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The Mechanisms Underlying Nonuniform Tendon  
Displacements

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1 A 3D Model of the Achilles Tendon to Determine  
2 The Mechanisms Underlying Nonuniform Tendon Displacements  
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42 ABSTRACT

43 The Achilles is the thickest tendon in the body and is the primary elastic energy-storing  
44 component during running. The form and function of the human Achilles is complex: twisted  
45 structure, intratendinous interactions, and differential motor control from the triceps surae muscles  
46 make Achilles behavior difficult to intuit. Recent *in vivo* imaging of the Achilles has revealed  
47 nonuniform displacement patterns that are not fully understood and may result from complex  
48 architecture and musculotendon interactions. In order to understand which features of the Achilles  
49 tendon give rise to the nonuniform deformations observed *in vivo*, we used computational modeling  
50 to predict the mechanical contributions from different features of the tendon. The aims of this study  
51 are to: (i) build a novel computational model of the Achilles tendon based on ultrashort echo time  
52 MRI, (ii) compare simulated displacements with published *in vivo* ultrasound measures of  
53 displacement, and (iii) use the model to elucidate the effects of tendon twisting, intratendon sliding,  
54 retrocalcaneal insertion, and differential muscle forces on tendon deformation. Intratendon sliding  
55 and differential muscle forces were found to be the largest factors contributing to displacement  
56 nonuniformity between tendon regions. Elimination of intratendon sliding or muscle forces reduced  
57 displacement nonuniformity by 96% and 85%, respectively, while elimination of tendon twist and  
58 the retrocalcaneal insertion reduced displacement nonuniformity by only 35% and 3%. These results  
59 suggest that changes in the complex internal structure of the tendon alter the interaction between  
60 muscle forces and tendon behavior and therefore may have important implications on muscle  
61 function during movement.

62

## 63 INTRODUCTION

64 Tendons are vital for human movement as they transmit forces from muscles to bones. In addition  
65 to this fundamental role, tendons are important for the dynamics of coordinated movement as they  
66 serve to store and release elastic energy during cyclic movements (Alexander and Bennet-Clark,  
67 1977; Alexander, 1991; Ishikawa et al., 2005; Sasaki and Neptune, 2006), protect muscle fibers from  
68 damage due to stretch (Griffiths, 1991), allow muscles to work at favorable power outputs (Cronin et  
69 al., 2013; Lichtwark and Wilson, 2006), and enhance muscle performance by enabling muscle-  
70 tendon-unit performance that exceeds the capabilities of muscles alone (Roberts, 2002). As the  
71 largest and strongest tendon in the human body (Järvinen et al., 2005), the Achilles tendon is  
72 important for walking and running, sustaining loads many times body weight during these tasks  
73 (Giddings et al., 2000).

74         Recent research has demonstrated the complexity of *in vivo* behavior of the Achilles tendon.  
75 Using dynamic ultrasound imaging, Arndt et al. (2012) showed nonuniform tissue displacements  
76 within the Achilles tendon during cyclic passive ankle dorsiflexion. In that study, tissue displacements  
77 were greater in the anterior portion of the tendon as compared to the displacements in the  
78 posterior portion of the tendon, an interesting result considering that the moment arm is greater for  
79 the posterior portion (Fig. 1). Slane and Thelen (2014) observed similar nonuniformity in both  
80 passive and eccentric loading conditions, with less extreme nonuniformity during eccentric trials.  
81 Bojsen-Møller et al. (2004) showed nonuniform displacements between the soleus distal  
82 aponeurosis and the gastrocnemius distal aponeurosis (proximal to the Achilles tendon) during  
83 isometric ankle plantarflexion. Finally, Franz et al. (2015) observed nonuniform displacements of the  
84 Achilles tendon in healthy subjects during normal treadmill walking.

85         What gives rise to these observations of nonuniform displacements? Structurally, the  
86 tendon is composed of three subtendons<sup>1</sup> (Handsfield et al., 2016), each of which is controlled by a  
87 different muscle belly in the triceps surae (Bojsen-Møller and Magnusson, 2015; Edama et al., 2014;  
88 Sarrafian, 1993; Szaro et al., 2009). These subtendons twist about one another in an internal  
89 direction (Bojsen-Møller and Magnusson, 2015; Edama et al., 2014; van Gils et al., 1996; White,  
90 1943), but the functional significance of this twist is not well understood. Additionally, recent work  
91 has suggested that sliding of tendon substructures may occur in healthy and young, but not older,  
92 animals (Thorpe et al., 2013). Interestingly, Slane and Thelen (2015) observed reduced regional  
93 nonuniformity in middle-aged subjects, as compared to young subjects.

94         As an explanation for the nonuniformity observed in the Achilles, Arndt et al. (2012) and  
95 Bojsen-Møller et al. (2004) have proposed differential force contributions from the triceps surae  
96 muscles. Due to the complexity of Achilles behavior and structure such as twisted subtendons, intra-  
97 tendon sliding, and an osseous insertion behind the calcaneus (retrocalcaneal), it is difficult to intuit  
98 how these and other features of the tendon contribute to patterns of deformation. While a useful  
99 tool for dynamic musculoskeletal imaging, ultrasound studies are generally limited to a small  
100 imaging region within some portion of the tendon, making it difficult to appreciate the behavior of  
101 the entire tendon. Taken together, there are questions that cannot be answered from imaging

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<sup>1</sup> Previous authors have used different terminology to describe the portions of the Achilles arising from the distinct muscular heads of the triceps surae. Here, we use the term “subtendon” for this structure. For a brief review and comparison of different terminologies used, see Handsfield et al. (2016).

102 studies alone: *i*) do differential muscle forces give rise to the tissue displacements observed  
103 experimentally? *ii*) what role does inter-subtendon sliding have in creating tissue displacement  
104 profiles? *iii*) what are the mechanical contributions of various features of the Achilles tendon such as  
105 twisting and a retrocalcaneal insertion?

