# Christa E MÃ¼ller

List of Publications by Year
in descending order

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|---|---|---|---|
| 465 papers | 23,532 citations | 6250 80 h-index | 17580 121 g-index |
| 482 all docs | 482 docs citations | 482 times ranked | 19630 citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | 3CL Protease Inhibitors with an Electrophilic Arylketone Moiety as Anti-SARS-CoV-2 Agents. Journal of Medicinal Chemistry, 2022, 65, 2926-2939. | 2.9 | 75 |
| 2 | Andrographolide Derivatives Target the KEAP1/NRF2 Axis and Possess Potent Antiâ€SARSâ€CoVâ€2 Activity. ChemMedChem, 2022, 17, e202100732. | 1.6 | 6 |
| 3 | Development of high-affinity fluorinated ligands for cannabinoid subtype 2 receptor, and inÂvitro evaluation of a radioactive tracer for imaging. European Journal of Medicinal Chemistry, 2022, 232, 114138. | 2.6 | 5 |
| 4 | Structureâ€“Activity Relationship of 3-Methylcytidine-5â€²-Î±,Î²-methylenediphosphates as CD73 Inhibitors. Journal of Medicinal Chemistry, 2022, 65, 2409-2433. | 2.9 | 5 |
| 5 | GPR18-Mediated Relaxation of Human Isolated Pulmonary Arteries. International Journal of Molecular Sciences, 2022, 23, 1427. | 1.8 | 7 |
| 6 | P2X4 and P2X7 are essential players in basal T cell activity and Ca <sup>2+</sup> signaling milliseconds after T cell activation. Science Advances, 2022, 8, eabl9770. | 4.7 | 20 |
| 7 | CD73 controls ocular adenosine levels and protects retina from light-induced phototoxicity. Cellular and Molecular Life Sciences, 2022, 79, 152. | 2.4 | 5 |
| 8 | International Union of Basic and Clinical Pharmacology. CXII: Adenosine Receptors: A Further Update. Pharmacological Reviews, 2022, 74, 340-372. | 7.1 | 67 |
| 9 | Discovery of P2Y<sub>2</sub> Receptor Antagonist Scaffolds through Virtual High-Throughput Screening. Journal of Chemical Information and Modeling, 2022, 62, 1538-1549. | 2.5 | 6 |
| 10 | Single Stabilizing Point Mutation Enables Highâ€Resolution Coâ€Crystal Structures of the Adenosine A<sub>2A</sub> Receptor with Preladenant Conjugates. Angewandte Chemie - International Edition, 2022, 61, . | 7.2 | 14 |
| 11 | Heterotrimeric G Protein Î±-Subunits - Structures, Peptide-Derived Inhibitors, and Mechanisms. Current Medicinal Chemistry, 2022, 29, 6359-6378. | 1.2 | 7 |
| 12 | ACS Pharmacology & Translational Science in 2022. ACS Pharmacology and Translational Science, 2022, 5, 1-2. | 2.5 | 0 |
| 13 | Chemistry and Analysis of Organic Compounds in Dinosaurs. Biology, 2022, 11, 670. | 1.3 | 11 |
| 14 | Agonist-Dependent Coupling of the Promiscuous Adenosine A<sub>2B</sub> Receptor to GÎ± Protein Subunits. ACS Pharmacology and Translational Science, 2022, 5, 373-386. | 2.5 | 8 |
| 15 | Innentitelbild: Eine einzige stabilisierende Punktmutation ermÃ¶glicht hochaufgelÃ¶ste Coâ€Kristallstrukturen des Adenosinâ€A<sub>2A</sub>â€Rezeptors mit Preladenantâ€Konjugaten (Angew.) Tj ET Q q1 1 0.78431 4 rg BT | 2.4 | 0 |
| 16 | Caffeine intake exerts dual genome-wide effects on hippocampal metabolism and learning-dependent transcription. Journal of Clinical Investigation, 2022, 132, . | 3.9 | 22 |
| 17 | Irreversible Antagonists for the Adenosine A2B Receptor. Molecules, 2022, 27, 3792. | 1.7 | 6 |
