

**OBJECTIVE:** Bone contains exclusively genetic Type I collagen. These collagens are identical sequences of amino acids that form the tropocollagen molecules linked by cross-links found at the end of each tropocollagen molecules. Collagen cross-linking, a major post-translational modification of collagen, plays important roles in the biological and biomechanical features of bone. The aim of this work is to use a 3D nano-scale finite element model of a sub structure in collagen fibril called mineralized collagen microfibril and to investigate the important roles of cross-links in the expression of bone strength and its capacity and ability to absorb energy. METHOD Finite element modelling of collagen Finite element modelling of mineralzed microfibril collagen cross-links --WW--Unfolding Mineralized collagen microfibril Frictional 2πR "Delay" • 3D finite element model mesh **3D** finte element model of microfibril **Illustration of real microfibril Rheological properties** Value Source





**Buehler (2008)** 

Buehler et al. (2008)

Uzel and Buehler, (2011)

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1) Equivalent Young's modulus of collagen microfibril as a function of the cross-link number under varying Young's modulus of mineral E<sub>m</sub>



3)Equivalent fracture stress of collagen microfibril as a function of the cross-link number under varying Young's modulus of mineral E<sub>m</sub>



2) Poisson's ratio of collagen microfibril as a function of the cross-link number under varying Young's modulus of mineral Em

CONCLUSION: In this study results were obtained under tensile loading with symmetric and periodic boundary conditions. These results show that the number of cross-link has a most influence on the increase of the elastic and failure proprieties.

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4)Equivalent damping capacity of collagen microfibril as a function of the cross-link number under varying Young's modulus of mineral E<sub>m</sub>

## **REFERENCE:**

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