106 The overall goal of this work was to create a finite element model of the Achilles tendon to  
107 investigate the contributions of Achilles tendon geometry, twisting tendon morphology, and  
108 differential muscle forces on tendon displacement profiles. The model geometry was constructed  
109 from images obtained using a customized ultrashort echo time (UTE) MRI sequence. The specific  
110 goals were to use the model to replicate tissue displacements observed in the literature; estimate  
111 muscle force contributions; and use the model to understand the interactions of muscle forces,  
112 subtendon twisting and sliding, and a retrocalcaneal insertion on tendon displacements and strains.

Accepted manuscript

## 113 METHODS

## 114 Ultrashort echo time magnetic resonance imaging

115 A healthy 27 year-old female volunteer (height: 1.63m, mass: 52 kg) provided informed  
116 consent to the following protocol, which was approved by the University of Virginia Institutional  
117 Review Board for Human Subjects Research. We scanned the volunteer using a 3T Siemens  
118 (Erlangen, Germany) Trio MRI scanner with a combination ultrashort echo time (UTE) and short echo  
119 time (shTE) dual-echo pulse sequence utilizing a spoke-radial 3D k-space trajectory which was  
120 originally developed for imaging cortical bone and is fully described elsewhere (Miller et al., 2015).  
121 We placed a flex coil around the subject's right ankle which was manually positioned to 25° of  
122 plantarflexion. We used the following sequence parameters optimized for tendon imaging-  
123  $TE_1(\text{UTE})/TE_2(\text{shTE})/TR/\alpha$ : 0.08ms/2.54ms/6ms/10°. Field of view was 192x192x192 mm<sup>3</sup> and spatial  
124 resolution was 0.8mm isotropic. The 3D radial k-space trajectory consisted of 91,088 individual  
125 spokes, corresponding to 50% undersampling at this field of view and resolution. The shTE  
126 magnitude images were subtracted pixel-by-pixel from inherently co-registered UTE magnitude  
127 images to maximize tendon contrast. To reduce noise, we averaged and resampled images at a slice  
128 thickness of 2.4mm in the axial plane using Osirix imaging software (Rosset et al., 2004).

## 129 Model geometry

130 We manually outlined the Achilles tendon in MR images from the soleus muscle-tendon  
131 junction to the extent of tendon insertion on the calcaneus (Fig. 2) using in-house segmentation  
132 software written in Matlab (The Mathworks, Natick, MA). The control points that defined the tendon  
133 external geometry were exported to Autodesk Inventor Professional (Autodesk Inc, San Rafael, CA).  
134 To reconstruct axial cross-sections of tendon, we interpolated control points with closed cubic  
135 splines. 3D tendon geometry was constructed by lofting over all axial cross-sections.

136 In order to model the Achilles sub-structure, we defined 2D surfaces in Autodesk Inventor that  
137 subdivided the 3D tendon geometry into three subtendons. Each of the three subtendons  
138 corresponded to one of the three muscles of the triceps surae (Fig. 3) (Bojsen-Møller and  
139 Magnusson, 2015; Edama et al., 2014; Sarrafian, 1993; Szaro et al., 2009). At the proximal aspect of  
140 the tendon, we defined the anterior portion as the subtendon extending from the soleus (hereafter  
141 referred to as SOL subtendon), the posterior medial portion as the subtendon extending from the  
142 medial gastrocnemius (MG subtendon), and the posterior lateral portion as the subtendon extending  
143 from the lateral gastrocnemius (LG subtendon), as these are the relative locations of the triceps  
144 surae muscles informed by MRI (Handsfield et al., 2014). At the distal aspect of the free tendon, we  
145 defined the location of each subtendon according to literature reports of dissection studies that  
146 separated the three portions of the Achilles tendon by peeling away the portions of the tendon from  
147 the proximal end of the tendon where the muscles' distal aponeuroses are separate and  
148 distinguishable (Bojsen-Møller and Magnusson, 2015; Edama et al., 2014; Sarrafian, 1993; Szaro et  
149 al., 2009). This positioning resulted in an internal twisting of the Achilles subtendons, which is  
150 consistent with literature (Bojsen-Møller and Magnusson, 2015; Edama et al., 2014; van Gils et al.,  
151 1996; White, 1943).

152 The morphology of the Achilles subtendons in the calcaneal region is not well defined in the  
153 literature; therefore, in the present model, the cross-sectional morphology at the end of the free

154 tendon was maintained through the tendon's insertion on the calcaneus. This morphology is  
155 consistent with sagittal imaging studies depicting collagen fibers with a longitudinal orientation in  
156 the insertional region (Milz et al., 2002) as well as photographs of the posterior collagen fiber  
157 directions of the Achilles tendon (Dalmau-Pastor et al., 2014). The model geometry was transferred  
158 to AMPSolid (AMPS Technologies, Pittsburgh, PA) and meshed into tetrahedral solid elements.

159 Constitutive model

160 Each subtendon was modeled as an incompressible Neo-Hookean solid with strain energy  
161 density function,  $W$ , given by Equation 1:

$$W = C(I_1 - 3) \quad (\text{Equation 1})$$

162 where  $C$  is a material constant and  $I_1$  is the first invariant of the right Cauchy-Green deformation  
163 tensor. The first invariant,  $I_1$ , is defined as the sum of squares of the principal stretches,  $\lambda_1$ ,  $\lambda_2$  &  $\lambda_3$ ,  
164 according to Equation 2:

$$I_1 = (\lambda_1)^2 + (\lambda_2)^2 + (\lambda_3)^2 \quad (\text{Equation 2})$$

165 The material parameter  $C$  was set to a value of 45 MPa for all three subtendons. This parameter  
166 corresponds to a Young's modulus of approximately 270 MPa which is consistent with previously  
167 reported Young's moduli for the human Achilles tendon *in vivo* (Kubo et al., 2002).

