An advanced glycation endproduct (AGE)-rich diet promotes accumulation of AGEs in Achilles tendon

Skovgaard, Dorthe Charlotte; Svensson, René B.; Scheijen, Jean; Eliasson, Pernilla; Mogensen, Pernille; Hag, Anne Mette Fisker; Kjær, Michael; Schalkwijk, Casper G; Schjerling, Peter; Magnusson, Stig Peter; Couppé, Christian

Published in:
Physiological reports

DOI:
10.14814/phy2.13215

Publication date:
2017

Document Version
Publisher’s PDF, also known as Version of record

Citation for published version (APA):
Skovgaard, D., Svensson, R. B., Scheijen, J., Eliasson, P., Mogensen, P., Hag, A. M. F., ... Couppé, C. (2017). An advanced glycation endproduct (AGE)-rich diet promotes accumulation of AGEs in Achilles tendon. DOI: 10.14814/phy2.13215
An advanced glycation endproduct (AGE)-rich diet promotes accumulation of AGEs in Achilles tendon

Dorthe Skovgaard1, Rene B. Svensson2, Jean Scheijen3, Pernilla Eliasson2, Pernille Mogensen2, Anne Mette F. Hag1, Michael Kjær2, Casper G. Schalkwijk3, Peter Schjerling2, Stig P. Magnusson2,4 & Christian Coupp2,4

1 Department of Clinical Physiology, Nuclear Medicine & PET and Cluster for Molecular Imaging, Rigshospitalet and University of Copenhagen, Copenhagen, Denmark
2 Department of Orthopedic Surgery M, Institute of Sports Medicine and IOC Research Centre Copenhagen, Bispebjerg Hospital and Center for Healthy Aging, Faculty of Health and Medical Sciences, University of Copenhagen, Copenhagen, Denmark
3 Experimental Internal Medicine at the Faculty of Health, Medicine and Life Sciences, Maastricht University Medical Center, Copenhagen, The Netherlands
4 Department of Physical Therapy, Musculoskeletal Rehabilitation Research Unit, Bispebjerg Hospital, University of Copenhagen, Copenhagen, Denmark

Keywords
Achilles tendon, advanced glycation endproducts, AGE-rich diet, aging.

Abstract
Advanced Glycation Endproducts (AGEs) accumulate in long-lived tissue proteins like collagen in bone and tendon causing modification of the biomechanical properties. This has been hypothesized to raise the risk of orthopedic injury such as bone fractures and tendon ruptures. We evaluated the relationship between AGE content in the diet and accumulation of AGEs in weight-bearing animal Achilles tendon. Two groups of mice (C57BL/6Ntac) were fed with either high-fat diet low in AGEs high-fat diet (HFD) (n = 14) or normal diet high in AGEs (ND) (n = 11). AGE content in ND was six to 50-fold higher than HFD. The mice were sacrificed at week 40 and Achilles and tail tendons were carefully excised to compare weight and nonweight-bearing tendons. The amount of the AGEs carboxymethyllysine (CML), methylglyoxal-derived hydroimidazolone (MG-H1) and carboxyethyllysine (CEL) in Achilles and tail tendon was measured using ultraperformance liquid chromatography tandem mass spectrometry (UPLC-MS/MS) and pentosidine with high-pressure liquid chromatography (HPLC) with fluorescent detection. AGEs in Achilles tendon were higher than in tail tendon for CML (P < 0.0001), CEL (P < 0.0001), MG-H1 and pentosidine (for both ND and HFD) (P < 0.0001).

The AGE-rich diet (ND) resulted in an increase in CML (P < 0.0001), MG-H1 (P < 0.001) and pentosidine (P < 0.0001) but not CEL, in Achilles and tail tendons. This is the first study to provide evidence for AGE accumulation in injury-prone, weight-bearing Achilles tendon associated with intake of an AGE-rich diet. This indicates that food-derived AGES may alter tendon properties and the development of tendon injuries.

