Dennis Schade

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/5036192/publications.pdf

Version: 2024-02-01

47 1,266 18 34 papers citations h-index g-index

50 50 50 2274 all docs docs citations times ranked citing authors

#	Article	IF	CITATIONS
1	Differentiation of cardiomyocytes and generation of human engineered heart tissue. Nature Protocols, 2017, 12, 1177-1197.	5.5	197
2	Small Molecule-Mediated TGF- \hat{l}^2 Type II Receptor Degradation Promotes Cardiomyogenesis in Embryonic Stem Cells. Cell Stem Cell, 2012, 11, 242-252.	5.2	119
3	Cell Permeable Stapled Peptide Inhibitor of Wnt Signaling that Targets β-Catenin Protein-Protein Interactions. Cell Chemical Biology, 2017, 24, 958-968.e5.	2.5	92
4	Stepwise Clearance of Repressive Roadblocks Drives Cardiac Induction in Human ESCs. Cell Stem Cell, 2016, 18, 341-353.	5 . 2	89
5	Reduction of Nï‰-hydroxy-L-arginine by the mitochondrial amidoxime reducing component (mARC). Biochemical Journal, 2011, 433, 383-391.	1.7	80
6	Development of Novel Potent Orally Bioavailable Oseltamivir Derivatives Active against Resistant Influenza A. Journal of Medicinal Chemistry, 2014, 57, 759-769.	2.9	77
7	Wnt Inhibition Correlates with Human Embryonic Stem Cell Cardiomyogenesis: A Structure–Activity Relationship Study Based on Inhibitors for the Wnt Response. Journal of Medicinal Chemistry, 2012, 55, 697-708.	2.9	63
8	Synthesis and SAR of $\langle i \rangle b \langle i \rangle$ -Annulated 1,4-Dihydropyridines Define Cardiomyogenic Compounds as Novel Inhibitors of TGF1² Signaling. Journal of Medicinal Chemistry, 2012, 55, 9946-9957.	2.9	62
9	Discovery of Inhibitor of Wnt Production 2 (IWP-2) and Related Compounds As Selective ATP-Competitive Inhibitors of Casein Kinase 1 (CK1) Î/ε. Journal of Medicinal Chemistry, 2018, 61, 4087-4102.	2.9	42
10	A protein tertiary structure mimetic modulator of the Hippo signalling pathway. Nature Communications, 2020, 11, 5425.	5.8	38
11	Small Molecules Targeting <i>in Vivo</i> Tissue Regeneration. ACS Chemical Biology, 2014, 9, 57-71.	1.6	36
12	Modulating the NO generating system from a medicinal chemistry perspective: Current trends and therapeutic options in cardiovascular disease., 2010, 126, 279-300.		33
13	Structure–activity relationship of novel and known inhibitors of human dimethylarginine dimethylaminohydrolase-1: Alkenyl-amidines as new leads. Bioorganic and Medicinal Chemistry, 2008, 16, 10205-10209.	1.4	30
14	Synthetic Approaches to <i>N</i> ^Î -Methylated <scp>l</scp> -Arginine, <i>N</i> ^{I‰} -Hydroxy- <scp>l</scp> -arginine, <scp>l</scp> -Citrulline, and <i>N</i> ^Î -Cyano- <scp>l-Cyano-<scp>l-Organic Chemistry, 2008, 73, 1025-1030.</scp></scp>	1.7	24
15	Design, synthesis and 3D-QSAR studies of novel 1,4-dihydropyridines as $TGF\hat{l}^2/Smad$ inhibitors. European Journal of Medicinal Chemistry, 2015, 95, 249-266.	2.6	23
16	Medicinal Chemistry Approaches to Heart Regeneration. Journal of Medicinal Chemistry, 2015, 58, 9451-9479.	2.9	22
17	New Prodrugs of the Antiprotozoal Drug Pentamidine. ChemMedChem, 2011, 6, 2233-2242.	1.6	21
18	Zanamivir Amidoxime- and N-Hydroxyguanidine-Based Prodrug Approaches to Tackle Poor Oral Bioavailability. Journal of Pharmaceutical Sciences, 2015, 104, 3208-3219.	1.6	19

