

Gretchen J Mahler

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5882782/publications.pdf>

Version: 2024-02-01

36
papers

2,236
citations

377584

21
h-index

406436

35
g-index

38
all docs

38
docs citations

38
times ranked

3795
citing authors

#	ARTICLE	IF	CITATIONS
1	Microfluidic modeling of the glomerulus and tubular apparatus. , 2022, , 353-366.		0
2	Shear and endothelial induced late-stage calcific aortic valve disease-on-a-chip develops calcium phosphate mineralizations. Lab on A Chip, 2022, 22, 1374-1385.	3.1	6
3	Modelling Renal Filtration and Reabsorption Processes in a Human Glomerulus and Proximal Tubule Microphysiological System. Micromachines, 2021, 12, 983.	1.4	8
4	Critical Considerations for the Design of Multi-Organ Microphysiological Systems (MPS). Frontiers in Cell and Developmental Biology, 2021, 9, 721338.	1.8	17
5	Chondroitin Sulfate Promotes Interstitial Cell Activation and Calcification in an In Vitro Model of the Aortic Valve. Cardiovascular Engineering and Technology, 2021, , 1.	0.7	4
6	Intra-amniotic administration (Gallus gallus) of TiO ₂ , SiO ₂ , and ZnO nanoparticles affect brush border membrane functionality and alters gut microflora populations. Food and Chemical Toxicology, 2020, 135, 110896.	1.8	16
7	The role of metal oxide nanoparticles, Escherichia coli, and Lactobacillus rhamnosus on small intestinal enzyme activity. Environmental Science: Nano, 2020, 7, 3940-3964.	2.2	11
8	Biofluidic Permeable Electronics: Electronic ECM: A Permeable Microporous Elastomer for an Advanced Bio-Integrated Continuous Sensing Platform (Adv. Mater. Technol. 7/2020). Advanced Materials Technologies, 2020, 5, 2070043.	3.0	1
9	Bacteria Remediate the Effects of Food Additives on Intestinal Function in an in vitro Model of the Gastrointestinal Tract. Frontiers in Nutrition, 2020, 7, 131.	1.6	10
10	TiO ₂ Nanoparticles and Commensal Bacteria Alter Mucus Layer Thickness and Composition in a Gastrointestinal Tract Model. Small, 2020, 16, e2000601.	5.2	29
11	Electronic ECM: A Permeable Microporous Elastomer for an Advanced Bio-Integrated Continuous Sensing Platform. Advanced Materials Technologies, 2020, 5, 2000242.	3.0	14
12	A human proximal tubule-on-a-chip to study renal disease and toxicity. Biomicrofluidics, 2019, 13, 014107.	1.2	39
13	ZnO nanoparticles affect nutrient transport in an in vitro model of the small intestine. Food and Chemical Toxicology, 2019, 124, 112-127.	1.8	26
14	Endothelial barrier dysfunction induced by nanoparticle exposure through actin remodeling via caveolae/raft-regulated calcium signalling. NanoImpact, 2018, 11, 82-91.	2.4	22
15	Silicon dioxide nanoparticle exposure affects small intestine function in an in vitro model. Nanotoxicology, 2018, 12, 485-508.	1.6	63
16	Titanium dioxide nanoparticle exposure alters metabolic homeostasis in a cell culture model of the intestinal epithelium and Drosophila melanogaster. Nanotoxicology, 2018, 12, 390-406.	1.6	46
17	Effect of dietary additives on intestinal permeability in both Drosophila and a human cell co-culture. DMM Disease Models and Mechanisms, 2018, 11, .	1.2	34
18	A novel microfluidic device to model the human proximal tubule and glomerulus. RSC Advances, 2017, 7, 4216-4225.	1.7	31

#	ARTICLE	IF	CITATIONS
19	Titanium dioxide nanoparticle ingestion alters nutrient absorption in an in vitro model of the small intestine. <i>NanoImpact</i> , 2017, 5, 70-82.	2.4	136
20	Endothelial to mesenchymal transformation is induced by altered extracellular matrix in aortic valve endothelial cells. <i>Journal of Biomedical Materials Research - Part A</i> , 2017, 105, 2729-2741.	2.1	40
21	Nanoparticle size-specific actin rearrangement and barrier dysfunction of endothelial cells. <i>Nanotoxicology</i> , 2017, 11, 846-856.	1.6	27
22	The role of shear stress and altered tissue properties on endothelial to mesenchymal transformation and tumor-endothelial cell interaction. <i>Biomicrofluidics</i> , 2017, 11, 044104.	1.2	34
23	Body-on-a-Chip Systems for Animal-free Toxicity Testing. <i>ATLA Alternatives To Laboratory Animals</i> , 2016, 44, 469-478.	0.7	12
24	Modeling Barrier Tissues In Vitro: Methods, Achievements, and Challenges. <i>EBioMedicine</i> , 2016, 5, 30-39.	2.7	94
25	Shear stress magnitude and transforming growth factor- β 1 regulate endothelial to mesenchymal transformation in a three-dimensional culture microfluidic device. <i>RSC Advances</i> , 2016, 6, 85457-85467.	1.7	14
26	Detection of outer membrane vesicles in <i>Synechocystis</i> PCC 6803. <i>FEMS Microbiology Letters</i> , 2015, 362, fmv163.	0.7	18
27	Effects of shear stress pattern and magnitude on mesenchymal transformation and invasion of aortic valve endothelial cells. <i>Biotechnology and Bioengineering</i> , 2014, 111, 2326-2337.	1.7	110
28	Body-on-a-chip simulation with gastrointestinal tract and liver tissues suggests that ingested nanoparticles have the potential to cause liver injury. <i>Lab on A Chip</i> , 2014, 14, 3081-3092.	3.1	225
29	Inflammatory Cytokines Promote Mesenchymal Transformation in Embryonic and Adult Valve Endothelial Cells. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2013, 33, 121-130.	1.1	176
30	Paracellular Transport of Soybean β -Conglycinin using a Caco-2/HT29-MTX Co-Culture. <i>FASEB Journal</i> , 2013, 27, 794.10.	0.2	0
31	Oral exposure to polystyrene nanoparticles affects iron absorption. <i>Nature Nanotechnology</i> , 2012, 7, 264-271.	15.6	293
32	Inflammatory Regulation of Valvular Remodeling: The Good(?), the Bad, and the Ugly. <i>International Journal of Inflammation</i> , 2011, 2011, 1-13.	0.9	41
33	Aortic valve disease and treatment: The need for naturally engineered solutions. <i>Advanced Drug Delivery Reviews</i> , 2011, 63, 242-268.	6.6	168
34	Cardiac developmental toxicity. <i>Birth Defects Research Part C: Embryo Today Reviews</i> , 2011, 93, 291-297.	3.6	18
35	Characterization of a gastrointestinal tract microscale cell culture analog used to predict drug toxicity. <i>Biotechnology and Bioengineering</i> , 2009, 104, 193-205.	1.7	199
36	Characterization of Caco-2 and HT29-MTX cocultures in an in vitro digestion/cell culture model used to predict iron bioavailability. <i>Journal of Nutritional Biochemistry</i> , 2009, 20, 494-502.	1.9	246