| 18 | Small-Molecule Thioesters as SARS-CoV-2 Main Protease Inhibitors: Enzyme Inhibition, Structureâ€“Activity Relationships, Antiviral Activity, and X-ray Structure Determination. Journal of Medicinal Chemistry, 2022, 65, 9376-9395. | 2.9 | 35 |

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|---|---|---|---|
| 19 | Agonists, Antagonists, and Modulators of P2X7 Receptors. Methods in Molecular Biology, 2022, , 31-52. | 0.4 | 8 |
| 20 | Apoptotic brown adipocytes enhance energy expenditure via extracellular inosine. Nature, 2022, 609, 361-368. | 13.7 | 53 |
| 21 | Sphingosine-1-phosphate induces migration of microglial cells via activation of volume-sensitive anion channels, ATP secretion and activation of purinergic receptors. Biochimica Et Biophysica Acta - Molecular Cell Research, 2021, 1868, 118915. | 1.9 | 13 |
| 22 | Update of P2X receptor properties and their pharmacology: IUPHAR Review 30. British Journal of Pharmacology, 2021, 178, 489-514. | 2.7 | 165 |
| 23 | Medicinal chemistry of P2 and adenosine receptors: Common scaffolds adapted for multiple targets. Biochemical Pharmacology, 2021, 187, 114311. | 2.0 | 29 |
| 24 | Sulfated Polysaccharides from Macroalgae Are Potent Dual Inhibitors of Human ATP-Hydrolyzing Ectonucleotidases NPP1 and CD39. Marine Drugs, 2021, 19, 51. | 2.2 | 8 |
| 25 | Thioesterase-mediated side chain transesterification generates potent Gq signaling inhibitor FR900359. Nature Communications, 2021, 12, 144. | 5.8 | 32 |
| 26 | A Cellular Assay for the Identification and Characterization of Connexin Gap Junction Modulators. International Journal of Molecular Sciences, 2021, 22, 1417. | 1.8 | 7 |
| 27 | Chemistry of porphyrins in fossil plants and animals. RSC Advances, 2021, 11, 7552-7563. | 1.7 | 26 |
| 28 | Macrocyclic Gq Protein Inhibitors FR900359 and/or YM-254890â€“Fit for Translation?. ACS Pharmacology and Translational Science, 2021, 4, 888-897. | 2.5 | 17 |
| 29 | Thermal proteome profiling identifies the membrane-bound purinergic receptor P2X4 as a target of the autophagy inhibitor indophagolin. Cell Chemical Biology, 2021, 28, 1750-1757.e5. | 2.5 | 22 |
| 30 | Effects of GPR18 Ligands on Body Weight and Metabolic Parameters in a Female Rat Model of Excessive Eating. Pharmaceuticals, 2021, 14, 270. | 1.7 | 7 |
| 31 | Targeting the Main Protease of SARSâ€CoVâ€2: From the Establishment of High Throughput Screening to the Design of Tailored Inhibitors. Angewandte Chemie - International Edition, 2021, 60, 10423-10429. | 7.2 | 95 |
| 32 | Die Hauptprotease von SARSâ€CoVâ€2 als Zielstruktur: Von der Etablierung eines Hochdurchsatzâ€Screenings zum Design maÃŸgeschneiderter Inhibitoren. Angewandte Chemie, 2021, 133, 10515-10521. | 1.6 | 3 |
| 33 | PSB 603 â€“ a known selective adenosine A2B receptor antagonist â€“ has anti-inflammatory activity in mice. Biomedicine and Pharmacotherapy, 2021, 135, 111164. | 2.5 | 21 |
| 34 | Accelerating Translation of Innovative Drugs from Bench to Patients: ACS Pharmacology & Translational Science to Evolve, Grow, and Bridge the Gap between Chemistry and Biology in Drug Research and Development. ACS Pharmacology and Translational Science, 2021, 4, 1026-1027. | 2.5 | 2 |
| 35 | Synthesis of Novel Fluorinated Xanthine Derivatives with High Adenosine A2B Receptor Binding Affinity. Pharmaceuticals, 2021, 14, 485. | 1.7 | 1 |