Introduction
Advanced Glycation Endproducts (AGEs) are chemically modified molecules with pathogenic implications for aging, diabetes and several other chronic diseases (Brownlee 2001; Monnier et al. 2005). AGEs are formed through a nonenzymatic reaction between reducing sugars and free
amino groups of proteins, lipids, or nucleic acids. This reaction occurs endogenously in the body, and it is long-lived proteins in the extracellular matrix (ECM) that are particularly susceptible to AGE modifications with aging (Monnier et al. 2005). In collagen-rich structures like bone and tendon, AGEs form cross-links, which morphological modify the collagen network. This alters the biomechanical properties of the tissue with increased stiffness and fragility (Eriksen et al. 1985; Monnier et al. 2005; Zimmermann et al. 2011; Grasa et al. 2013). It has been hypothesized that this accumulation can raise the risk of orthopedic injury such as bone fractures and tendon ruptures (Kjaer 2004; Monnier et al. 2005).

In recent years, much focus has been on the negative effects of dietary intake of AGEs on healthy tissues. Current evidence support that ingestion of AGE-rich diet increases the AGE content in major arteries, heart, and bone that lead to age-related changes, and pathology in both animals and humans (Eriksen et al. 1985; Koschinsky et al. 1997; Shapiro et al. 2008; Uribarri et al. 2011). However, in the present substudy we only focused on the impact of the dietary intake of AGEs, and therefore only WT mice were investigated. The mice were obtained at 8 week of age (Taconic Europe, Lille Skensved, Denmark), from which point one-half of the mice (#D12079B, Research Diets, New Brunswick, NJ) to induce obesity, and the other half (n = 11) were fed ad libitum normal diet (ND) (protein, 27 cal%; carbohydrate, 60 cal%; fat, 13 cal%); #1310, Altromin Spezialfutter & KG, Germany). The difference of the group size is due to loss of tissue samples. As previously published, the amount of AGEs were 6–50 times higher in ND compared to HFD (CML, CEL and MG-H1) had higher content of AGEs in their nonweight-bearing tail tendon compared to mice ingesting a diet with low amounts of AGEs (Eriksen et al. 1985). The tail tendons of mice with high AGE intake also displayed increased stiffness, which indicates that AGE-rich diet leads to functionally important accumulation of AGEs in the connective tissue that do not see large loads (Eriksen et al. 1985). However, as tail tendon is only mechanically loaded to a very moderate degree the findings do not reveal whether or not dietary AGE accumulation will be the same in a weight-bearing and thus more mechanically loaded tendon (Svensson et al. 2016). Recently it was found, that in master athletes the age-related AGE accumulation in patellar tendon was less pronounced compared to untrained but healthy counterparts (Couppe et al. 2014). Furthermore, mechanical loading during treadmill training in mice increases collagen turnover and reduces AGEs in tibialis anterior tendon (Wood and Brooks 2016). Thus, it is conceivable that weight-bearing tendons have lower AGE content than less loaded tendons, and we wanted in the present study to investigate whether a AGE-rich diet promotes AGE accumulation also in tendons that are subjected to high mechanical loading, or whether this mechanical loading rather counteracts the diet induced AGE accumulation.

Objective/Purpose

The purpose of this study was to investigate if a diet containing high amounts of AGEs (CML, CEL and MG-H1) results in AGE accumulation in the clinically important Achilles tendon. In addition, we wanted to compare the level of AGEs with data from the previous published study on tail tendon to elucidate possible differences in the quantitative level of AGEs in tendons with different functional importance.

We hypothesized that AGE accumulation would be more pronounced with high dietary AGE intake in Achilles tendon, similarly to that earlier found in the tail tendon. In addition, we hypothesized that the amount of AGEs in Achilles tendon would be lower compared to tail tendon tissue because of a mechanically induced higher collagen turnover in the weight-bearing Achilles tendon.