#	Article	IF	CITATIONS
19	Prodrug design for the potent cardiovascular agent Nï‰-hydroxy-l-arginine (NOHA): Synthetic approaches and physicochemical characterization. Organic and Biomolecular Chemistry, 2011, 9, 5249.	1.5	15
20	Nδ-Methylated l-arginine derivatives and their effects on the nitric oxide generating system. Bioorganic and Medicinal Chemistry, 2008, 16, 2305-2312.	1.4	13
21	Synthesis and biological evaluation of I-valine-amidoximeesters as double prodrugs of amidines. Bioorganic and Medicinal Chemistry, 2011, 19, 1907-1914.	1.4	13
22	Combined Proteomic and In Silico Target Identification Reveal a Role for 5-Lipoxygenase in Developmental Signaling Pathways. Cell Chemical Biology, 2018, 25, 1095-1106.e23.	2.5	13
23	Tetrahydroindoles as Multipurpose Screening Compounds and Novel Sirtuin Inhibitors. ChemMedChem, 2019, 14, 853-864.	1.6	13
24	The Peptidylglycine αâ€Amidating Monooxygenase (PAM): A Novel Prodrug Strategy for Amidoximes and <i>N</i> â€Hydroxyguanidines?. ChemMedChem, 2009, 4, 1595-1599.	1.6	12
25	Pentafluoroâ€3â€hydroxyâ€pentâ€2â€enâ€1â€ones Potently Inhibit FNTâ€Type Lactate Transporters from all Five Humanâ€Pathogenic <i>Plasmodium</i> Species. ChemMedChem, 2021, 16, 1283-1289.	21.6	12
26	Toward Secondâ€Generation Cardiomyogenic and Antiâ€cardiofibrotic 1,4â€Dihydropyridineâ€Class TGFβ Inhibitors. ChemMedChem, 2019, 14, 810-822.	1.6	11
27	Crataegus Extract WS®1442 Stimulates Cardiomyogenesis and Angiogenesis From Stem Cells: A Possible New Pharmacology for Hawthorn?. Frontiers in Pharmacology, 2019, 10, 1357.	1.6	11
28	Designing modulators of dimethylarginine dimethylaminohydrolase (DDAH): A focus on selectivity over arginase. Journal of Enzyme Inhibition and Medicinal Chemistry, 2012, 27, 24-28.	2.5	9
29	Platform for determining the inhibition profile of neuraminidase inhibitors in an influenza virus N1 background. Journal of Virological Methods, 2016, 237, 192-199.	1.0	7
30	Probing Embryonic Development Enables the Discovery of Unique Small-Molecule Bone Morphogenetic Protein Potentiators. Journal of Medicinal Chemistry, 2022, 65, 3978-3990.	2.9	7
31	Design, Synthesis, and Bioactivation of <i>O</i> -Glycosylated Prodrugs of the Natural Nitric Oxide Precursor <i>N</i> ^ω -Hydroxy- <scp>I</scp> -arginine. Journal of Medicinal Chemistry, 2016, 59, 8030-8041.	2.9	6
32	Efficient Synthesis of Optically Pure Nï‰-Alkylated l-Arginines. Synthesis, 2008, 2008, 2391-2397.	1.2	5
33	Metabolism and distribution of two highly potent and selective peptidomimetic inhibitors of matriptase. Xenobiotica, 2010, 40, 93-101.	0.5	5
34	Small-molecule probe reveals a kinase cascade that links stress signaling to TCF/LEF and Wnt responsiveness. Cell Chemical Biology, 2021, 28, 625-635.e5.	2.5	5
35	Unique photoaffinity probes to study $TGF\hat{l}^2$ signaling and receptor fates. Chemical Communications, 2019, 55, 4323-4326.	2.2	3
36	Discovery of <i>N</i> -(4-Aminobutyl)- <i>N</i> ′-(2-methoxyethyl)guanidine as the First Selective, Nonamino Acid, Catalytic Site Inhibitor of Human Dimethylarginine Dimethylaminohydrolase-1 (<i>h</i> DDAH-1). Journal of Medicinal Chemistry, 2020, 63, 425-432.	2.9	3

#	Article	IF	CITATIONS
37	Growth factor mimetics for skin regeneration: In vitro profiling of primary human fibroblasts and keratinocytes. Archiv Der Pharmazie, 2021, 354, e2100082.	2.1	3
38	High-Throughput Screening Platform in Postnatal Heart Cells and Chemical Probe Toolbox to Assess Cardiomyocyte Proliferation. Journal of Medicinal Chemistry, 2022, 65, 1505-1524.	2.9	3
39	Phenotypic screen identifies FOXO inhibitor to counteract maturation and promote expansion of human iPS cell-derived cardiomyocytes. Bioorganic and Medicinal Chemistry, 2022, 65, 116782.	1.4	3
40	Arylazoamidoximes and Related Compounds as NOâ€modulators. Archiv Der Pharmazie, 2010, 343, 9-16.	2.1	1
41	An Efficient Synthesis of Optically Pure N Î-Monomethylated l-Arginine and l-Ornithine. Synthesis, 2016, 48, 723-729.	1.2	1
42	Higher Carbon Analogues of 1,4â€Dihydropyridines as Potent TGFβ/Smad Inhibitors. European Journal of Inorganic Chemistry, 2020, 2020, 176-181.	1.0	1
43	A phytomedicine approach to stem cell modulation for heart regeneration. Planta Medica, 2015, 81, .	0.7	1
44	Abstract 135: A Novel TGFb Selective Inhibitor Drives Cardiogenesis in Embryonic Stem Cells. Circulation Research, 2012, 111, .	2.0	0
45	Crataegus ssp. promotes late-stage cardiac differentiation and regeneration. Planta Medica, 2015, 81, .	0.7	О
46	A new pharmacology for Crataegus ssp.? The standardized extract WS®1442 promotes cardiogenesis from stem cells in vitro. Planta Medica, 2016, 81, S1-S381.	0.7	0
47	Abstract 132: Imaging-Based Assay for Screening of Cell Cycle Modifying Substances in Postnatal Cardiomyocytes. Circulation Research, 2019, 125, .	2.0	О