| 36 | Adenosine A<sub>2A</sub>R/A<sub>1</sub>R Antagonists Enabling Additional H<sub>3</sub>R Antagonism for the Treatment of Parkinsonâ€™s Disease. Journal of Medicinal Chemistry, 2021, 64, 8246-8262. | 2.9 | 6 |

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|---|---|---|---|
| 37 | Iodine-catalyzed electrophilic substitution of indoles: Synthesis of (un)symmetrical diindolylmethanes with a quaternary carbon center. Beilstein Journal of Organic Chemistry, 2021, 17, 1464-1475. | 1.3 | 5 |
| 38 | Feature-Based Molecular Networking for the Targeted Identification of G<sub>q</sub>-Inhibiting FR900359 Derivatives. Journal of Natural Products, 2021, 84, 1941-1953. | 1.5 | 7 |
| 39 | Europe Virtual Issue: Pharmacology and Translational Science in Europe. ACS Pharmacology and Translational Science, 2021, 4, 1264-1264. | 2.5 | 0 |
| 40 | Virtual Issue: Oncology, Immunology, and Immuno-oncology. ACS Pharmacology and Translational Science, 2021, 4, 1475. | 2.5 | 0 |
| 41 | The GPR18 Agonist PSB-KD-107 Exerts Endothelium-Dependent Vasorelaxant Effects. Pharmaceuticals, 2021, 14, 799. | 1.7 | 7 |
| 42 | Fintiamin: A diketopiperazine from the marine spongeâ€derived fungus Eurotium sp.. Archiv Der Pharmazie, 2021, 354, e2100206. | 2.1 | 6 |
| 43 | A group of cationic amphiphilic drugs activates MRGPRX2 and induces scratching behavior in mice. Journal of Allergy and Clinical Immunology, 2021, 148, 506-522.e8. | 1.5 | 29 |
| 44 | New Drug Modalities in Medicinal Chemistry, Pharmacology, and Translational Science: Joint Virtual Special Issue by Journal of Medicinal Chemistry, ACS Medicinal Chemistry Letters, and ACS Pharmacology & Translational Science. Journal of Medicinal Chemistry, 2021, 64, 13935-13936. | 2.9 | 3 |
| 45 | Recommended tool compounds and drugs for blocking P2X and P2Y receptors. Purinergic Signalling, 2021, 17, 633-648. | 1.1 | 21 |
| 46 | New Drug Modalities in Medicinal Chemistry, Pharmacology, and Translational Science: Joint Virtual Special Issue by Journal of Medicinal Chemistry, ACS Medicinal Chemistry Letters, and ACS Pharmacology & Translational Science. ACS Medicinal Chemistry Letters, 2021, 12, 1508-1509. | 1.3 | 2 |
| 47 | New Drug Modalities in Medicinal Chemistry, Pharmacology, and Translational Science: Joint Virtual Special Issue by Journal of Medicinal Chemistry, ACS Medicinal Chemistry Letters, and ACS Pharmacology & Translational Science. ACS Pharmacology and Translational Science, 2021, 4, 1712-1713. | 2.5 | 0 |
| 48 | THE CONCISE GUIDE TO PHARMACOLOGY 2021/22: G proteinâ€coupled receptors. British Journal of Pharmacology, 2021, 178, S27-S156. | 2.7 | 337 |
| 49 | Unraveling binding mechanism and kinetics of macrocyclic GÎ±q protein inhibitors. Pharmacological Research, 2021, 173, 105880. | 3.1 | 10 |
| 50 | Structure-activity relationships of agonists for the orphan G protein-coupled receptor GPR27. European Journal of Medicinal Chemistry, 2021, 225, 113777. | 2.6 | 9 |
| 51 | An experimental strategy to probe Gq contribution to signal transduction in living cells. Journal of Biological Chemistry, 2021, 296, 100472. | 1.6 | 22 |