168 Model boundary conditions

169 A frictionless contact interface was defined between the three subtendons. This interface  
170 was chosen to simulate subtendon sliding, a property demonstrated in fascicles of healthy energy-  
171 storing equine tendons (Thorpe et al., 2013) that may be relevant for human Achilles subtendons.  
172 While the mechanics of the interfascicular matrix have been explored previously (Thorpe et al.,  
173 2015), the mechanics of the human Achilles intersubtendon matrix are not known. A frictionless  
174 boundary condition was used here to be computationally simple; in conjunction with a nonsliding  
175 boundary condition (see *Variation of model morphology* section below) these conditions represent  
176 the extreme examples of the sliding that may occur between Achilles subtendons. The proximal  
177 aspect of the Achilles was constrained to only move in the proximal-distal direction. The anterior-  
178 distal aspect of the calcaneal insertion was assigned to rotate 25° to simulate the rotation of the  
179 ankle during the dorsiflexion portion of the loading conditions described by Slane and Thelen (2014).  
180 The center of rotation was defined as the center of the line joining the subject's medial and lateral  
181 malleoli determined from MR images. The axis of rotation was perpendicular to the sagittal imaging  
182 plane and passed through the center of rotation.

183 To simulate the forces applied to the tendon by the three triceps surae muscles during both  
184 passive and active eccentric ankle dorsiflexion, nodal stiffness and pressure boundaries were applied  
185 to the proximal surface of each subtendon (Fig. 3D). Linear nodal stiffness boundary conditions were  
186 used to simulate the passive forces in muscle, which increase with increasing stretch (Herzog et al.,  
187 1991). The stiffness parameters were tuned based on comparison with *in vivo* data of passive  
188 dorsiflexion, which is described below. Linearly ramped pressure boundaries were used to simulate  
189 the active component of muscle forces on the ascending limb of the muscle force-length curve

190 (Maganaris, 2003); for eccentric simulations, both passive and active properties of muscle were  
191 modeled.

192 Comparison with *in vivo* data and muscle force determination

193 To compare the present modeling results with the tissue displacements from Slane and Thelen  
194 (2014), we defined a region of interest within the model that was consistent with the region of  
195 interest from their imaging study (Fig. 4). We projected 3D nodal displacements from the region of  
196 interest into the sagittal plane to obtain sagittal displacement profiles. Sagittal displacement profiles  
197 were smoothed with a quadratic surface fit as was done to ultrasound data from Slane and Thelen  
198 (2014). Nodes were binned into Anterior, Middle, and Posterior regions of the tendon (Fig. 4B and  
199 4C) and displacements were averaged within these bins.

200 To approximate the muscle forces relevant during passive and eccentric ankle dorsiflexion, we first  
201 tuned the stiffness coefficients corresponding to the proximal boundary conditions according to the  
202 displacement profiles from passive trials reported in (Slane and Thelen, 2014). The stiffness  
203 boundary conditions in the passive model were varied until the nodal displacements in the Anterior,  
204 Middle, and Posterior regions were within one standard deviation of experimental values. After  
205 stiffness boundaries had been set, we tuned pressure boundaries using data from eccentric trials  
206 reported in (Slane and Thelen, 2014) following the same protocol. Stiffness coefficients and pressure  
207 magnitudes for the LG subtendon were each set as one-half the coefficients and magnitudes used  
208 for the MG subtendon. This proportion reflects the relative PCSAs of these two muscles (Handsfield  
209 et al., 2014). In addition to the passive and eccentric tuned models, we ran an additional simulation  
210 with no force boundary conditions on the subtendons. These *no muscle force* models were used in  
211 order to observe the isolated effects of ankle kinematics on Achilles displacements, in the absence of  
212 any muscle force contribution.

213 Variation of model morphology

214 In order to explore the contributions of different geometric and functional features of the Achilles  
215 tendon, we created additional models that systematically eliminated various features. The first  
216 model variation—*no insertion*—was identical to the full model in the free tendon region, but did not  
217 include the calcaneal region of the tendon. The second model variation—*no insertion no twist*—  
218 similarly did not include the calcaneal region, but also lacked any twisting of the subtendons. The  
219 third model variation—*no sliding*— included the full tendon morphology but did not allow sliding  
220 between subtendons. External model geometry and all proximal and distal boundary conditions  
221 were maintained across the model variations.

222 RESULTS

223 Muscle force determination and comparison with experimental results

224 The model generated sagittal plane displacement profiles similar to those reported in Slane  
225 and Thelen (2014), and reproduced the result (Fig. 1) that distal displacements were greater in the  
226 anterior region of the free tendon than in the middle and posterior regions (Fig. 4)(Arndt et al., 2012;  
227 Franz et al., 2015; Slane and Thelen, 2015, 2014). Displacement magnitudes in the simulations were  
228 within one-third of one standard deviation of the mean from experimental ultrasound data (Fig. 5)  
229 and the profile of the displacements shows the same qualitative trend as the data in Slane and

230 Thelen for both passive and eccentric cases. The maximum simulated muscle forces are given in  
231 Table 1. Passive muscle forces in the soleus represented 34% of the total passive forces in the triceps  
232 surae, passive forces in the medial gastrocnemius represented 39% of the total, and passive forces in  
233 the lateral gastrocnemius represented 27% of the total (Table 1). In eccentric simulations, forces in  
234 the soleus represented 63% of the total triceps surae force, medial gastrocnemius force represented  
235 23% of the total, and lateral gastrocnemius force represented 14% of the total (Table 1).

