

Sergey Leikin

List of Publications by Year in descending order

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38742

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docs citations

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times ranked

9212
citing authors

#	ARTICLE	IF	CITATIONS
1	4PBA reduces growth deficiency in osteogenesis imperfecta by enhancing transition of hypertrophic chondrocytes to osteoblasts. JCI Insight, 2022, 7, .	5.0	16
2	Procollagen Trafficking and its Implications in Osteogenesis Imperfecta. Biology of Extracellular Matrix, 2021, , 23-53.	0.3	0
3	The regulatory role of matrix proteins in mineralization of bone. , 2021, , 165-187.		2
4	Guidelines for the use and interpretation of assays for monitoring autophagy (4th) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 622 Td (edition 9.1	9.1	1,430
5	Noncanonical ERâ€“Golgi trafficking and autophagy of endogenous procollagen in osteoblasts. Cellular and Molecular Life Sciences, 2021, 78, 8283-8300.	5.4	12
6	Mechanisms of procollagen and HSP47 sorting during ER-to-Golgi trafficking. Matrix Biology, 2020, 93, 79-94.	3.6	25
7	Substitution of murine type I collagen A1 3-hydroxylation site alters matrix structure but does not recapitulate osteogenesis imperfecta bone dysplasia. Matrix Biology, 2020, 90, 20-39.	3.6	11
8	COL1A1 C-propeptide mutations cause ER mislocalization of procollagen and impair C-terminal procollagen processing. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2019, 1865, 2210-2223.	3.8	18
9	Evidence of biomechanical and collagen heterogeneity in uterine fibroids. PLoS ONE, 2019, 14, e0215646.	2.5	32
10	Endoplasmic reticulum stress is induced in growth plate hypertrophic chondrocytes in G610C mouse model of osteogenesis imperfecta. Biochemical and Biophysical Research Communications, 2019, 509, 235-240.	2.1	33
11	Noncanonical autophagy at ER exit sites regulates procollagen turnover. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, E10099-E10108.	7.1	136
12	Substitutions for arginine at position 780 in triple helical domain of the Î±1(I) chain alter folding of the type I procollagen molecule and cause osteogenesis imperfecta. PLoS ONE, 2018, 13, e0200264.	2.5	16
13	The chaperone activity of 4PBA ameliorates the skeletal phenotype of Chihuahua, a zebrafish model for dominant osteogenesis imperfecta. Human Molecular Genetics, 2017, 26, 2897-2911.	2.9	68
14	P4HA1 mutations cause a unique congenital disorder of connective tissue involving tendon, bone, muscle and the eye. Human Molecular Genetics, 2017, 26, 2207-2217.	2.9	37
15	Celecoxib treatment of fibrous dysplasia (FD) in a human FD cell line and FD-like lesions in mice with protein kinase A (PKA) defects. Molecular and Cellular Endocrinology, 2017, 439, 165-174.	3.2	5
16	Absence of the ER Cation Channel TMEM38B/TRIC-B Disrupts Intracellular Calcium Homeostasis and Dysregulates Collagen Synthesis in Recessive Osteogenesis Imperfecta. PLoS Genetics, 2016, 12, e1006156.	3.5	49
17	Osteoblast Malfunction Caused by Cell Stress Response to Procollagen Misfolding in Î±2(I)-G610C Mouse Model of Osteogenesis Imperfecta. Journal of Bone and Mineral Research, 2016, 31, 1608-1616.	2.8	71
18	Makings of a brittle bone: Unexpected lessons from a low protein diet study of a mouse OI model. Matrix Biology, 2016, 52-54, 29-42.	3.6	15