| 52 | Discovery of potent nucleotide pyrophosphatase/phosphodiesterase3 (NPP3) inhibitors with ancillary carbonic anhydrase inhibition for cancer (immuno)therapy. RSC Medicinal Chemistry, 2021, 12, 1187-1206. | 1.7 | 5 |
| 53 | A novel P2X2â€dependent purinergic mechanism of enteric gliosis in intestinal inflammation. EMBO Molecular Medicine, 2021, 13, e12724. | 3.3 | 35 |
| 54 | CD73-mediated adenosine production by CD8 T cell-derived extracellular vesicles constitutes an intrinsic mechanism of immune suppression. Nature Communications, 2021, 12, 5911. | 5.8 | 66 |

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| 55 | 2â€Substituted thienotetrahydropyridine derivatives: Allosteric ectonucleotidase inhibitors. Archiv Der Pharmazie, 2021, 354, e2100300. | 2.1 | 4 |
| 56 | Involvement of GPR17 in Neuronal Fibre Outgrowth. International Journal of Molecular Sciences, 2021, 22, 11683. | 1.8 | 3 |
| 57 | Quantification of Degradation Products Formed during Heat Sterilization of Glucose Solutions by LC-MS/MS: Impact of Autoclaving Temperature and Duration on Degradation. Pharmaceuticals, 2021, 14, 1121. | 1.7 | 10 |
| 58 | Age-related shift in LTD is dependent on neuronal adenosine A2A receptors interplay with mGluR5 and NMDA receptors. Molecular Psychiatry, 2020, 25, 1876-1900. | 4.1 | 129 |
| 59 | An Agonist Radioligand for the Proinflammatory Lipid-Activated G Protein-Coupled Receptor GPR84 Providing Structural Insights. Journal of Medicinal Chemistry, 2020, 63, 2391-2410. | 2.9 | 21 |
| 60 | Substituted 4-phenylthiazoles: Development of potent and selective A1, A3 and dual A1/A3 adenosine receptor antagonists. European Journal of Medicinal Chemistry, 2020, 186, 111879. | 2.6 | 9 |
| 61 | Design, synthesis and biological evaluation of suramin-derived dual antagonists of the proinflammatory G protein-coupled receptors P2Y2 and GPR17. European Journal of Medicinal Chemistry, 2020, 186, 111789. | 2.6 | 10 |
| 62 | Ligand binding and activation of UTP-activated G protein-coupled P2Y2 and P2Y4 receptors elucidated by mutagenesis, pharmacological and computational studies. Biochimica Et Biophysica Acta - General Subjects, 2020, 1864, 129501. | 1.1 | 6 |
| 63 | A2A and A2B adenosine receptors: The extracellular loop 2 determines high (A2A) or low affinity (A2B) for adenosine. Biochemical Pharmacology, 2020, 172, 113718. | 2.0 | 24 |
| 64 | Cellâ€permeable highâ€affinity tracers for G<sub>q</sub> proteins provide structural insights, reveal distinct binding kinetics and identify small molecule inhibitors. British Journal of Pharmacology, 2020, 177, 1898-1916. | 2.7 | 21 |
| 65 | Sensitive LC-MS/MS Method for the Quantification of Macrocyclic GÎ±q Protein Inhibitors in Biological Samples. Frontiers in Chemistry, 2020, 8, 833. | 1.8 | 4 |
| 66 | Update of P2Y receptor pharmacology: IUPHAR Review 27. British Journal of Pharmacology, 2020, 177, 2413-2433. | 2.7 | 151 |
| 67 | 2-Substituted Î±,Î²-Methylene-ADP Derivatives: Potent Competitive Ecto-5â€²-nucleotidase (CD73) Inhibitors with Variable Binding Modes. Journal of Medicinal Chemistry, 2020, 63, 2941-2957. | 2.9 | 37 |
| 68 | Dissection of P2X4 and P2X7 Receptor Current Components in BV-2 Microglia. International Journal of Molecular Sciences, 2020, 21, 8489. | 1.8 | 15 |