Methods

Dietary mouse model

This investigation builds on a previously published study, in which atherosclerosis and AGE accumulation in tail tendon tissue were investigated using both homozygous ApoE−/− mice (“atherosclerotic phenotype”) and wild type (WT) mice (C57BL/6Ntac) on different diets (Eriksen et al. 1985). However, in the present substudy we only focused on the impact of the dietary intake of AGEs, and therefore only WT mice were investigated. The mice were obtained at 8 week of age (Taconic Europe, Lille Skensved, Denmark), from which point one-half of the mice (n = 14) were fed ad libitum HFD (protein, 17 cal%; carbohydrate, 43 cal%; fat, 41 cal%; Western Diet, #D12079B, Research Diets, New Brunswick, NJ) to induce obesity, and the other half (n = 11) were fed ad libitum normal diet (ND) (protein, 27 cal%; carbohydrate, 60 cal%; fat, 13 cal%); #1310, Altromin Spezialfutter & KG, Germany). The difference of the group size is due to loss of tissue samples. As previously published, the amount of AGEs were 6–50 times higher in ND compared to HFD (ND vs. HFD: CML: 206.1 vs. 4.7 nmol/g; CEL: 86.5 vs. 3.0 nmol/g; MG-H1: 554.0 vs. 34.5 nmol/g;) (Eriksen et al. 1985). Mice were sacrificed at 40 weeks of age by decapitation after having fasted overnight and being anesthetized for a period of 4 h, during which time the mice were 18F-Fluoro-Deoxy-Glucose Positron Emission Tomography (PET) scanned as part of the main study looking at the progression of atherosclerosis (Hag et al. 2012). The hind legs of each mouse were removed and stored at −20°C until the experimental day. One experienced researcher (PE) carefully excised the Achilles tendon in toto, being
conscientious not to include muscle fibers in the sample using a technique that was performed in previous studies (Eliasson et al. 2007, 2009).

Animal care and experimental procedures were performed under the approval of the Animal Experiments Inspectorate in Denmark (permit number 2011/561-14).

AGE analysis

Each Achilles tendon sample was incubated in 1 mL of 1 mol/L NaCl at 4°C for 24 h with at least one fluid exchange. After removal of NaCl, the tendon samples were incubated in 1 mL of 0.5 mol/L acetic acid at 4°C for 24 h with one fluid exchange. The acetic acid was removed and the tendon samples were incubated with 1 mL of chloroform-methanol (2:1 vol/vol) at 4°C for 24 h with one fluid exchange (Eriksen et al. 1985; Dunn et al. 1991). After removal of the organic solvent, the tendon samples were dried under a gentle stream of nitrogen at room temperature. To avoid AGE formation during hydrolysis, samples were reduced with 0.2 mL of 0.1 mol/L NaBH4 in 0.2 mol/L borate buffer (pH 9.2) for 2 h at room temperature. The reduced samples were deproteinized in 1 mL of cold (4°C) 20% (vol/vol) trifluoroacetic acid and centrifuged at 4300 g, 4°C for 20 min, and the supernatant was discarded. The remaining pellets were hydrolyzed in 0.5 mL of 6 mol/L HCl. From each hydrolysate, 40 μL were taken and mixed with 20 μL of internal standard before analysis. CML, CEL, and MG-H1 were hydrolyzed in 0.5 mL of 6 mol/L HCl. From each hydrolysate, 40 μL were taken and mixed with 20 μL of internal standard before analysis. CML, CEL, and MG-H1 were hydrolyzed in 0.5 mL of 6 mol/L HCl. From each hydrolysate, 40 μL were taken and mixed with 20 μL of internal standard before analysis. CML, CEL, and MG-H1 were hydrolyzed in 0.5 mL of 6 mol/L HCl. From each hydrolysate, 40 μL were taken and mixed with 20 μL of internal standard before analysis. CML, CEL, and MG-H1 were hydrolyzed in 0.5 mL of 6 mol/L HCl. From each hydrolysate, 40 μL were taken and mixed with 20 μL of internal standard before analysis.

Data reduction and statistics

Results were analyzed using a two-way mixed model ANOVA (tissue type, diet) with repeated measures for the tissue type since Achilles and tail tendons were from the same animal. Initially the analysis included interactions, if the interaction was not significant, data were finally analyzed using only main effects. Tukey method for multiple comparisons was used in post hoc tests.

Results

Baseline characteristics

As previously reported (Eriksen et al. 1985), mice fed with HFD had higher body weight compared to mice consuming ND (HFD: 47.4 ± 3.6 vs. ND: 33.9 ± 3.4 gram, P < 0.0001), but there were no significant differences in plasma glucose (HFD: 15.8 ± 5.5 vs. ND: 13.3 ± 4.8 mmol/L, P = 0.35) or cholesterol level (HFD: 15.9 ± 5.5 vs. ND: 12.0 ± 4.2 mmol/L, P = 0.104).

