Tendon cell ciliary length as a biomarker of *in situ* cytoskeletal tensional homeostasis

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Summary

To determine if tendon cell ciliary length could be used as a biomarker of cytoskeletal tensional homeostasis, 20 mm long adult rat tail tendons were placed in double artery clamps set 18 mm apart to create a 10% laxity. The tendons were allowed to contract over a 7 day period under culture conditions. At 0, 1, 5, and 7 days the tendon cell cilia were stained and ciliary length measured using confocal imaging. There was a significant (p<0.001) increase in ciliary length at 1 day. At day 5 (when the tendon became visibly taut) there was a significant (p<0.001) decrease in ciliary length compared to day 1. By day 7 the tendon remained taut and ciliary length returned to day zero values (p=0.883). These results suggest that cilia length reflects the local mechanobiological environment of tendon cells and could be used as a potential *in situ* biomarker of altered cytoskeletal tensional homeostasis.

KEY WORDS: tendon, cilia, recalibration, tensional homeostasis, marker.

Introduction

Tendon cells are capable of generating a tensional load within their cytoskeleton through an α-smooth muscle actin-mediated mechanism, which can generate a considerable traction force against the cell’s extracellular matrix. This cell-based contraction produces a cytoskeletal tensional homeostasis or set point, that calibrates the cell’s response to changing extracellular matrix strain through the regulation of gene expression. Loss of this cellular tensional homeostasis is thought to occur in tendon cells when collagen fibrils associated with those cells are damaged, or become lax, and lose the ability to bear load. This loss of mechanobiological homeostasis has been shown to stimulate catabolic processes within tendon cells and has been postulated as a causative factor in the development of tendinopathy. The ability to identify *in situ* alterations in the mechanobiologic environment of tendon samples may provide further insight into the role of mechanobiology in the development and progression of tendinopathy.

Our lab has demonstrated that loss of tensional homeostasis in tendon cells is accompanied by an increase in the length of the cells’ primary cilia. Primary cilia are sensory organelles which detect and transmit mechanical and chemical information between the extracellular environment and the cell. Tendon cell cilia have been shown to adapt their length to changes in their local loading environments; increasing their length in response to decreased mechanobiologic stimulus and shortening to their resting homeostatic length following application of externally applied cyclic loads. However, the ability of intrinsically generated (i.e. cell-based) tensional homeostasis to regulate ciliary length in tendon cells is unknown.

Therefore, the purpose of the current study was to determine if elongated tendon cell cilia in lax rat tail tendon fascicles (RTTfs) can be returned to control lengths through a previously described actin-based cellular contraction mechanism. A previous study has demonstrated that tendon cells are able to actively contract RTTfs with a 10% laxity over a 7 day period and recover normal tendon cell tensional homeostasis as determined by inhibition of collagenase synthesis. In the current study, we hypothesize that the primary cilia of RTTfs tendon cells will significantly increase in length following creation of a 10% laxity in the tendon. In addition, we hypothesize that the length of preliminary cilia will return to control values following the cell-based contraction of the tendon over 7 days and the return of the previously documented tensional homeostasis. Validation of these hypotheses could support the validity of ciliary length as a potential biomarker of the *in situ* mechanobiologic status of tendon cells.

Methods

Following Institutional Animal Care and Use Committee approval, tendons were obtained from the tails of
adult 3-month-old Sprague-Dawley rats immediately after euthanasia and maintained in DMEM supplemented with 10% FBS, 1% antibiotic/antimycotic and 0.75% ascorbic acid while incubated at 37°C and 10% CO2. Twenty-four rat tail tendon fascicles (RTTs) (20 mm long) from each of 5 animals were placed in double artery clamps set 18 mm apart so as to create a grip to grip laxity of 10% of total RTT length. The RTTs were divided evenly into groups and analyzed at one of four different time periods as follows: fresh control (Day 0) or allowed to freely contract between the clamps for 1 day (Day 1), 5 days (Day 5), or 7 days (Day 7). A previous study investigated the re-establishment of tensional homeostasis in lax tendons utilizing younger 1-month-old rat tail tendons at 3 and 5 days13. The tendons used in this study were from older 3-month-old rats that have been shown to have a slower rate of cell-based tendon contraction than 1-month-old rats17. Therefore, in the current study the later time points of 5 and 7 days were used as this corresponds to when the tendons became taut between the clamps for 3-month-old rats. One centimeter segments from each of the six RTTs were harvested per treatment group and stained with 250 nM Tubulin Tracker Green (Life Technologies) for 30 minutes at 37°C. Stained RTTs were washed vigorously in dPBS for 30 minutes, then placed into two subsequent non-vigorous dPBS washes for a total of 3-5 hours of washing. Cilia were visualized using Pro Long Gold with DAPI (Life Technologies) and confocal images were created using an Olympus Fluo view 1000 laser scanning microscope. The length of each primary cilia, from the centriole to the tip of the ciliary axenome, was measured using Image J as previously performed11. Cilia lengths were presented as the average and standard deviation of the 5 rats and analyzed using a two factor ANOVA and Tukey’s post hoc test with significance set at p< 0.05.