| 69 | Synthesis and structure-activity relationships of cerebroside analogues as substrates of cerebroside sulphotransferase and discovery of a competitive inhibitor. Journal of Enzyme Inhibition and Medicinal Chemistry, 2020, 35, 1503-1512. | 2.5 | 4 |
| 70 | Nucleotide Analog ARL67156 as a Lead Structure for the Development of CD39 and Dual CD39/CD73 Ectonucleotidase Inhibitors. Frontiers in Pharmacology, 2020, 11, 1294. | 1.6 | 23 |
| 71 | Nucleotide P2Y1 receptor agonists are in vitro and in vivo prodrugs of A1/A3 adenosine receptor agonists: implications for roles of P2Y1 and A1/A3 receptors in physiology and pathology. Purinergic Signalling, 2020, 16, 543-559. | 1.1 | 17 |
| 72 | Fluorescent Probes for Ecto-5â€²-nucleotidase (CD73). ACS Medicinal Chemistry Letters, 2020, 11, 2253-2260. | 1.3 | 10 |

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| 73 | Development of Anthraquinone Derivatives as Ectonucleoside Triphosphate Diphosphohydrolase (NTPDase) Inhibitors With Selectivity for NTPDase2 and NTPDase3. Frontiers in Pharmacology, 2020, 11, 1282. | 1.6 | 12 |
| 74 | Design, Synthesis and Biological Evaluation of Highly Potent Simplified Archazolids. ChemMedChem, 2020, 15, 1348-1363. | 1.6 | 5 |
| 75 | P2Y2 Receptor Promotes High-Fat Diet-Induced Obesity. Frontiers in Endocrinology, 2020, 11, 341. | 1.5 | 23 |
| 76 | Development of a Radiofluorinated Adenosine A2B Receptor Antagonist as Potential Ligand for PET Imaging. International Journal of Molecular Sciences, 2020, 21, 3197. | 1.8 | 3 |
| 77 | Discovery of Tricyclic Xanthines as Agonists of the Cannabinoid-Activated Orphan G-Protein-Coupled Receptor GPR18. ACS Medicinal Chemistry Letters, 2020, 11, 2024-2031. | 1.3 | 16 |
| 78 | Novel, Dual Targetâ€Directed Annelated Xanthine Derivatives Acting on Adenosine Receptors and Monoamine Oxidase B. ChemMedChem, 2020, 15, 772-786. | 1.6 | 9 |
| 79 | P2Y<sub>1</sub>â€like nucleotide receptorsâ€”Structures, molecular modeling, mutagenesis, and oligomerization. Wiley Interdisciplinary Reviews: Computational Molecular Science, 2020, 10, e1464. | 6.2 | 12 |
| 80 | 8-Benzylaminoxanthine scaffold variations for selective ligands acting on adenosine A2A receptors. Design, synthesis and biological evaluation. Bioorganic Chemistry, 2020, 101, 104033. | 2.0 | 5 |
| 81 | Synthesis of Novel Potent Archazolids: Pharmacology of an Emerging Class of Anticancer Drugs. Journal of Medicinal Chemistry, 2020, 63, 1684-1698. | 2.9 | 14 |
| 82 | Discovery and Structure Relationships of Salicylanilide Derivatives as Potent, Non-acidic P2X1 Receptor Antagonists. Journal of Medicinal Chemistry, 2020, 63, 6164-6178. | 2.9 | 10 |
| 83 | Computational Investigations on the Binding Mode of Ligands for the Cannabinoid-Activated G Protein-Coupled Receptor GPR18. Biomolecules, 2020, 10, 686. | 1.8 | 13 |
| 84 | Agonists and Antagonists for Purinergic Receptors. Methods in Molecular Biology, 2020, 2041, 45-64. | 0.4 | 37 |
| 85 | Recombinant expression of ecto-nucleotide pyrophosphatase/phosphodiesterase 4 (NPP4) and development of a luminescence-based assay to identify inhibitors. Analytical Biochemistry, 2020, 603, 113774. | 1.1 | 14 |