Advanced glycation end products

Data and effect sizes are shown in Figure 1 and Table 1, respectively (see below). There were more AGEs in the Achilles tendon than the tail tendon for CML (P < 0.0001) (Fig. 1A), CEL (P < 0.0001) (Fig. 1B) and MG-H1 (P < 0.0001) (Fig. 1C) for both ND and HFD. This was also the case for pentosidine in both the ND (P < 0.0001) and HFD (P < 0.0001) group (Fig. 1D).

The ND led to increased CML (P < 0.0001) (Fig. 1A) and MG-H1 (P < 0.001) (Fig. 1C) in both the Achilles and tail tendons. CEL in the tendons was not significantly affected by ND (Fig. 1B). Except for pentosidine, none of the measured AGEs displayed any significant interaction term in the ANOVA, indicating that the effect of diet was...
identical for the Achilles and tail tendons. For pentosidine there was an interaction \((P < 0.01)\) (Fig. 1D), which showed an increase in pentosidine for the ND in the Achilles \((P < 0.0001)\) but not in the tail tendon \((P = 0.27)\).

**Discussion**

To the best of our knowledge, this is the first study to provide evidence of AGE accumulation in Achilles tendons only as a function of intake of an AGE-rich diet compared to high-fat (but low AGE) diet, thereby extending our previous findings with dietary AGEs as an important source of AGE accumulation in tendon tissue. Furthermore, contrary to our hypothesis, Achilles tendon demonstrated greater AGE content (CML, MG CEL and pentosidine) than tail tendon. This is somewhat surprising, as mechanical loading earlier has been shown to reduce accumulation of AGE in the weight-bearing patellar tendon (Kongsgaard et al. 2009; Couppe et al. 2014). Tendon mechanical properties are determined by the underlying composition and structure, although details still remains to be described. Type I collagen is the primary components of tendon ECM, constituting up to 85% of the tissue dry mass (Kjaer 2004; Svensson et al. 2016). Covalent, intermolecular cross-links associated with the assembly of the collagen fibrils are believed to be a major source of mechanical strength, and improved mechanical properties during development are likely related to maturation of these enzymatic cross-links (Eyre 1987; Bailey et al. 1998). However, the slow turnover of collagen allows for the accumulation of AGEs with aging that reduce collagen fibril sliding leading to impaired tendon mechanics (Verzijl et al. 2000; Li et al. 2013; Fessel et al. 2014). Although the biomechanical properties were not investigated in the present study, others and we have previously reported, that AGE-rich diet leads to AGE accumulation and increased mechanical properties (mainly in the region of plastic deformation, where AGEs

---

*Figure 1. Advanced glycation endproduct content in tail tendon and Achilles tendon. Data given as Mean ± SEM. Fat: \(n = 11\), Chow: \(n = 14\), (A): CML: ***diff from tail \(P < 0.0001\) (main effect), ##diff from HFD \(P < 0.001\) (main effect). (B): CEL: ***diff from tail \(P < 0.0001\) (main effect). (C): MG-H1: ***diff from tail \(P < 0.0001\) (main effect). (D): Pentosidine: There was an interaction \((P < 0.01)\), which showed an increase in pentosidine for the ND group in the Achilles (### diff from HFD \(P < 0.0001\)), but not in the tail tendon \((P = 0.27)\). For both diets there was an effect of tissue (***diff from tail \(P < 0.0001\)). HFD, high-fat diet.*
might play a role) in non weight-bearing tail tendons (Eriksen et al. 1985; Roncero-Ramos et al. 2014). In line with the study hypothesis, the present data provides evidence of an AGE-rich diet leading to AGE accumulation in Achilles tendon and thereby has an aging effect on tissue as previously reported (Eriksen et al. 1985). Consequently, AGE deposition in tendons might contribute to the higher incidence of tendinopathy and tendon rupture, as was already suggested in patients with diabetes and with aging (Kjaer 2004; de Jonge et al. 2011, 2015; Ranger et al. 2016; Svensson et al. 2016). However, future studies will have to address this issue also in normal individuals of different ages without diabetes.