236 Strains, displacements, and forces in tendon

237 Simulation of increased muscle force conditions resulted in an overall decrease in tissue  
238 displacements and increase in average strains in the free Achilles tendon (Fig. 6). In spite of a  
239 uniform stress-strain relationship in all subtendons, the force-strain relationships differed across  
240 subtendons and between *sliding* and *no sliding* models (Fig. 6) as a result of differences in geometry  
241 between the subtendons and differences in overall loading related to *sliding vs. no sliding* models.  
242 Displacement nonuniformity between subtendons was greatest for the passive force sliding models  
243 (Fig. 6A). For both the passive and eccentric simulations in the sliding model, strains were highest in  
244 the MG subtendon followed by the LG subtendon and SOL subtendon (Fig. 6A). Non-zero strains  
245 were observed in the *no muscle force* condition as a result of boundary conditions such as the  
246 constrained path of the proximal free tendon and contact between subtendons, which caused small  
247 strains in the subtendons. Differential strains were highest between the SOL and MG subtendons in  
248 the passive muscle force condition. Strains in the SOL subtendon were consistently the lowest, even  
249 when simulated soleus muscle forces were greater than either gastrocnemius muscle (Fig. 6A). For  
250 the non-sliding models at all force conditions, displacement nonuniformity was negligible (Fig. 6B).

251 Displacement nonuniformity: contribution of muscle forces, insertion, twist, and sliding

252 Displacement nonuniformity in the Achilles tendon varied throughout the length of the tendon, was  
253 greatest near the proximal aspect of the free tendon, and diminished toward the calcaneal insertion  
254 (Fig. 7A). When no forces were applied to the subtendons, displacement nonuniformity was  
255 negligible. In the sliding models, displacement nonuniformity between subtendons was non-  
256 negligible when any muscle forces (passive or eccentric) were applied, and were the greatest in the  
257 passive force condition (Fig. 7A). In the absence of the retrocalcaneal tendon insertion (i.e. free  
258 tendon only), displacements increased throughout the length of the free tendon but displacement  
259 nonuniformity between the subtendons did not differ greatly compared to the model containing an  
260 insertion region (Fig. 7). In the absence of both the retrocalcaneal tendon insertion and the twisting  
261 subtendons, displacements decreased in the SOL subtendon but increased in the LG and MG  
262 subtendons, resulting in an overall decrease in displacement nonuniformity (Fig. 7). For all of the  
263 sliding models, variations in the force boundary conditions had a greater effect on displacement  
264 nonuniformity than any of the morphological variations. Displacement nonuniformity was negligible  
265 through the length of the tendon for the non-sliding models, especially between MG and SOL  
266 subtendons (Fig. 7D).

267

268 DISCUSSION

269 Nonuniform displacements within the tendon resulted primarily from different applied muscle  
270 forces between subtendons. The highest nonuniformity occurred during passive muscle force  
271 simulations when the MG muscle contributed the most (39%) to total Achilles tendon force,  
272 followed by the soleus (34%) and the LG muscle (27%). In the eccentric condition, displacements  
273 were more uniform and the soleus contributed the most force (63%), followed by the MG (23%) and  
274 the LG (14%). Thus, differences in relative force contribution from the muscles may explain  
275 differences between passive and eccentric trials. There are known architectural and functional  
276 differences between the soleus and gastrocnemius muscles which could explain this phenomenon  
277 (Arndt et al., 1998; Cronin et al., 2013a). The higher passive force contribution from the medial  
278 gastrocnemius compared to the soleus implies that these muscles may differ in passive force curves  
279 or slack lengths relative to ankle position. In the present model, the stiffness boundary condition  
280 used to simulate medial gastrocnemius passive force was nearly two-and-a-half times the stiffness  
281 that simulated soleus passive force. The hypothesized greater passive stiffness in the medial  
282 gastrocnemius subtendon may promote higher stretching and elastic storage of energy in the  
283 posterior aspect of the Achilles tendon, a potentiality which warrants further study.

284 Elimination of intra-tendon sliding altered displacement nonuniformity more than variations in  
285 muscle force or tendon morphology. Sliding was a necessary condition in order to recapitulate the  
286 displacement profiles from the literature. Elimination of sliding severely reduced displacement  
287 nonuniformity in the tendon. In contrast to the *nonsliding models*, models with no retrocalcaneal  
288 insertion and no subtendon twist diminished displacement nonuniformity only slightly. This suggests  
289 that sliding is a much more important contributor to nonuniformity than twisting or insertion  
290 location. Taken together, sliding and non-zero muscle forces were both necessary to induce  
291 noticeable nonuniform displacements within the tendon model. Interestingly, sliding appears to be a  
292 phenomenon present in healthy young, but is diminished in older, horse fascicles (Thorpe et al.,  
293 2013) and human tendons (Slane and Thelen, 2015). In the present work, we lacked values or  
294 estimates of the material stiffness of the matrix between Achilles subtendons; therefore, we used  
295 frictionless sliding and nonsliding as bounds for the sliding that likely occurs between subtendons. In  
296 a previous work, Thorpe et al. investigated the mechanics of the matrix between tendon fascicles  
297 (Thorpe et al., 2015); a similar study of the matrix between human Achilles subtendons would  
298 enable a deeper understanding of the behavior of these structures.

299 In this study, we estimated triceps surae muscle forces by tuning force boundary condition  
300 magnitudes to displacement profiles observed in (Slane and Thelen, 2014). Peak soleus forces  
301 simulated in this study are on the order of 200N passive and 1000N active. Peak gastrocnemius  
302 forces simulated in this study are about 350N passive and 450N active. Slane & Thelen (2014)  
303 reported net eccentric plantarflexion moments in excess of 30Nm, implying plantarflexion forces  
304 around 600N. The forces in the current model exceeded these net force estimates from Slane &  
305 Thelen (2014) but were physiologically reasonable and consistent with several *in vivo* literature  
306 reports (Arndt et al., 1998; Finni and Komi, 1998; Mian et al., 2007; Rubenson et al., 2012). It should  
307 be noted that our boundary conditions ignored any resistive forces from fluid and connective tissue  
308 within the ankle.