| 86 | Extracellular adenosine reversibly inhibits the activation of human regulatory T cells and negatively influences the achievement of the operational tolerance in liver transplantation. American Journal of Transplantation, 2019, 19, 48-61. | 2.6 | 19 |
| 87 | Tools and Drugs for Purine-Binding Targetsâ€”Important Players in Inflammation and Cancer. Proceedings (mdpi), 2019, 22, . | 0.2 | 0 |
| 88 | Decarboxylative Coupling Reaction of 2â€(1Hâ€Indolâ€3â€yl)acetic Acids with Indole, Azaindole, Benzimidazole and Indazole Derivatives. Advanced Synthesis and Catalysis, 2019, 361, 4286-4293. | 2.1 | 20 |
| 89 | Xâ€Ray Coâ€Crystal Structure Guides the Way to Subnanomolar Competitive Ectoâ€5â€²â€Nucleotidase (CD73) Inhibitors for Cancer Immunotherapy. Advanced Therapeutics, 2019, 2, 1900075. | 1.6 | 33 |
| 90 | Identification of aurintricarboxylic acid as a potent allosteric antagonist of P2X1 and P2X3 receptors. Neuropharmacology, 2019, 158, 107749. | 2.0 | 38 |

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| 91 | Structural and Conformational Studies on Carboxamides of 5,6-Diaminouracilsâ€"Precursors of Biologically Active Xanthine Derivatives. Molecules, 2019, 24, 2168. | 1.7 | 2 |
| 92 | Memory deficits induced by chronic cannabinoid exposure are prevented by adenosine A2AR receptor antagonism. Neuropharmacology, 2019, 155, 10-21. | 2.0 | 21 |
| 93 | Identification of adenine-N9-(methoxy)ethyl-Î²-bisphosphonate as NPP1 inhibitor attenuates NPPase activity in human osteoarthritic chondrocytes. Purinergic Signalling, 2019, 15, 247-263. | 1.1 | 6 |
| 94 | Structureâ€"Activity Relationship of Purine and Pyrimidine Nucleotides as Ecto-5â€²-Nucleotidase (CD73) Inhibitors. Journal of Medicinal Chemistry, 2019, 62, 3677-3695. | 2.9 | 53 |
| 95 | A<sub>2B</sub> Adenosine Receptor Antagonists with Picomolar Potency. Journal of Medicinal Chemistry, 2019, 62, 4032-4055. | 2.9 | 17 |
| 96 | Fast, Efficient, and Versatile Synthesis of 6-amino-5-carboxamidouracils as Precursors for 8-Substituted Xanthines. Frontiers in Chemistry, 2019, 7, 56. | 1.8 | 14 |
| 97 | Coordination of capsule assembly and cell wall biosynthesis in Staphylococcus aureus. Nature Communications, 2019, 10, 1404. | 5.8 | 66 |
| 98 | Novel multi-target directed ligands based on annelated xanthine scaffold with aromatic substituents acting on adenosine receptor and monoamine oxidase B. Synthesis, in vitro and in silico studies. Bioorganic and Medicinal Chemistry, 2019, 27, 1195-1210. | 1.4 | 17 |
| 99 | Development of Chromen-4-one Derivatives as (Ant)agonists for the Lipid-Activated G Protein-Coupled Receptor GPR55 with Tunable Efficacy. ACS Omega, 2019, 4, 4276-4295. | 1.6 | 9 |
| 100 | Modulating P1 Adenosine Receptors in Disease Progression of SOD1G93A Mutant Mice. Neurochemical Research, 2019, 44, 1037-1042. | 1.6 | 7 |
| 101 | Adenine-(methoxy)-ethoxy-PÎ±,Î±-dithio-triphosphate inhibits pathologic calcium pyrophosphate deposition in osteoarthritic human chondrocytes. Organic and Biomolecular Chemistry, 2019, 17, 9913-9923. | 1.5 | 3 |
| 102 | Elucidating the active Î´-opioid receptor crystal structure with peptide and small-molecule agonists. Science Advances, 2019, 5, eaax9115. | 4.7 | 81 |
| 103 | Chromenones as Multineurotargeting Inhibitors of Human Enzymes. ACS Omega, 2019, 4, 22161-22168. | 1.6 | 17 |