Our second hypothesis was that AGE-content would be lower in Achilles tendon compared to tendon tail, irrespective of the diet, due to a presumed load-induced higher collagen turnover (Wood and Brooks 2016). Interestingly, the present data does not support this hypothesis.

An obvious explanation is that AGE cross-linking is the result of a balance between exposure and rates of collagen synthesis/turnover, with the half-life of collagen being one important determinant (Baynes 2000). Thorpe et al. found a clear difference in AGE content in two weight-bearing equine tendons with aging (Thorpe et al. 2010). The high strain and injury-prone superficial digital flexor tendon (comparable to the human Achilles tendon) displayed a higher AGE content and much lower collagen turnover (estimated half-life of collagen 197 years) compared to the low strain and rarely injured common digital extensor tendon, which had an estimated collagen half-life of 34 years (Thorpe et al. 2010). Our data therefore indicates that Achilles tendon, despite being subjected to greater loading, has a lower collagen turnover than tail tendon allowing larger amounts of AGEs to accumulate. The Achilles tendon is the largest tendon in the body and is very important during locomotion. It is possible that a low collagen turnover is necessary for optimal integrity and function of this important tendon. In fact, human Achilles tendon collagen fibrils (core part of tendon) have been found to be practically inert throughout adult life (Heinemeyer et al. 2013).

An important aspect is to what extent AGE accumulation can be counteracted with regular exercise/loading of the tissue. We recently demonstrated that individuals with type 2 diabetes have similar AGE content (pentosidine) as healthy controls in their Achilles tendons although they have a markedly higher AGE content in the skin (Couppe et al. 2014). These data suggest that weight-bearing connective tissue may respond differently to glycermic loading than non-weight-bearing connective tissue. It has been suggested that mechanical loading can increase collagen turnover and thereby prevent AGE accumulation (Avery and Bailey 2005; Kongsgaard et al. 2009; Couppe et al. 2014). In support of this, life-long running and resistance training have shown to reduce the age-associated accumulation of the AGE level in the human patellar tendon (Kongsgaard et al. 2009; Couppe et al. 2014). It is therefore possible that mechanical strain of the tendon and/or just a general systemic effect (better glycemic control) associated physical activity could help offset age-induced accumulation of AGE in tendons (Odetti et al. 1996; Boor et al. 2009). It is also possible that AGE-related pathogenic processes and the aging effect on tissue including those of tendons may be influenced by dietary AGE restriction. Several recent studies have implicated that low AGE-diet can reduce AGEs (pentosidine) in rodent tail tendon tissue and on whole-body level lower the markers of oxidative stress, inflammation and insulin resistance (Sell and Monnier 1997; Lingelbach et al. 2000; Vlassara et al. 2016). In the present study, the difference in AGE content is likely related to lower carbohydrate (sugar) content in the HFD and the processing of the two types of diet, where ND was manufactured with heating, which is well known to induce AGEs (Maillard reaction) (Maillard 1912; Urbirri et al. 2010).

### Table 1. Biochemical parameters.

|                     | Effect of Achilles tendon | Effect of ND |
|---------------------|---------------------------|--------------|
| CML (nmol/mmol hydroxyproline) | 14.9 ± 0.7*** | 7.4 ± 1.4**  |
| CEL (nmol/mmol hydroxyproline)  | 10.5 ± 0.9*** | 0.6 ± 0.9   |
| MG-H1 (nmol/mmol hydroxyproline) | 177.4 ± 13.6*** | 39.6 ± 10.1** |
| Pentosidine (within Tail) | 0.010 ± 0.005 |              |
| Pentosidine (within Achilles) | 0.031 ± 0.005*** |        |
| Pentosidine (within ND) | 0.056 ± 0.005*** |        |
| Pentosidine (within HFD) | 0.035 ± 0.005*** |        |

Mean ± SE for each group. Mean difference for effects of Achilles tendon and ND. Significant difference (***P < 0.001, **P < 0.01, *P < 0.05; 0.001).

