

# Kathryn S E Cheah

## List of Publications by Year in descending order

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78  
papers

7,864  
citations

87888

38  
h-index

74163

75  
g-index

86  
all docs

86  
docs citations

86  
times ranked

9047  
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	Hypertrophic chondrocytes serve as a reservoir for marrow-associated skeletal stem and progenitor cells, osteoblasts, and adipocytes during skeletal development. ELife, 2022, 11, .	6.0	28
3	Hedgehog signaling orchestrates cartilage-to-bone transition independently of Smoothed. Matrix Biology, 2022, 110, 76-90.	3.6	5
4	Deciphering osteoarthritis genetics across 826,690 individuals from 9 populations. Cell, 2021, 184, 4784-4818.e17.	28.9	188
5	PRIMUS: Comprehensive proteomics of mouse intervertebral discs that inform novel biology and relevance to human disease modelling. Matrix Biology Plus, 2021, 12, 100082.	3.5	13
6	Cellular Plasticity in Musculoskeletal Development, Regeneration, and Disease. Journal of Orthopaedic Research, 2020, 38, 708-718.	2.3	4
7	$\beta$ 1 integrin regulates convergent extension in mouse notogenesis, ensures notochord integrity and the morphogenesis of vertebrae and intervertebral discs. Development (Cambridge), 2020, 147, .	2.5	2
8	Transformation of resident notochordâ€descendent nucleus pulposus cells in mouse injuryâ€induced fibrotic intervertebral discs. Aging Cell, 2020, 19, e13254.	6.7	16
9	<scp>IRX3</scp> and <scp>IRX5</scp> Inhibit Adipogenic Differentiation of Hypertrophic Chondrocytes and Promote Osteogenesis. Journal of Bone and Mineral Research, 2020, 35, 2444-2457.	2.8	31
10	Deep-learning-assisted biophysical imaging cytometry at massive throughput delineates cell population heterogeneity. Lab on A Chip, 2020, 20, 3696-3708.	6.0	41
11	Directed Differentiation of Notochord-like and Nucleus Pulposus-like Cells Using Human Pluripotent Stem Cells. Cell Reports, 2020, 30, 2791-2806.e5.	6.4	48
12	Fbxo9 functions downstream of Sox10 to determine neuron-glia fate choice in the dorsal root ganglia through Neurog2 destabilization. Proceedings of the National Academy of Sciences of the United States of America, 2020, 117, 4199-4210.	7.1	13
13	DIPPER, a spatiotemporal proteomics atlas of human intervertebral discs for exploring ageing and degeneration dynamics. ELife, 2020, 9, .	6.0	37
14	Acquisition of multipotent and migratory neural crest cells in vertebrate evolution. Current Opinion in Genetics and Development, 2019, 57, 84-90.	3.3	7
15	The extended chondrocyte lineage: implications for skeletal homeostasis and disorders. Current Opinion in Cell Biology, 2019, 61, 132-140.	5.4	20
16	Lgr5 and Col22a1 Mark Progenitor Cells in the Lineage toward Juvenile Articular Chondrocytes. Stem Cell Reports, 2019, 13, 713-729.	4.8	35
17	Unique and overlapping GLI1 and GLI2 transcriptional targets in neoplastic chondrocytes. PLoS ONE, 2019, 14, e0211333.	2.5	22
18	Quantitative Phase Imaging Flow Cytometry for Ultraâ€Largeâ€Scale Singleâ€Cell Biophysical Phenotyping. Cytometry Part A: the Journal of the International Society for Analytical Cytology, 2019, 95, 510-520.	1.5	60