| 104 | Koala and Wombat Gammaherpesviruses Encode the First Known Viral NTPDase Homologs and Are Phylogenetically Divergent from All Known Gammaherpesviruses. Journal of Virology, 2019, 93, . | 1.5 | 2 |
| 105 | Investigation on 2â€²,3â€²-O-Substituted ATP Derivatives and Analogs as Novel P2X3 Receptor Antagonists. ACS Medicinal Chemistry Letters, 2019, 10, 493-498. | 1.3 | 8 |
| 106 | Soluble and membrane-bound adenylate kinase and nucleotidases augment ATP-mediated inflammation in diabetic retinopathy eyes with vitreous hemorrhage. Journal of Molecular Medicine, 2019, 97, 341-354. | 1.7 | 21 |
| 107 | Diabetes-induced Neuropathic Mechanical Hyperalgesia Depends on P2X4 Receptor Activation in Dorsal Root Ganglia. Neuroscience, 2019, 398, 158-170. | 1.1 | 38 |
| 108 | Antithrombotic P2Y12 receptor antagonists: recent developments in drug discovery. Drug Discovery Today, 2019, 24, 325-333. | 3.2 | 41 |

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| 109 | P2Y receptors (version 2019.4) in the IUPHAR/BPS Guide to Pharmacology Database. IUPHAR/BPS Guide To Pharmacology CITE, 2019, 2019, . | 0.2 | 6 |
| 110 | Synthesis and structure-activity relationships of quinolinone and quinoline-based P2X7 receptor antagonists and their anti-sphere formation activities in glioblastoma cells. European Journal of Medicinal Chemistry, 2018, 151, 462-481. | 2.6 | 24 |
| 111 | Genome Editing in Neuroepithelial Stem Cells to Generate Human Neurons with High Adenosine-Releasing Capacity. Stem Cells Translational Medicine, 2018, 7, 477-486. | 1.6 | 8 |
| 112 | Mechanism underlying the contractile activity of UTP in the mammalian heart. European Journal of Pharmacology, 2018, 830, 47-58. | 1.7 | 12 |
| 113 | Fluorescent-Labeled Selective Adenosine A<sub>2B</sub> Receptor Antagonist Enables Competition Binding Assay by Flow Cytometry. Journal of Medicinal Chemistry, 2018, 61, 4301-4316. | 2.9 | 24 |
| 114 | Tools and drugs for uracil nucleotide-activated P2Y receptors. , 2018, 190, 24-80. | | 42 |
| 115 | Pharmacological evaluation of new constituents of â€œSpiceâ€: synthetic cannabinoids based on indole, indazole, benzimidazole and carbazole scaffolds. Forensic Toxicology, 2018, 36, 385-403. | 1.4 | 88 |
| 116 | 6-(Ar)Alkylamino-Substituted Uracil Derivatives: Lipid Mimetics with Potent Activity at the Orphan G Protein-Coupled Receptor 84 (GPR84). ACS Omega, 2018, 3, 3365-3383. | 1.6 | 30 |
| 117 | Adenosine A<sub>2A</sub> receptor agonists with potent antiplatelet activity. Platelets, 2018, 29, 292-300. | 1.1 | 20 |
| 118 | Structural characterization and pharmacological evaluation of the new synthetic cannabinoid CUMYLâ€PEGACLONE. Drug Testing and Analysis, 2018, 10, 597-603. | 1.6 | 37 |
| 119 | Mechanisms of the action of adenine on antiâ€allergic effects in mast cells. Immunity, Inflammation and Disease, 2018, 6, 97-105. | 1.3 | 6 |
| 120 | Heterologous Expression, Biosynthetic Studies, and Ecological Function of the Selective Gqâ€Signaling Inhibitor FR900359. Angewandte Chemie - International Edition, 2018, 57, 836-840. | 7.2 | 57 |
| 121 | Heterologe Expression, Biosynthese und Ã¶kologische Funktion des selektiven Gqâ€Signaltransduktionsinhibitors FR900359. Angewandte Chemie, 2018, 130, 844-849. | 1.6 | 5 |