For pentosidine there was an interaction (P < 0.01), which showed a significant increase in pentosidine for the ND group in the Achilles (P < 0.0001), but not in the tail tendon (P = 0.27). For both diets there was an effect of tissue (P < 0.0001). HFD, High-Fat Diet; ND, Normal Diet.
We cannot exclude the possibility that other differences between the diets (ND vs. HFD), for example, carbohydrate concentration, macronutrients and total energy intake, also contributed to the AGE accumulation in the tendons of the present study. However, others have demonstrated that with AGE-rich but isocaloric diet intake, pathologic changes in serum and tissue are still induced (Cai et al. 2007). Of note, dicarbonyl stress caused by the most reactive AGE precursor methylglyoxal (MGO) measured in this study can without sugar such as glucose or without going through the classical Maillard reaction pathways induce AGE formation (Rabbani and Thornalley 2012). Inflammation per se may also form AGEs that again induce inflammation creating a vicious cycle (Hanssen et al. 2014). The study was limited by a small sample size that possibly led to failure to demonstrate a significant difference in pentosidine for the tail tendon, which was actually demonstrated in our previous report (Eriksen et al. 1985).

In conclusion, we have found evidence for an accumulation of AGEs in the weight-bearing Achilles tendon due to intake of AGE-rich diet. This suggests that food-related intake of AGEs could potentially alter tendon properties and thereby represent a risk factor in the development of tendon injuries.

**Conflict of Interest**

None declared.

**References**

Avery, N. C., and A. J. Bailey. 2005. Enzymic and non-enzymic cross-linking mechanisms in relation to turnover of collagen: relevance to aging and exercise. Scand. J. Med. Sci. Sports 15:231–240.

Bailey, A. J., R. G. Paul, and L. Knott. 1998. Mechanisms of maturation and ageing of collagen. Mech. Ageing Dev. 106:1–56.

Baynes, J. W. 2000. From life to death—the struggle between chemistry and biology during aging: the Maillard reaction as an amplifier of genomic damage. Biogerontology 1:235–246.

Boor, P., P. Celec, M. Behuliak, P. Grancic, A. Kebis, M. Kukan, et al. 2009. Regular moderate exercise reduces advanced glycation and ameliorates early diabetic nephropathy in obese Zucker rats. Metabolism 58:1669–1677.

Brownlee, M. 2001. Biochemistry and molecular cell biology of diabetic complications. Nature 414:813–820.

Cai, W., J. C. He, L. Zhu, X. Chen, S. Wallenstein, G. E. Striker, et al. 2007. Reduced oxidant stress and extended lifespan in mice exposed to a low glycoxyx diet: association with increased AGER1 expression. Am. J. Pathol. 170:1893–1902.

Couppe, C., R. B. Svensson, J. F. Grosset, V. Kovanen, R. H. Nielsen, M. R. Olsen, et al. 2014. Life-long endurance running is associated with reduced glycation and mechanical stress in connective tissue. Age (Dordr) 36: 9665.

Dunn, J. A., D. R. McCance, S. R. Thorpe, T. J. Lyons, and J. W. Baynes. 1991. Age-dependent accumulation of N epsilon-(carboxymethyl)lysine and N epsilon-(carboxymethyl)hydroxylysine in human skin collagen. Biochemistry 30:1205–1210.

Eliasson, P., A. Fahlgren, B. Pastersk, and P. Aspenberg. 2007. Unloaded rat Achilles tendons continue to grow, but lose viscoelasticity. J. Appl. Physiol. 103:459–463.

Eliasson, P., T. Andersson, J. Kulas, P. Seemann, and P. Aspenberg. 2009. Myostatin in tendon maintenance and repair. Growth Factors 27:247–254.

Erikson, C., R. B. Svensson, J. Scheijen, A. M. Mag, C. Schalkwijk, S. Paeft, et al. 1985. Systemic Stiffening of Mouse Tail Tendon is related to Dietary Advanced Glycation End-products but not High Fat Diet or Cholesterol. J. Appl. Physiol. 2014.

Eyre, D. 1987. Collagen cross-linking amino acids. Methods Enzymol. 144:115–139.