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19	Zebrafish Collagen Type I: Molecular and Biochemical Characterization of the Major Structural Protein in Bone and Skin. <i>Scientific Reports</i> , 2016, 6, 21540.	3.3	97
20	MBTPS2 mutations cause defective regulated intramembrane proteolysis in X-linked osteogenesis imperfecta. <i>Nature Communications</i> , 2016, 7, 11920.	12.8	112
21	Genetic Defects in TAPT1 Disrupt Ciliogenesis and Cause a Complex Lethal Osteochondrodysplasia. <i>American Journal of Human Genetics</i> , 2015, 97, 521-534.	6.2	39
22	Haploinsufficiency for either one of the type-II regulatory subunits of protein kinase A improves the bone phenotype of Prkar1a+/~ mice. <i>Human Molecular Genetics</i> , 2015, 24, 6080-6092.	2.9	9
23	Abnormal Type I Collagen Post-translational Modification and Crosslinking in a Cyclophilin B KO Mouse Model of Recessive Osteogenesis Imperfecta. <i>PLoS Genetics</i> , 2014, 10, e1004465.	3.5	98
24	Pulse-chase analysis of procollagen biosynthesis by azidohomoalanine labeling. <i>Connective Tissue Research</i> , 2014, 55, 403-410.	2.3	10
25	Collagen Structure, Folding and Function. , 2014, , 71-84.		15
26	Collagen degradation by tumor-associated trypsins. <i>Archives of Biochemistry and Biophysics</i> , 2013, 535, 111-114.	3.0	11
27	Mutations in WNT1 Cause Different Forms of Bone Fragility. <i>American Journal of Human Genetics</i> , 2013, 92, 565-574.	6.2	240
28	Helical Structure Determines Different Susceptibilities of dsDNA, dsRNA, and tsDNA to Counterion-Induced Condensation. <i>Biophysical Journal</i> , 2013, 104, 2031-2041.	0.5	19
29	Kuskokwim Syndrome, a Recessive Congenital Contracture Disorder, Extends the Phenotype of FKBP10 Mutations. <i>Human Mutation</i> , 2013, 34, 1279-1288.	2.5	53
30	Deficiency of CRTAP in non-lethal recessive osteogenesis imperfecta reduces collagen deposition into matrix. <i>Clinical Genetics</i> , 2012, 82, 453-459.	2.0	30
31	Absence of FKBP10 in recessive type XI osteogenesis imperfecta leads to diminished collagen cross-linking and reduced collagen deposition in extracellular matrix. <i>Human Mutation</i> , 2012, 33, 1589-1598.	2.5	86
32	Deficient degradation of homotrimeric type I collagen, $\alpha 1(I)3$ glomerulopathy in oim mice. <i>Molecular Genetics and Metabolism</i> , 2011, 104, 373-382.	1.1	10
33	Electrostatic Braiding and Homologous Pairing of DNA Double Helices. <i>Biophysical Journal</i> , 2011, 101, 875-884.	0.5	28
34	Chaperoning osteogenesis: new protein-folding disease paradigms. <i>Trends in Cell Biology</i> , 2011, 21, 168-176.	7.9	70
35	COL1 C-propeptide cleavage site mutations cause high bone mass osteogenesis imperfecta. <i>Human Mutation</i> , 2011, 32, 598-609.	2.5	119
36	Signatures of DNA flexibility, interactions and sequence-related structural variations in classical X-ray diffraction patterns. <i>Nucleic Acids Research</i> , 2011, 39, 7289-7299.	14.5	13

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37	Fluctuations and interactions of semi-flexible polyelectrolytes in columnar assemblies. <i>Journal of Physics Condensed Matter</i> , 2010, 22, 072202.	1.8	9
38	Variable bone fragility associated with an Amish <i>COL1A2</i> variant and a knock-in mouse model. <i>Journal of Bone and Mineral Research</i> , 2010, 25, 247-261.	2.8	98
39	Carcinomas Contain a Matrix Metalloproteinase-Resistant Isoform of Type I Collagen Exerting Selective Support to Invasion. <i>Cancer Research</i> , 2010, 70, 4366-4374.	0.9	89
40	Alternate protein kinase A activity identifies a unique population of stromal cells in adult bone. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010, 107, 8683-8688.	7.1	42
41	Lack of Cyclophilin B in Osteogenesis Imperfecta with Normal Collagen Folding. <i>New England Journal of Medicine</i> , 2010, 362, 521-528.	27.0	158
42	Undulations Enhance the Effect of Helical Structure on DNA Interactions. <i>Journal of Physical Chemistry B</i> , 2010, 114, 11668-11680.	2.6	22
43	Molecular Mechanism of Type I Collagen Homotrimer Resistance to Mammalian Collagenases. <i>Journal of Biological Chemistry</i> , 2010, 285, 22276-22281.	3.4	100
44	In utero transplantation of adult bone marrow decreases perinatal lethality and rescues the bone phenotype in the knockin murine model for classical, dominant osteogenesis imperfecta. <i>Blood</i> , 2009, 114, 459-468.	1.4	93
45	DNA Double Helices Recognize Mutual Sequence Homology in a Protein Free Environment. <i>Journal of Physical Chemistry B</i> , 2008, 112, 1060-1064.	2.6	73
46	Segregation of Type I Collagen Homo- and Heterotrimers in Fibrils. <i>Journal of Molecular Biology</i> , 2008, 383, 122-132.	4.2	28
47	Structural Heterogeneity of Type I Collagen Triple Helix and Its Role in Osteogenesis Imperfecta. <i>Journal of Biological Chemistry</i> , 2008, 283, 4787-4798.	3.4	81
48	Helical coherence of DNA in crystals and solution. <i>Nucleic Acids Research</i> , 2008, 36, 5540-5551.	14.5	36
49	Defective C-propeptides of the Pro α 2(I) Chain of Type I Procollagen Impede Molecular Assembly and Result in Osteogenesis Imperfecta. <i>Journal of Biological Chemistry</i> , 2008, 283, 16061-16067.	3.4	57
50	Selective retention and degradation of molecules with a single mutant α 1(I) chain in the <i>Brl IV</i> mouse model of <i>OI</i> . <i>Matrix Biology</i> , 2007, 26, 604-614.	3.6	52
51	Structure and interactions of biological helices. <i>Reviews of Modern Physics</i> , 2007, 79, 943-996.	45.6	285
52	Prolyl 3-hydroxylase 1 deficiency causes a recessive metabolic bone disorder resembling lethal/severe osteogenesis imperfecta. <i>Nature Genetics</i> , 2007, 39, 359-365.	21.4	429
53	Y-position cysteine substitution in type I collagen (α 1(I) R888C/p.R1066C) is associated with osteogenesis imperfecta/Ehlers-Danlos syndrome phenotype. <i>Human Mutation</i> , 2007, 28, 396-405.	2.5	63
54	Procollagen Triple Helix Assembly: An Unconventional Chaperone-Assisted Folding Paradigm. <i>PLoS ONE</i> , 2007, 2, e1029.	2.5	56