Results

The average length of the primary cilia in tendon cells of fresh (Day 0) rat tail tendon fascicles (RTTs) was 1.35 ± 0.11 µm. One day following creation of a 10% laxity in their grip to grip length the RTTs remained visibly lax and the length of the primary cilia significantly increased (p<0.001) to 2.76 ± 0.19 µm (Figs. 1, 2). By day 5 the RTTs appeared visually taut between the clamps and the cilia had significantly increased.

![Figure 1. Confocal images of in situ tendon cells with associated nuclei (DAPI) and cilia (Tubulin Tracker Green) from tendons allowed to freely contract between clamps at Day 0, Day 1, Day 5, and Day 7 with the tendon remaining lax at Day 1, but visibly taut at Day 5 and Day 7.](image1)

![Figure 2. Graph showing the average cilia length of in situ tendon cells from tendons allowed to freely contract between clamps at A) Day 0, B) Day 1, C) Day 5, and D) Day 7 with the tendon remaining lax at Day 1, but visibly taut at Day 5 and Day 7. Lines above bars indicate significantly different values, p<0.05.](image2)
(p<0.001) shortened (1.65 ± 0.19 µm) with respect to day 1 lengths. The RTTs remained taut at day 7 and had an average length of 1.40 ± 0.13 µm which was not significantly (p=0.883) different from 0 day (fresh) samples (Figs. 1, 2).

Discussion

The ability of tendon cells to maintain their normal cytoskeletal tensional homeostasis has been shown to play a critical role in regulating catabolic gene expression and protein synthesis5,6,8,10. Loss of cytoskeletal homeostasis may occur as a result of an absence of the normal, residual tension in the pericellular matrix of tendons and ligaments caused by injury or repetitive exercise18-20. Such alterations in the local mechanobiologic environment of tendon cells have been implicated as the inciting factor in the pathogenesis of tendinopathy7,21. However, since an upregulation of catabolic factors associated with tendinopathy can also be triggered by other humoral stimuli (i.e. IL-1b, PGE2)22-24 a biomarker which specifically reflects the local, in situ mechanobiologic status of the cell could be helpful in supporting this theory. The primary cilium of tendon cells have been shown to adapt their length to changes in their local loading environments; increasing their length in response to decreased mechanobiologic stimulus and shortening to their resting homeostatic length following application of externally applied cyclic loads11.

The results of the current study demonstrate that cilia elongate within 24 hours following induced loss of normal cytoskeletal tension through tendon laxity. This outcome confirms a previous study which also demonstrated cilia elongation with tendon laxity11. In addition, the same previous study also documented that with continuous loss of cytoskeletal tension, tendon cilial length does not change significantly after the initial 24 hour lengthening11. This “set control length” has also been demonstrated in chondrocyte cilia12 suggesting that the average cilial length would not increase beyond this set length once cytoskeletal tension has been lost. This growth in ciliary length has also been documented in other tissues that have lost external stimuli or internal tension25-28. The precise reason for primary cilia to lengthen in the absence of physical stimuli is unclear. However, it may represent an attempt by the tendon cell to “amplify” the transmission of weakened mechanotransduction signals11,29.

Externally applied forces transmitted through the extracellular matrix have been shown to reverse cilia elongation both in tendon11 and other tissues25-27,29,30. However, the current study is unique as it demonstrates that cilia elongation can also be reversed through internally generated cell traction. This actin-mediated cellular force contracts the extracellular matrix to eliminate the laxity in the extracellular matrix and allows the tendon cells to regain their normal internal tensional homeostasis5. Previous studies from our lab have documented a similar contraction pattern of lax RTTs and a return to cellular homeostasis between 5 and 7 days, as manifested by an inhibition of interstitial collagenase synthesis3,17.

The current study demonstrates that as the cellular tensional homeostasis is restored between 5 and 7 days, the length of the cilia decreases until it returns to a length not statistically different than control values. Since cilial length is reflected by the local mechanobiologic environment of the cell, the findings of the current study suggest that increased cilial length may be an effective biomarker with which to document alterations in the in situ mechanobiologic signaling environment of tendon cells. The histological assessment of cilial length in histologic sections of tendinopathic tissues may help clarify the potential role of mechanobiologic understimulation7 in the etiopathogenesis of the catabolic changes associated with tendinopathy31.

Conclusion

In conclusion, the current study demonstrates the relationship of tendon laxity and tendon cell tensional homeostasis to the length of their primary cilia. In addition, results support the potential use of tendon cell cilary length as a biomarker of the in situ mechanobiologic environment of tendon cells.

References

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