Fessel, G., Y. Li, V. Diederich, M. Guizar-Sicairos, P. Schneider, D. R. Sell, et al. 2014. Advanced glycation end-products reduce collagen molecular sliding to affect collagen fibril damage mechanisms but not stiffness. PLoS ONE 9:e110948.

Grasa, J., B. Calvo, C. Delgado-Andrade, and M. P. Navarro. 2013. Variations in tendon stiffness due to diets with different glycoxins affect mechanical properties in the muscle-tendon unit. Ann. Biomed. Eng. 41:488–496.

Hag, A. M., S. F. Pedersen, C. Christoffersen, T. Binderup, M. M. Jensen, J. T. Jorgensen, et al. 2012. (18)F-FDG PET imaging of murine atherosclerosis: association with gene expression of key molecular markers. PLoS ONE 7:e50908.

Hanssen, N. M., L. Engelen, I. Ferreira, J. L. Scheijen, M. S. Huijberts, M. J. T. Jorgensen, et al. 2013. Plasma levels of advanced glycation endproducts Neptosilin-(carboxymethyl)lysine, Neptosilin-(carboxyethyl)lysine, and pentosidine are not independently associated with cardiovascular disease in individuals with or without type 2 diabetes: the Hoorn and CODAM studies. J. Clin. Endocrinol. Metab. 98:E1369–E1373.

Hanssen, N. M., K. Wouters, M. S. Huijberts, M. J. Gijbels, J. C. Sluimer, J. L. Scheijen, et al. 2014. Higher levels of advanced glycation endproducts in human carotid atherosclerotic plaques are associated with a rupture-prone phenotype. Eur. Heart J. 35:1137–1146.

