion

The aim of this study was to evaluate the effects of different US application times on tendon repair in tenotomized rats, and the results suggested that application times of less than three minutes were insufficient to promote adequate organization of the collagen fibers. The literature also does not provide clear definitions regarding US therapy treatment times. US doses are the most frequently studied

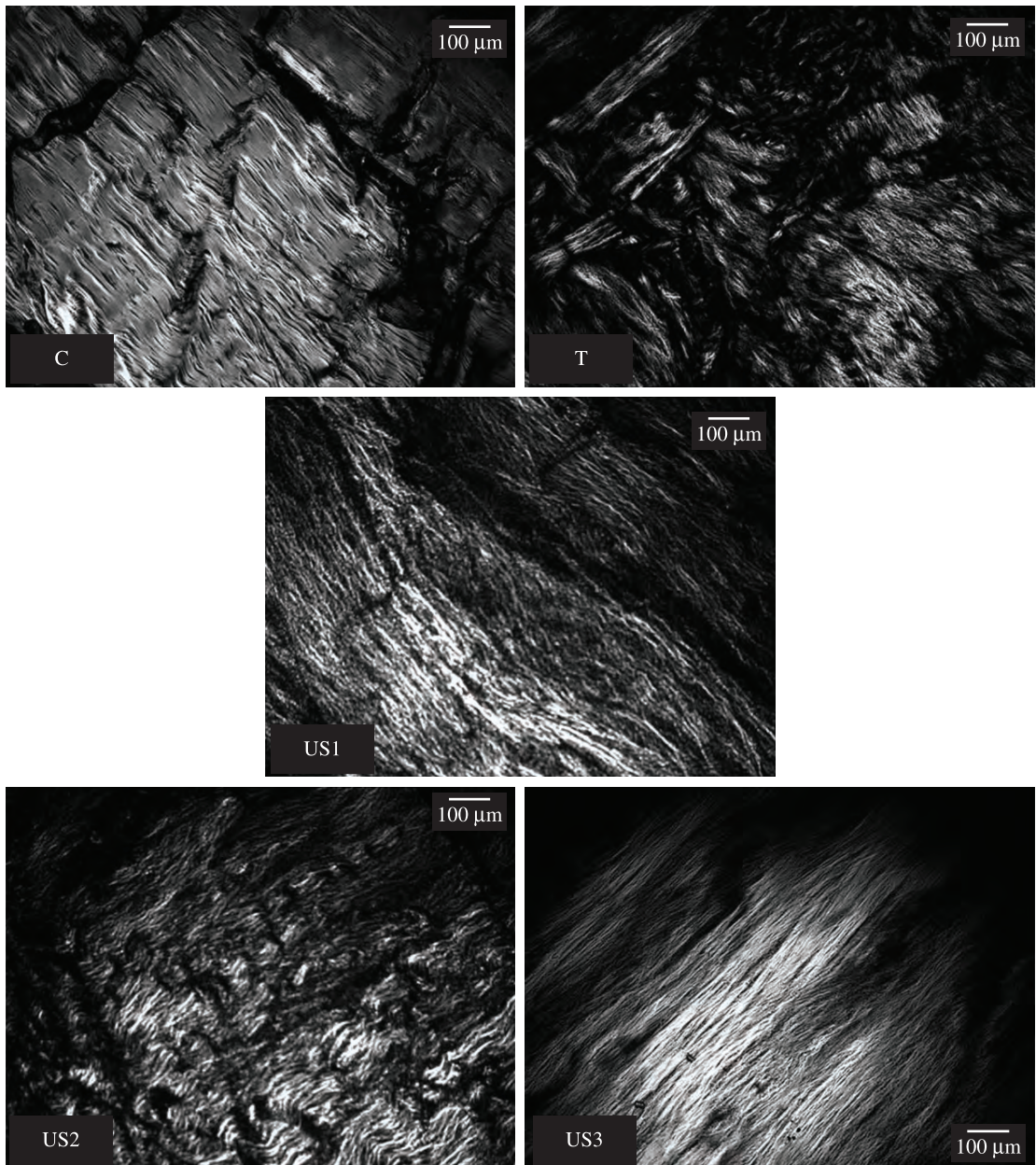


Figure 5. Images related to the qualitative observations of the birefringence analyses of the tenotomized rat tendons. Specimens were positioned with the longitudinal axes of the collagen fibers at 45 degrees. In the US1, US2 and tenotomy (T) images, deficiencies in the organization of the collagen fibers can be observed. The US3 image demonstrates better organization of the collagen fibers, similar to that of the control group (C).

variables^{15,30}, and there is evidence that lower doses are more effective for tissue repair⁴⁴.