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19	Mechanistic insights into skeletal development gained from genetic disorders. <i>Current Topics in Developmental Biology</i> , 2019, 133, 343-385.	2.2	17
20	Multi-ATOM: Ultrahigh-throughput single-cell quantitative phase imaging with subcellular resolution. <i>Journal of Biophotonics</i> , 2019, 12, e201800479.	2.3	34
21	Histological and reference system for the analysis of mouse intervertebral disc. <i>Journal of Orthopaedic Research</i> , 2018, 36, 233-243.	2.3	72
22	Early onset of disc degeneration in SM/J mice is associated with changes in ion transport systems and fibrotic events. <i>Matrix Biology</i> , 2018, 70, 123-139.	3.6	41
23	Synergistic co-regulation and competition by a SOX9-GLI-FOXA phasic transcriptional network coordinate chondrocyte differentiation transitions. <i>PLoS Genetics</i> , 2018, 14, e1007346.	3.5	56
24	Reprogramming of Mouse Calvarial Osteoblasts into Induced Pluripotent Stem Cells. <i>Stem Cells International</i> , 2018, 2018, 1-11.	2.5	0
25	Inhibiting the integrated stress response pathway prevents aberrant chondrocyte differentiation thereby alleviating chondrodysplasia. <i>ELife</i> , 2018, 7, .	6.0	59
26	Asymmetric localization of DLC1 defines avian trunk neural crest polarity for directional delamination and migration. <i>Nature Communications</i> , 2017, 8, 1185.	12.8	16
27	Activating the unfolded protein response in osteocytes causes hyperostosis consistent with craniodiaphyseal dysplasia. <i>Human Molecular Genetics</i> , 2017, 26, 4572-4587.	2.9	28
28	SOXE neofunctionalization and elaboration of the neural crest during chordate evolution. <i>Scientific Reports</i> , 2016, 6, 34964.	3.3	16
29	Label-Free Quantitative Proteomics Reveals Survival Mechanisms Developed by Hypertrophic Chondrocytes under ER Stress. <i>Journal of Proteome Research</i> , 2016, 15, 86-99.	3.7	14
30	SOXE transcription factors form selective dimers on non-compact DNA motifs through multifaceted interactions between dimerization and high-mobility group domains. <i>Scientific Reports</i> , 2015, 5, 10398.	3.3	65
31	Fate of growth plate hypertrophic chondrocytes: Death or lineage extension?. <i>Development Growth and Differentiation</i> , 2015, 57, 179-192.	1.5	90
32	Interplay between Genetic Risk Factors and Protective Mechanisms for Intervertebral Disc Degeneration in Mice. <i>Global Spine Journal</i> , 2015, 5, s-0035-1554500-s-0035-1554500.	2.3	0
33	Predicting the spatiotemporal dynamics of hair follicle patterns in the developing mouse. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014, 111, 2596-2601.	7.1	31
34	The chondrocytic journey in endochondral bone growth and skeletal dysplasia. <i>Birth Defects Research Part C: Embryo Today Reviews</i> , 2014, 102, 52-73.	3.6	67
35	Hypertrophic chondrocytes can become osteoblasts and osteocytes in endochondral bone formation. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014, 111, 12097-12102.	7.1	589
36	A meta-analysis identifies adolescent idiopathic scoliosis association with <i>LBX1</i> locus in multiple ethnic groups. <i>Journal of Medical Genetics</i> , 2014, 51, 401-406.	3.2	79