Heinemeier, K. M., P. Schjerling, J. Heinemeier, S. P. Magnusson, and M. Kjaer. 2013. Lack of tissue renewal in...
human adult Achilles tendon is revealed by nuclear bomb
(14)C. FASEB J. 27:2074–2079.
de Jonge, S., C. van den Berg, R. J. de Vos, H. J. van der
Heide, A. Weir, J. A. Verhaar, et al. 2011. Incidence of midportion Achilles tendinopathy in the general population.
Br. J. Sports Med. 45:1026–1028.
de Jonge, S., R. Rozenberg, B. Vieyra, H. J. Stam, H. J.
Aanstoot, H. Weinsans, et al. 2015. Achilles tendons in
people with type 2 diabetes show mildly compromised
structure: an ultrasound tissue characterisation study. Br. J.
Sports Med. 49:999–999.
Kjaer, M. 2004. Role of extracellular matrix in adaptation of
tendon and skeletal muscle to mechanical loading. Physiol.
Rev. 84:649–698.
Kongsgaard, M., V. Kovonen, P. Aagaard, S. Doessing, P.
Hansen, A. H. Laursen, et al. 2009. Corticosteroid
injections, eccentric decline squat training and heavy slow
resistance training in patellar tendinopathy. Scand. J. Med.
Sci. Sports 19:790–802.
Koschinsky, T., C. J. He, T. Mitsuhashi, R. Bucala, C. Liu, C.
Buening, et al. 1997. Orally absorbed reactive glycation
products (glycotoxins): an environmental risk factor in
diabetic nephropathy. Proc. Natl Acad. Sci. USA 94:6474–
6479.
Li, Y., G. Fessel, M. Georgiadis, and J. G. Snedeker. 2013.
Advanced glycation end-products diminish tendon collagen
fiber sliding. Matrix Biol. 32:169–177.
Lingelbach, L. B., A. E. Mitchell, R. B. Rucker, and R. B.
McDonald. 2000. Accumulation of advanced glycation
endproducts in aging male Fischer 344 rats during long-
term feeding of various dietary carbohydrates. J. Nutr.
130:1247–1255.
Maillard, L. C. 1912. Action des acides aminés sur les sucres:
formation des mélanoidines par voie méthodique. C. R.
Acad. Sci. 154:66–68.
Monnier, V. M., G. T. Mustata, K. L. Biemel, O. Reihl, M. O.
Lederer, D. Zheng, et al. 2005. Cross-linking of the
extracellular matrix by the maillard reaction in aging and
diabetes: an update on “a puzzle nearing resolution”. Ann.
N. Y. Acad. Sci. 1043:533–544.
Odetti, P., N. Traverso, L. Cosso, G. Noberasco, M. A.
Pronzato, and U. M. Marinari. 1996. Good glycemic
control reduces oxidation and glycation end-products in
collagen of diabetic rats. Diabetologia 39:1440–1447.
Rabbani, N., and P. J. Thorlalley. 2012. Glycation research in
amino acids: a place to call home. Amino Acids 42:1087–1096.
Ranger, T. A., M. A. Wong, J. L. Cook, and J. E. Gaida. 2016.
Is there an association between tendinopathy and diabetes
mellitus? A systematic review with meta-analysis. Br. J.
Sports Med. 50:982–989.
Roncero-Ramos, I., C. Niquet-Leridon, C. Strauch, V. M.
Monnier, F. I. Tessier, M. P. Navarro, et al. 2014. An
advanced glycation end product (AGE)-rich diet promotes
No-carboxymethyl-lysine accumulation in the cardiac tissue
and tendons of rats. J. Agric. Food Chem. 62:6001–6006.
Scheijen, J. L., M. P. van de Waarenburg, C. D. Stehouwer,
and C. G. Schalkwijk. 2009. Measurement of pentosidine in
human plasma protein by a single-column high-
performance liquid chromatography method with
fluorescence detection. J. Chromatogr. B Analyt. Technol.
Biomed. Life Sci. 877:610–614.
Sell, D. R., and V. M. Monnier. 1997. Age-related associated
tail tendon break time with tissue pentosidine in DBA/2
vs C57BL/6 mice: the effect of dietary restriction. J.
Gerontol. A Biol. Sci. Med. Sci. 52:B277–B284.
Shapiro, B. P., T. E. Owan, S. F. Mohammed, D. M. Meyer,
L. D. Mills, C. G. Schalkwijk, et al. 2008.
Advanced glycation end products accumulate in vascular
smooth muscle and modify vascular but not ventricular
properties in elderly hypertensive canines. Circulation
118:1002–1010.
Svensson, R. B., K. M. Heinemeier, C. Coupee, M. Kjaer,
and S. P. Magnusson. 2016. The effect of aging and exercise on
the tendon. J. Appl. Physiol. (1985) 126:1353–1362. jap
00328: 02016.
Thorpe, C. T., I. Streeter, G. L. Pinchbeck, A. E. Goodship, P.
D. Clegg, and H. L. Birch. 2010. Aspartic acid racemization
and collagen degradation markers reveal an accumulation of
damage in tendon collagen that is enhanced with aging. J.
Biol. Chem. 285:15674–15681.
Uribarri, J., S. Woodruff, S. Goodman, W. Cai, X. Chen, R.
Pyzik, et al. 2010. Advanced glycation end products in foods
and a practical guide to their reduction in the diet. J. Am.
Diet. Assoc. 110: 911–916e912.
Uribarri, J., W. Cai, M. Ramdas, S. Goodman, R. Pyzik, X.
Chen, et al. 2011. Restriction of advanced glycation end
products improves insulin resistance in human type 2
diabetes: potential role of AGER1 and SIRT1. Diabetes Care
34:1610–1616.
Verzijl, N., J. DeGroot, S. R. Thorpe, R. A. Bank, J. N. Shaw,
T. J. Lyons, et al. 2000. Effect of collagen turnover on the
accumulation of advanced glycation end products. J. Biol.
Chem. 275:39027–39031.
Vlassara, H., W. Cai, E. Tripp, R. Pyzik, K. Yee, L. Goldberg,
et al. 2016. Oral AGE restriction ameliorates insulin
resistance in obese individuals with the metabolic syndrome:
a randomised controlled trial. Diabetologia 59:2181–2192.
Wood, L. K., and S. V. Brooks. 2016. Ten weeks of
treadmill running decreases stiffness and increases collagen
turnover in tendons of old mice. J. Orthop. Res. 34:346–
353.
Zimmermann, E. A., E. Schaible, H. Bale, H. D. Barth, S. Y.
Tang, P. Reichert, et al. 2011. Age-related changes in
the plasticity and toughness of human cortical bone at
multiple length scales. Proc. Natl Acad. Sci. USA
108:14416–14421.