309 Subject-specific material properties were not available in this study. The material parameters chosen  
310 to model subtendons were based on literature reports of *in vivo* Achilles tendon material properties  
311 (Kubo et al., 2002). The Neo-Hookean model used was within the range of tendon material response

312 previously observed (Kubo et al., 2002; Lichtwark and Wilson, 2005; Stenroth et al., 2012). Our  
313 material model was less stiff than the average for young women observed by Stenroth et al. (2012),  
314 but it was within the broad range of previously reported tendon material responses (Kubo et al.,  
315 2002; Lichtwark and Wilson, 2005; Maganaris and Paul, 2002; Stenroth et al., 2012; Zajac, 1989).  
316 Because we tuned our forces to observed displacements, our result of high subtendon stretch,  
317 especially in the MG subtendon, is not sensitive to the specific material stiffness chosen in this study.  
318 Future incorporation of a transversely isotropic (Weiss et al., 1996) or biphasic (Yin and Elliott, 2004)  
319 material model may be used to model Achilles behavior in finer detail. We used an assumption that  
320 the ankle axis of rotation was perpendicular to the sagittal plane, which simplifies the kinematics of  
321 ankle motion that have been demonstrated previously (Lundberg et al., 1989). Incorporation of  
322 more sophisticated ankle kinematics, potentially also including inversion and eversion, may reveal  
323 more specific subtendon deformation patterns, especially increased displacements in the MG  
324 subtendon. In this study, we defined a region within the Achilles tendon model to be consistent with  
325 the region that was imaged using ultrasound in Slane and Thelen (2014). In light of the 3D structure  
326 of the Achilles tendon, it is likely that variations in ultrasound probe placement on the posterior  
327 ankle and morphological variations of subtendons across subjects may alter the tendon deformation  
328 patterns observed with ultrasound. In previous works, nonuniform deformations have been  
329 consistently observed (Arndt et al., 2012; Franz et al., 2015; Slane and Thelen, 2014) in spite of  
330 potential inter-subject variations and probe placement variations, implying that these variations may  
331 not contribute much to variations in deformation. In the future, the 3D computational models  
332 presented here could be used to explore the effect of anatomical variation and probe placement  
333 variation on the observed deformation of the Achilles tendon.

334 We used UTE MRI and literature information about subtendon morphology to develop an image-  
335 based model of the Achilles tendon. The imaging sequence used here is a 3D radial UTE sequence  
336 that was tuned to promote high contrast between tendon, cortical bone, fat, bursa, and muscle. UTE  
337 sequences may be used to image tissues with short  $T2^*$  times. The sequence used here worked well  
338 to distinguish the Achilles tendon from its neighboring structures. Finite element models of the  
339 Achilles tendon have been used previously to understand tendon loading (Shim et al., 2014) and the  
340 effects of surgery on the tendon (Von Forell and Bowden, 2014). This is the first work to our  
341 knowledge to use UTE MRI to inform Achilles model geometry. The UTE MR images did not have  
342 enough resolution to distinguish the separate Achilles subtendons, which necessitated our use of  
343 literature information to model these sub-structures. Future imaging and modeling approaches  
344 incorporating the meso- and micro-scale structures of tendon may reveal additional insights into the  
345 importance of these substructures.

346 This study may have implications to the aging tendon. Aging is reportedly related to a reduction in  
347 length of the retrocalcaneal insertion of the Achilles (Kim et al., 2011), possibly due to progressive  
348 ossification of the most distal regions of the insertion or calcaneal growth that alters the relative  
349 location of tendon insertion. Aging also may reduce intra-tendon sliding due to collagen cross-linking  
350 (Thorpe et al., 2013). Changes such as these would be expected to reduce displacement  
351 nonuniformity between subtendons. Age-related muscle weakening (Doherty, 2003) and changes to  
352 tendon stiffness (Narici et al., 2008; Stenroth et al., 2012) would also affect the degree of  
353 nonuniformity in subtendon displacements. Future work may suggest if and how subtendon sliding is  
354 related to tendon aging and may indicate if tendon health is related to the ability for sliding between  
355 subtendons.

356 The current work provides a foundation for how intra-tendon sliding, muscle forces, twisting  
357 subtendons, and insertional morphology may influence the extent of displacement nonuniformity  
358 within the Achilles tendon. Future modeling and experimental work are needed to further explore  
359 these and other morphological features of the tendon and to contextualize the mechanisms of  
360 differential muscle forces and intratendon sliding as contributors to nonuniform Achilles  
361 deformation.

362

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366

#### 367 CONFLICTS OF INTEREST

368 The authors have no conflicts of interest to disclose.

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- 495

## 496 FIGURE CAPTIONS

497 Figure 1: Illustration of tissue displacements in Achilles tendon. Kinematics alone suggests greater  
498 displacements in the posterior tendon during ankle dorsiflexion. In multiple experimental  
499 observations, greater displacement was seen in the anterior tendon (Arndt et al., 2012; Franz et al.,  
500 2015; Slane and Thelen, 2015, 2014). Figure adapted from Sobotta (1909).

501 Figure 2: Summary of imaging and segmentation of external tendon geometry. A: Sagittal and axial  
502 slices from 3D UTE-shTE images illustrate the location of the Achilles tendon and its segmentation in  
503 axial slices. B: 3D reconstruction of the segmented images reveals whole tendon shape from soleus  
504 myotendinous junction to retrocalcaneal insertion.

505 Figure 3: Illustration of the subtendons defined in the finite element model. A: Proximal view of  
506 model displays subtendons associated with the soleus, medial gastrocnemius, and lateral  
507 gastrocnemius muscles. B: distal view of the free tendon displays the location of the three  
508 subtendons at this level, which are rotated relative to the locations in A. C: Longitudinal views of the  
509 tendon reveal the twisted orientation of the subtendons. D: Passive and active muscle forces were  
510 modeled as stiffness and pressure boundary conditions, respectively. Passive simulations consisted  
511 of stiffness boundaries only while eccentric simulations consisted of both stiffness and pressure  
512 boundaries.