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37	Lumbar disc degeneration is linked to a carbohydrate sulfotransferase 3 variant. <i>Journal of Clinical Investigation</i> , 2013, 123, 4909-4917.	8.2	126
38	SNP rs11190870 near LBX1 is associated with adolescent idiopathic scoliosis in southern Chinese. <i>Journal of Human Genetics</i> , 2012, 57, 244-246.	2.3	64
39	Exhaustion of nucleus pulposus progenitor cells with ageing and degeneration of the intervertebral disc. <i>Nature Communications</i> , 2012, 3, 1264.	12.8	357
40	Indian hedgehog mutations causing brachydactyly type A1 impair Hedgehog signal transduction at multiple levels. <i>Cell Research</i> , 2011, 21, 1343-1357.	12.0	31
41	SOX9 Governs Differentiation Stage-Specific Gene Expression in Growth Plate Chondrocytes via Direct Concomitant Transactivation and Repression. <i>PLoS Genetics</i> , 2011, 7, e1002356.	3.5	174
42	The developmental roles of the extracellular matrix: beyond structure to regulation. <i>Cell and Tissue Research</i> , 2010, 339, 93-110.	2.9	144
43	Loss of procollagen IIA from the anterior mesendoderm disrupts the development of mouse embryonic forebrain. <i>Developmental Dynamics</i> , 2010, 239, 2319-2329.	1.8	22
44	In vivo cellular adaptation to ER stress: survival strategies with double-edged consequences. <i>Journal of Cell Science</i> , 2010, 123, 2145-2154.	2.0	120
45	Utility of HoxB2 enhancer-mediated Cre activity for functional studies in the developing inner ear. <i>Genesis</i> , 2009, 47, 361-365.	1.6	10
46	Genome-Wide Haplotype Association Mapping in Mice Identifies a Genetic Variant in <i>CER1</i> Associated With BMD and Fracture in Southern Chinese Women. <i>Journal of Bone and Mineral Research</i> , 2009, 24, 1013-1021.	2.8	21
47	Association between promoter -1607 polymorphism of MMP1 and Lumbar Disc Disease in Southern Chinese. <i>BMC Medical Genetics</i> , 2008, 9, 38.	2.1	44
48	Association of the Asporin D14 Allele with Lumbar-Disc Degeneration in Asians. <i>American Journal of Human Genetics</i> , 2008, 82, 744-747.	6.2	132
49	The molecular and cellular basis of exostosis formation in hereditary multiple exostoses. <i>International Journal of Experimental Pathology</i> , 2008, 89, 321-331.	1.3	35
50	Highly efficient deletion method for the engineering of plasmid DNA with single-stranded oligonucleotides. <i>BioTechniques</i> , 2008, 44, 217-224.	1.8	2
51	Surviving Endoplasmic Reticulum Stress Is Coupled to Altered Chondrocyte Differentiation and Function. <i>PLoS Biology</i> , 2007, 5, e44.	5.6	167
52	Association of the Taq I Allele in Vitamin D Receptor With Degenerative Disc Disease and Disc Bulge in a Chinese Population. <i>Spine</i> , 2006, 31, 1143-1148.	2.0	123
53	The TRP2 Allele of COL9A2 is an Age-Dependent Risk Factor for the Development and Severity of Intervertebral Disc Degeneration. <i>Spine</i> , 2005, 30, 2735-2742.	2.0	124
54	Genomic instability in laminopathy-based premature aging. <i>Nature Medicine</i> , 2005, 11, 780-785.	30.7	579