The dose standards employed in the present study were those most often used in US therapy studies, and the treatment times were the studied variables. The intensity (0.5 W/cm²) was based upon the research of Carvalho et al.⁴, Cunha et al.¹⁴, Reddy et al.³⁷, Koeke et al.³⁸, and Silva et al.⁴⁵. The form of ultrasonic pulsed emission was chosen based on research conducted by Carvalho et al.⁴, Cunha et al.¹⁴, Frasson et al.¹⁷, Blume et al.²², Koeke et al.³⁸, and Belanger et al.⁴⁶. The frequency of the equipment was justified by the experiments by Carvalho et al.⁴, Piedade et al.¹¹, Cunha et al.¹⁴, and Koeke et al.³⁸.

The treatment duration is known to depend upon the area of the injury. Oakley³¹ recommended that the treatment time be related to the treated area, with one or two minutes for each area that is 1½ times the size of the transducer. According to Oakley³¹, subsequent treatment times can be increased from 1 to 1½ minutes. This subsequent treatment time is based upon empirical studies or clinical observations. There are no clinical or experimental studies that support these findings, and this lack of information concerning treatment should be considered an issue for further study. The results of the present study do not corroborate the results of Oakley³¹ regarding the application times needed to promote better alignment of collagen fibers in rat tendons.

The application time is an important variable to be considered in US therapy dosimetry and determines the amount of energy applied to the treated tissue. Alexander et al.³⁰, in a systematic review of the use of US in various shoulder pathologies, noted that studies that employed the average energy of 4.2 J per treatment session were clinically effective, whereas those that applied an average of 2.019 J were not successful. The duration of application may have influenced these data.

The components of the extracellular matrix of the connective tissues present in tendons, particularly the collagen fibers, exhibit viscoelastic properties that depend on the amount of time the tissue is exposed to deformations or strains to keep their morphofunctional characteristics⁴⁷. There is a direct relationship between the alignment of collagen fibers of the extracellular matrix and the biomechanical properties of tendon: when the collagen fibers exhibit better organization, the flexibility and strength of these tendons also improve⁴⁸.

Thus, three minutes of US application could have sufficient biomechanical effects, compression and/or

tension to promote the mechanisms of cellular and matrix extracellular signaling for the realignment of the collagen fibers⁴⁹.

In this study, the OR in the tendon scar tissues of rats demonstrated statistically significant differences when 18 J of energy (group US3) was applied to the affected area. However, when other magnitudes of energy, namely, 6 J (US1 group) or 12 J (group US2), were deposited on the treated area, no differences were observed regarding the OR between the experimental and control groups.

The most commonly employed application time is five minutes, and the US is generally applied to an area twice the size of the transducer, resulting in an application time of 2.5 minutes per treated area^{6,38}. The size of the treated area is a variable that has not been described in studies related to US therapy and tendon repair. Wood et al.⁶, Cunha et al.¹⁴, and Koeke et al.³⁸ applied US therapy to an area of 1 cm², but these data were not described in the articles and were instead personally provided by the authors. The present study demonstrated that longer application times are needed to promote better alignment of the collagen fibers. One hypothesis to explain the differences between the results of this and previous studies is that the control of the treated areas may not have been accurately measured, making it impossible to compare findings. This variable must be considered when designing the experiment so that comparisons between studies can be made.

The relevance of studying and developing techniques to treat tendon injuries relates to the long recovery times and the resulting functional disabilities of tendon injuries, which may take weeks or even months to fully recover^{1,3,50}. The tensile strength of the tendons is promoted by the alignment of their collagen fibers and by the types and amount of fibers. Thus, this study used an experimental model already defined in the literature, described by Enwemeka and Reddy¹ and reproduced by other authors^{14,37-39,41}, that exhibits easy reproducibility, lower costs, and high levels of reliability.

The levels of collagen fiber organization in the injured area are measured according to the OR in the polarized light microscope, which provides quantitative data. The brightness of the birefringence image reveals the level of aggregation of the collagen fibers¹⁴. During the process of tissue repair, the OR values tend to decrease³⁹, corroborating the results of the present study, in which the US1 group demonstrated the lowest OR values, followed by the tenotomy, US2, US3, and control groups.

The findings of the present study suggest that three minutes of application per area of the US transducer results in positive effects on the collagen fiber organization in rat tendons during the tissue repair process, producing significant changes in the collagen optical retards. Shorter application times produced no significant changes in the optical delays when compared to the tenotomy group.

The transposition of the experimental findings to the clinical application of therapeutic ultrasound on tendon healing should be evaluated with respect to the difference between the application times of 2 and 3 minutes. Further studies are needed to assess the difference between these treatment times with regard to human healing.

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