513 Figure 4: Qualitative comparison of displacement profiles between experiment and finite element  
514 model. A: In Slane and Thelen (2014), the Achilles tendon was imaged with ultrasound while subjects  
515 went through cyclic ankle dorsiflexion (adapted from Sobotta (1909)). B: With dorsiflexion,  
516 ultrasound imaging reveals nonuniform displacements—anterior tissue displaced more than  
517 posterior tissue. (adapted from Slane and Thelen (2014)) C: In the FE model, nodal displacements in  
518 the anterior region were greater than those in the posterior region.

519 Figure 5: Quantitative comparison of regional displacements between experiment and finite element  
520 model. The peak distal displacements occurring in the anterior, middle, and posterior regions are  
521 shown for ultrasound experiments (Slane and Thelen, 2014) and for the present finite element  
522 model. For both passive and eccentric cases, finite element simulations predicted regional  
523 displacements within one standard deviation of experimental results.

524 Figure 6: Relationships between simulated muscle forces, tissue displacements, and strains in FE  
525 models. A: In the sliding model, distal displacements in the proximal free tendon decreased with  
526 increasing muscle forces and equivalent strains increased with muscle force. Force-strain  
527 relationships were similar between the gastrocnemius subtendons but differed in the soleus  
528 subtendon. B: In the absence of sliding subtendons, differential displacements were negligible and  
529 force-strain relationships were altered for all three subtendons.

530 Figure 7: Displacement profiles across morphological variations of the Achilles tendon. A: The  
531 reference Achilles model includes twisted subtendons, inter-subtendon sliding, and a retrocalcaneal  
532 insertion. B: Without this insertion, the model displays nonuniform displacements. C: Without  
533 twisted subtendons, the differential displacements in the free tendon are slightly reduced. D: In the  
534 absence of sliding, differential displacements between subtendons are negligible.

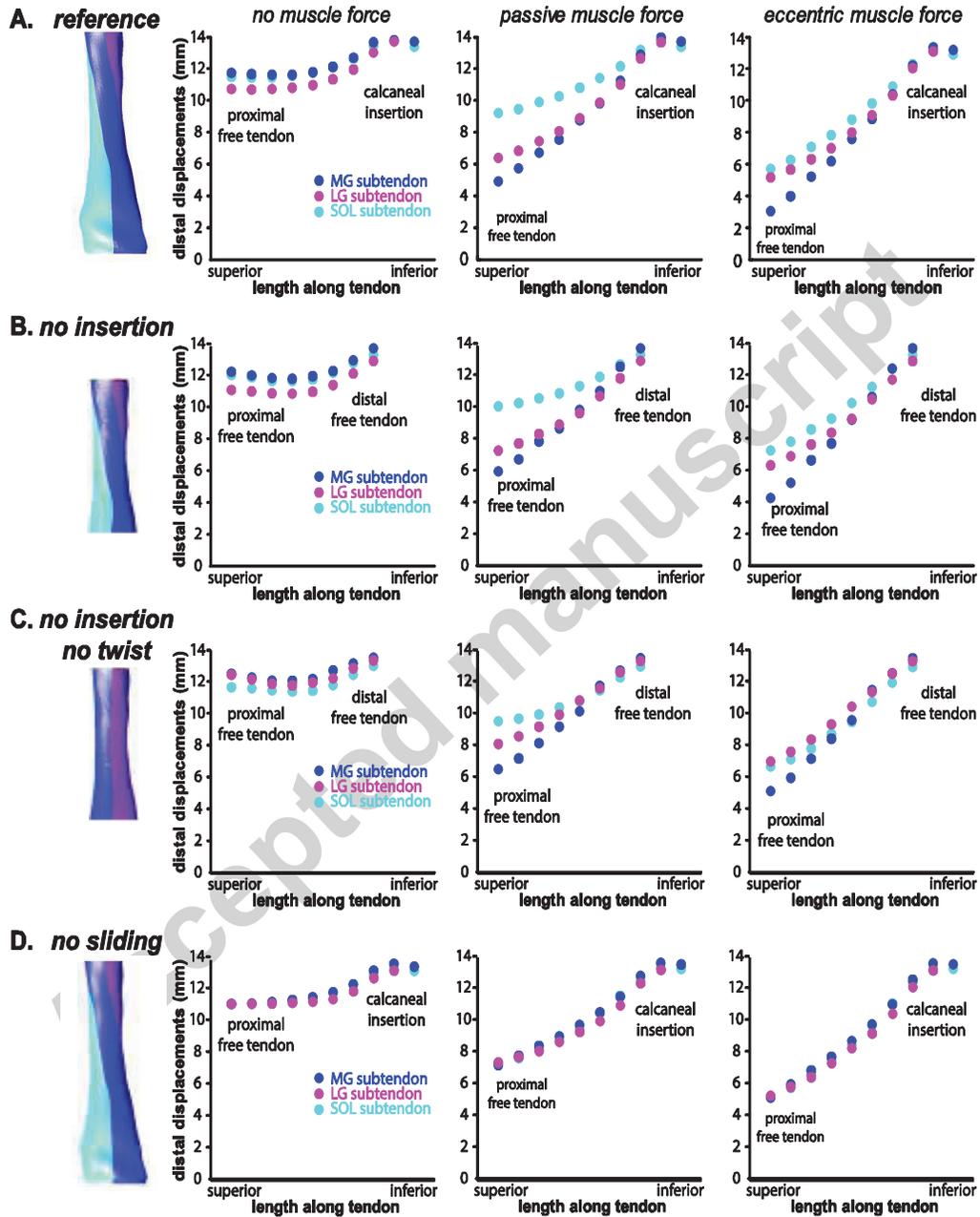
535 Table 1: Maximum muscle forces on Achilles subtendon in FE model