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55	Deficiency of Cartilage-Associated Protein in Recessive Lethal Osteogenesis Imperfecta. <i>New England Journal of Medicine</i> , 2006, 355, 2757-2764.	27.0	307
56	Molecular Mechanism of $\hat{I}\pm 1(I)$ -Osteogenesis Imperfecta/Ehlers-Danlos Syndrome. <i>Journal of Biological Chemistry</i> , 2006, 281, 6463-6470.	3.4	77
57	Direct Observation of Azimuthal Correlations between DNA in Hydrated Aggregates. <i>Physical Review Letters</i> , 2005, 95, 148102.	7.8	33
58	Mutations Near Amino End of $\hat{I}\pm 1(I)$ Collagen Cause Combined Osteogenesis Imperfecta/Ehlers-Danlos Syndrome by Interference with N-propeptide Processing. <i>Journal of Biological Chemistry</i> , 2005, 280, 19259-19269.	3.4	118
59	Torsional Deformation of Double Helix in Interaction and Aggregation of DNA. <i>Journal of Physical Chemistry B</i> , 2004, 108, 6508-6518.	2.6	63
60	Interactions of Inorganic Phosphate and Sulfate Anions with Collagen. <i>Biochemistry</i> , 2004, 43, 14901-14912.	2.5	67
61	Structure, stability and interactions of type I collagen with GLY349-CYS substitution in $\hat{I}\pm 1(I)$ chain in a murine Osteogenesis Imperfecta model. <i>Matrix Biology</i> , 2004, 23, 101-112.	3.6	32
62	Changes in Thermal Stability and Microunfolded Pattern of Collagen Helix Resulting from the Loss of $\hat{I}\pm 2(I)$ Chain in Osteogenesis Imperfecta Murine. <i>Journal of Molecular Biology</i> , 2003, 331, 191-200.	4.2	64
63	Type I Collagen Triplet Duplication Mutation in Lethal Osteogenesis Imperfecta Shifts Register of $\hat{I}\pm$ Chains throughout the Helix and Disrupts Incorporation of Mutant Helices into Fibrils and Extracellular Matrix. <i>Journal of Biological Chemistry</i> , 2003, 278, 10006-10012.	3.4	29
64	Type I collagen is thermally unstable at body temperature. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2002, 99, 1314-1318.	7.1	488
65	Temperature-Dependent DNA Condensation Triggered by Rearrangement of Adsorbed Cations. <i>Journal of Physical Chemistry B</i> , 2002, 106, 13362-13369.	2.6	64
66	Osteogenesis Imperfecta Murine: Interaction Between Type I Collagen Homotrimers. <i>Journal of Molecular Biology</i> , 2001, 309, 807-815.	4.2	22
67	Measurement of Forces between Hydroxypropylcellulose Polymers: Temperature Favored Assembly and Salt Exclusion. <i>Journal of Physical Chemistry B</i> , 2001, 105, 1877-1886.	2.6	37
68	Sequence Recognition in the Pairing of DNA Duplexes. <i>Physical Review Letters</i> , 2001, 86, 3666-3669.	7.8	94
69	Twist in Chiral Interaction between Biological Helices. <i>Physical Review Letters</i> , 2000, 84, 2537-2540.	7.8	35
70	Electrostatic interaction between long, rigid helical macromolecules at all interaxial angles. <i>Physical Review E</i> , 2000, 62, 2576-2596.	2.1	57
71	Does the Triple Helical Domain of Type I Collagen Encode Molecular Recognition and Fiber Assembly while Telopeptides Serve as Catalytic Domains?. <i>Journal of Biological Chemistry</i> , 1999, 274, 36083-36088.	3.4	97
72	Electrostatic Zipper Motif for DNA Aggregation. <i>Physical Review Letters</i> , 1999, 82, 4138-4141.	7.8	205