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55	Sox2 is required for sensory organ development in the mammalian inner ear. <i>Nature</i> , 2005, 434, 1031-1035.	27.8	485
56	Misfolding of Collagen X Chains Harboring Schmid Metaphyseal Chondrodysplasia Mutations Results in Aberrant Disulfide Bond Formation, Intracellular Retention, and Activation of the Unfolded Protein Response. <i>Journal of Biological Chemistry</i> , 2005, 280, 15544-15552.	3.4	58
57	Increased efficiency of oligonucleotide-mediated gene repair through slowing replication fork progression. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005, 102, 2508-2513.	7.1	59
58	Mammary gland-specific secretion of biologically active immunosuppressive agent cytotoxic-T-lymphocyte antigen 4 human immunoglobulin fusion protein (CTLA4Ig) in milk by transgenesis. <i>Journal of Immunological Methods</i> , 2003, 277, 171-183.	1.4	11
59	An externally fixed femoral fracture model for mice. <i>Journal of Orthopaedic Research</i> , 2003, 21, 685-690.	2.3	53
60	Identification of factors influencing strand bias in oligonucleotide-mediated recombination in <i>Escherichia coli</i> . <i>Nucleic Acids Research</i> , 2003, 31, 6674-6687.	14.5	90
61	The TATA-containing core promoter of the type II collagen gene (COL2A1) is the target of interferon-gamma-mediated inhibition in human chondrocytes: requirement for Stat1alpha, Jak1 and Jak2. <i>Biochemical Journal</i> , 2003, 369, 103-115.	3.7	56
62	Circling, Deafness, and Yellow Coat Displayed by Yellow Submarine (Ysb) and Light Coat and Circling (Lcc) Mice with Mutations on Chromosome 3. <i>Genomics</i> , 2002, 79, 777-784.	2.9	26
63	Chondrocyte antigen expression, immune response and susceptibility to arthritis. <i>International Immunology</i> , 2001, 13, 421-429.	4.0	4
64	Requirement for <i>Pbx1</i> in skeletal patterning and programming chondrocyte proliferation and differentiation. <i>Development (Cambridge)</i> , 2001, 128, 3543-3557.	2.5	266
65	Mechanism of Regulatory Target Selection by the SOX High-Mobility-Group Domain Proteins as Revealed by Comparison of SOX1/2/3 and SOX9. <i>Molecular and Cellular Biology</i> , 1999, 19, 107-120.	2.3	165
66	Disrupted expression of matrix genes in the growth plate of the mouse cartilage matrix deficiency ( <i>cmd</i> ) mutant. , 1998, 22, 349-358.		42
67	Different cis-Regulatory DNA Elements Mediate Developmental Stage- and Tissue-specific Expression of the Human COL2A1 Gene in Transgenic Mice. <i>Journal of Cell Biology</i> , 1998, 141, 1291-1300.	5.2	56
68	Abnormal Compartmentalization of Cartilage Matrix Components in Mice Lacking Collagen X: Implications for Function. <i>Journal of Cell Biology</i> , 1997, 136, 459-471.	5.2	188
69	SOX9 Binds DNA, Activates Transcription, and Coexpresses with Type II Collagen during Chondrogenesis in the Mouse. <i>Developmental Biology</i> , 1997, 183, 108-121.	2.0	640
70	SOX9 directly regulates the type-II collagen gene. <i>Nature Genetics</i> , 1997, 16, 174-178.	21.4	847
71	The Human $\alpha 2(\text{XI})$ Collagen Gene (COL11A2): Completion of Coding Information, Identification of the Promoter Sequence, and Precise Localization within the Major Histocompatibility Complex Reveal Overlap with the KE5 Gene. <i>Genomics</i> , 1996, 32, 401-412.	2.9	25
72	Extensive Alternative Splicing within the Amino-propeptide Coding Domain of $\alpha 2(\text{XI})$ Procollagen mRNAs. <i>Journal of Biological Chemistry</i> , 1996, 271, 16945-16951.	3.4	15

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73	Characterization of the Complete Genomic Structure of the Human WNT-5A Gene, Functional Analysis of its Promoter, Chromosomal Mapping, and Expression in Early Human Embryogenesis. <i>Journal of Biological Chemistry</i> , 1995, 270, 31225-31234.	3.4	46
74	Tissue-Specific and differential expression of alternatively spliced $\alpha 1(\text{II})$ collagen mRNAs in early human embryos. <i>Developmental Dynamics</i> , 1995, 203, 198-211.	1.8	94
75	Autosomal dominant and recessive osteochondrodysplasias associated with the COL11A2 locus. <i>Cell</i> , 1995, 80, 431-437.	28.9	390
76	Further evidence that the failure to cleave the aminopropeptide of type I procollagen is the cause of Ehlers-Danlos syndrome type VII. <i>Human Mutation</i> , 1994, 3, 358-364.	2.5	20
77	Influence of digits, ectoderm, and retinoic acid on chondrogenesis by mouse interdigital mesoderm in culture. <i>Developmental Dynamics</i> , 1994, 201, 297-309.	1.8	29
78	Intron-exon structure, alternative use of promoter and expression of the mouse collagen X gene, Col10a-1. <i>FEBS Journal</i> , 1993, 213, 99-111.	0.2	60