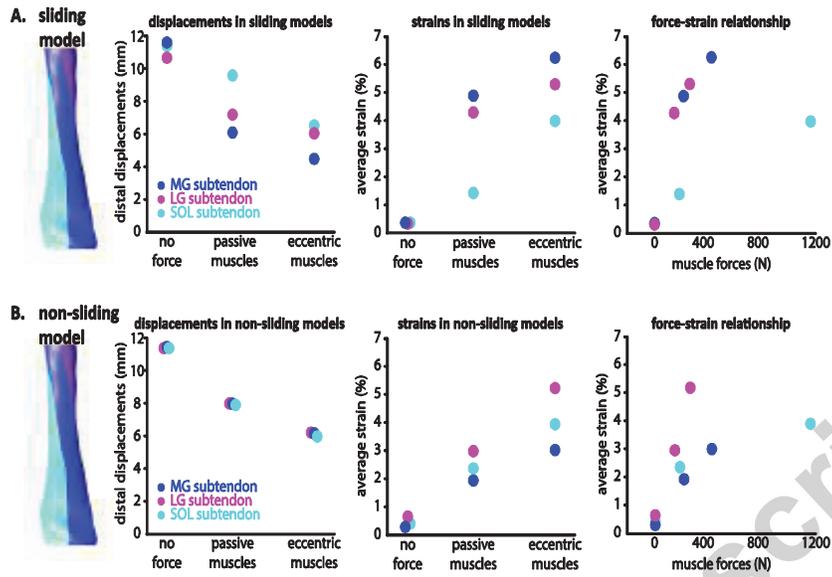
Maximum Forces During Simulation (Passive Forces) [% of total]			
Condition	Soleus Subtendon	MG Subtendon	LG Subtendon
Passive	(189 N) [34%]	(217 N) [39%]	(149 N) [27%]
Eccentric	1157 N (111 N) [63%]	423 N (132 N) [23%]	263 N (118 N) [14%]

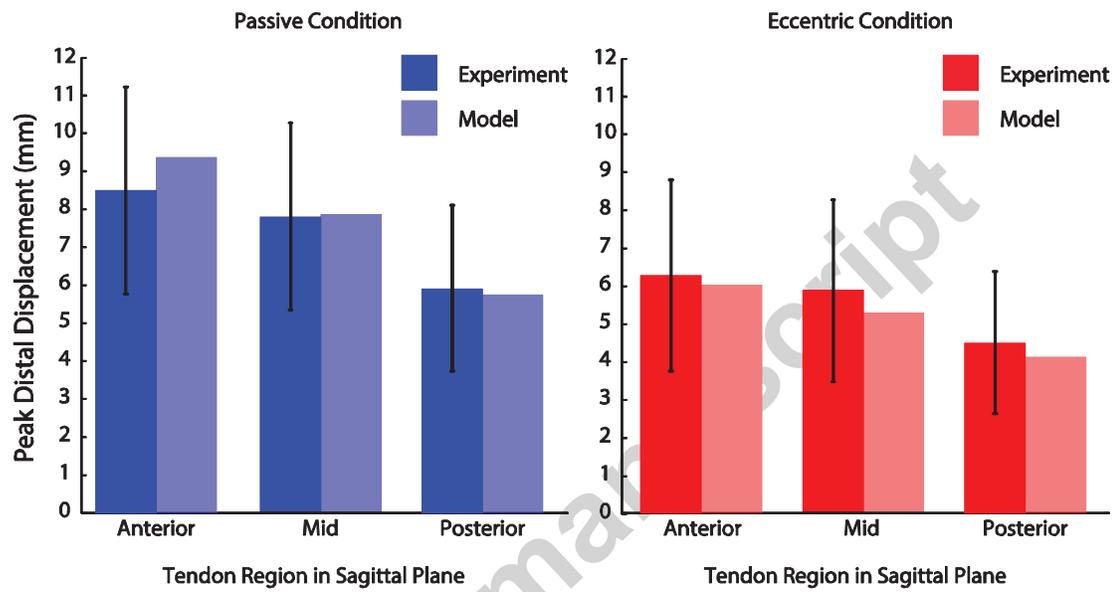
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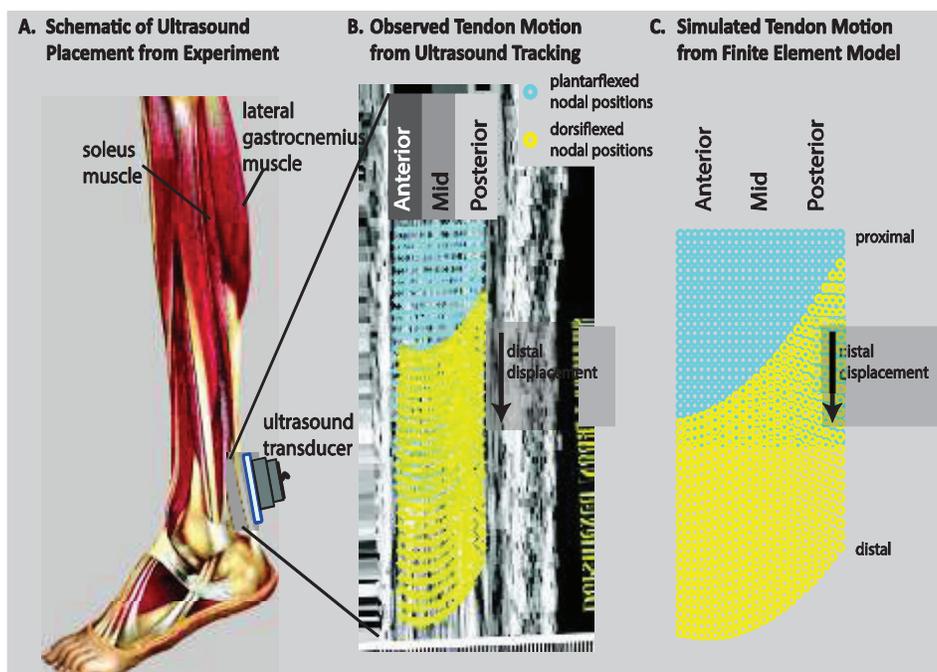
537 Maximum muscle forces in the passive and eccentric models are tabulated. Total force in each  
 538 subtendon are given in Newtons, passive force in parentheses; percent of total Achilles force is given  
 539 in brackets.