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73	Sugars and Polyols Inhibit Fibrillogenesis of Type I Collagen by Disrupting Hydrogen-Bonded Water Bridges between the Helices. <i>Biochemistry</i> , 1998, 37, 11888-11895.	2.5	90
74	Symmetry Laws for Interaction between Helical Macromolecules. <i>Biophysical Journal</i> , 1998, 75, 2513-2519.	0.5	40
75	Electrostatic interaction between helical macromolecules in dense aggregates: An impetus for DNA poly- and meso-morphism. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1998, 95, 13579-13584.	7.1	91
76	Raman spectral evidence for hydration forces between collagen triple helices. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1997, 94, 11312-11317.	7.1	155
77	Solvent hydrogen-bond network in protein self-assembly: solvation of collagen triple helices in nonaqueous solvents. <i>Biophysical Journal</i> , 1997, 72, 353-362.	0.5	56
78	“Overscreening” in a polar liquid as a result of coupling between polarization and density fluctuations. <i>Electrochimica Acta</i> , 1997, 42, 849-865.	5.2	43
79	Measured effects of diacylglycerol on structural and elastic properties of phospholipid membranes. <i>Biophysical Journal</i> , 1996, 71, 2623-2632.	0.5	240
80	Temperature-favoured assembly of collagen is driven by hydrophilic not hydrophobic interactions. <i>Nature Structural and Molecular Biology</i> , 1995, 2, 205-210.	8.2	150
81	Landau Theory of a System with Two Bilinearly Coupled Order Parameter sin External Field: Exact Mean Field Solution, Critical Properties and Isothermal Susceptibility. <i>Zeitschrift Fur Naturforschung - Section A Journal of Physical Sciences</i> , 1995, 50, 789-794.	1.5	0
82	Temperature-induced complementarity as a mechanism for biomolecular assembly. <i>Proteins: Structure, Function and Bioinformatics</i> , 1994, 19, 73-76.	2.6	21
83	Bending, hydration and interstitial energies quantitatively account for the hexagonal-lamellar-hexagonal reentrant phase transition in dioleoylphosphatidylethanolamine. <i>Biophysical Journal</i> , 1994, 67, 1603-1611.	0.5	134
84	Direct measurement of forces between self-assembled proteins: temperature-dependent exponential forces between collagen triple helices.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1994, 91, 276-280.	7.1	137
85	Hydration Forces. <i>Annual Review of Physical Chemistry</i> , 1993, 44, 369-395.	10.8	489
86	Chain-melting reentrant transition in bimolecular layers at large separations. <i>Physical Review Letters</i> , 1993, 70, 3623-3626.	7.8	7
87	Surface phase transitions and hydration forces. <i>Journal of Chemical Physics</i> , 1992, 97, 6809-6820.	3.0	17
88	On the theory of electrostatic interaction of neutral lipid bilayers separated by thin water film. <i>Journal of Chemical Physics</i> , 1991, 95, 5224-5229.	3.0	12
89	Measured entropy and enthalpy of hydration as a function of distance between DNA double helices. <i>Physical Review A</i> , 1991, 44, 5272-5278.	2.5	75
90	Mean-field theory of dehydration transitions. <i>Physical Review A</i> , 1991, 44, 1156-1168.	2.5	30

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91	Theory of hydration forces. Nonlocal electrostatic interaction of neutral surfaces. Journal of Chemical Physics, 1990, 92, 6890-6898.	3.0	62
92	Stalk mechanism of vesicle fusion. European Biophysics Journal, 1989, 17, 121-9.	2.2	151
93	Elastic properties of interfaces. Elasticity moduli and spontaneous geometric characteristics. Journal of the Chemical Society, Faraday Transactions 2, 1989, 85, 277.	1.1	27
94	Fluctuation theory of hydration forces: The dramatic effects of inhomogeneous boundary conditions. Physical Review A, 1989, 40, 6431-6437.	2.5	91
95	Reversible electrical breakdown of lipid bilayers: formation and evolution of pores. Biochimica Et Biophysica Acta - Biomembranes, 1988, 940, 275-287.	2.6	528
96	Definition of surface tension at a non-spherical interface. Journal of the Chemical Society, Faraday Transactions 2, 1988, 84, 1149.	1.1	32
97	Membrane fusion: Overcoming of the hydration barrier and local restructuring. Journal of Theoretical Biology, 1987, 129, 411-425.	1.7	142
98	Localization of nonlinear waves in randomly inhomogeneous media. Physics Letters, Section A: General, Atomic and Solid State Physics, 1984, 105, 31-33.	2.1	9