540



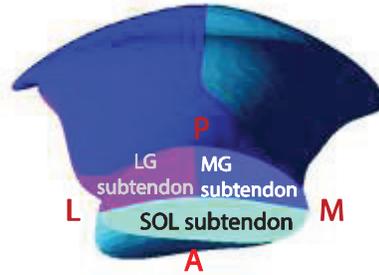




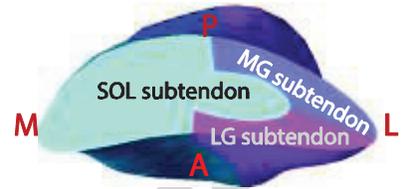


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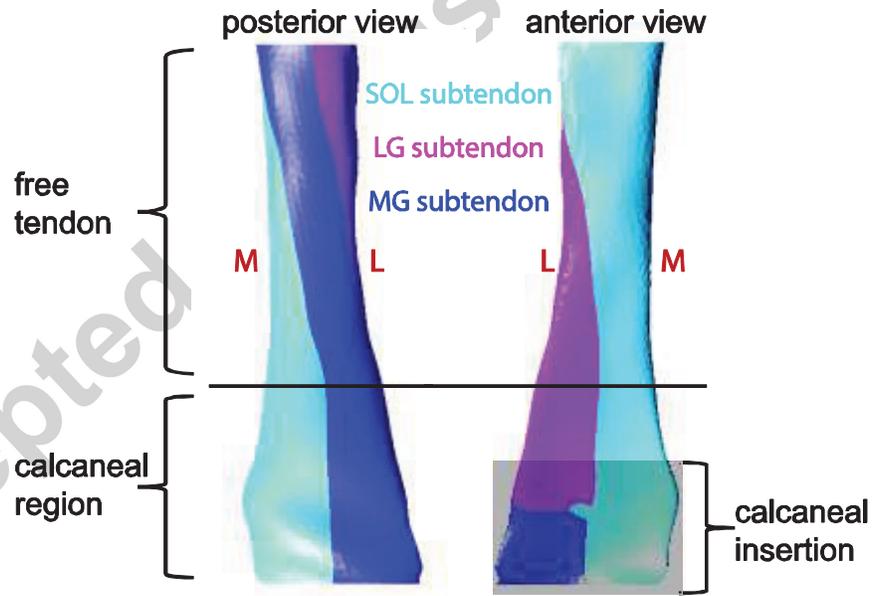
A. superior view



B. inferior view of free tendon

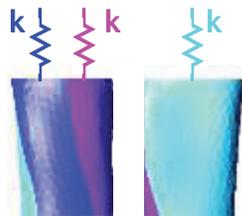


C. longitudinal views of subtendon/tendon model

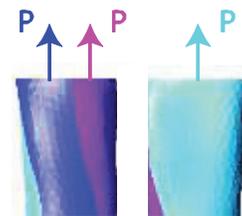


D. proximal boundary conditions

passive muscle modeled with stiffness boundary,  $k$

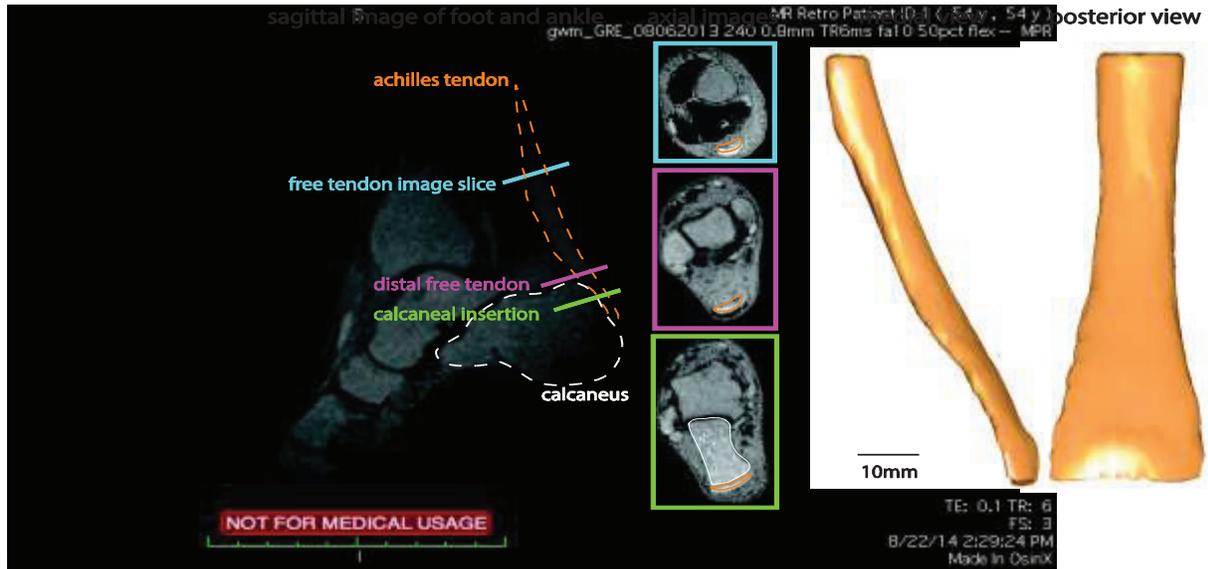


active muscle modeled with applied pressure,  $P$



**A. MR Images of Achilles tendon were segmented from the proximal free tendon through calcaneal insertion**

**B. 3D model of the Achilles tendon constructed from segmented images**



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**nonuniform motion expected**

more displacement in posterior portion  
due to larger moment arm

**nonuniform motion observed**

less displacement  
in posterior portion



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