| 122 | Structural Mapping of Adenosine Receptor Mutations: Ligand Binding and Signaling Mechanisms. Trends in Pharmacological Sciences, 2018, 39, 75-89. | 4.0 | 64 |
| 123 | Development of a selective and highly sensitive fluorescence assay for nucleoside triphosphate diphosphohydrolase1 (NTPDase1, CD39). Analyst, The, 2018, 143, 5417-5430. | 1.7 | 12 |
| 124 | Adenosine A2A receptor ligand recognition and signaling is blocked by A2B receptors. Oncotarget, 2018, 9, 13593-13611. | 0.8 | 77 |
| 125 | Cyclopropane-Containing Fatty Acids from the Marine Bacterium Labrenzia sp. 011 with Antimicrobial and GPR84 Activity. Marine Drugs, 2018, 16, 369. | 2.2 | 25 |
| 126 | Structure-activity relationships of imidazothiazinones and analogs as antagonists of the cannabinoid-activated orphan G protein-coupled receptor GPR18. European Journal of Medicinal Chemistry, 2018, 155, 381-397. | 2.6 | 22 |

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|---|---|---|---|
| 127 | Tritium-labeled agonists as tools for studying adenosine A2B receptors. Purinergic Signalling, 2018, 14, 223-233. | 1.1 | 16 |
| 128 | Applying Molecular Networking for the Detection of Natural Sources and Analogues of the Selective Gq Protein Inhibitor FR900359. Journal of Natural Products, 2018, 81, 1628-1635. | 1.5 | 27 |
| 129 | Medicinal Chemistry of A2B Adenosine Receptors. , 2018, , 137-168. | | 12 |
| 130 | Beneficial Effect of a Selective Adenosine A2A Receptor Antagonist in the APPswe/PS1dE9 Mouse Model of Alzheimerâ€™s Disease. Frontiers in Molecular Neuroscience, 2018, 11, 235. | 1.4 | 72 |
| 131 | 3-(2-Carboxyethyl)indole-2-carboxylic Acid Derivatives: Structural Requirements and Properties of Potent Agonists of the Orphan G Protein-Coupled Receptor GPR17. Journal of Medicinal Chemistry, 2018, 61, 8136-8154. | 2.9 | 19 |
| 132 | Understanding the Role of Adenosine A2AR Heteroreceptor Complexes in Neurodegeneration and Neuroinflammation. Frontiers in Neuroscience, 2018, 12, 43. | 1.4 | 44 |
| 133 | Preysslerâ€Popeâ€Jeannin Polyanions [NaP<sub>5</sub>W<sub>30</sub>O<sub>110</sub>]<sup>14â€“</sup> and [AgP<sub>5</sub>W<sub>30</sub>O<sub>110</sub>]<sup>14â€“</sup>: Microwaveâ€Assisted Synthesis, Structure, and Biological Activity. Zeitschrift Fur Anorganische Und Allgemeine Chemie, 2018, 644, 752-758. | 0.6 | 17 |
| 134 | General Synthesis of Unsymmetrical 3,3â€²-(Aza)diindolylmethane Derivatives. Journal of Organic Chemistry, 2018, 83, 9902-9913. | 1.7 | 35 |
| 135 | Probing Substituents in the 1- and 3-Position: Tetrahydropyrazino-Annelated Water-Soluble Xanthine Derivatives as Multi-Target Drugs With Potent Adenosine Receptor Antagonistic Activity. Frontiers in Chemistry, 2018, 6, 206. | 1.8 | 8 |
| 136 | Radiosynthesis and in vivo evaluation of a fluorine-18 labeled pyrazine based radioligand for PET imaging of the adenosine A2B receptor. Bioorganic and Medicinal Chemistry, 2018, 26, 4650-4663. | 1.4 | 17 |
| 137 | Molecular and functional interaction between GPR18 and cannabinoid CB2 G-protein-coupled receptors. Relevance in neurodegenerative diseases. Biochemical Pharmacology, 2018, 157, 169-179. | 2.0 | 47 |
| 138 | Tricyclic xanthine derivatives containing a basic substituent: adenosine receptor affinity and drug-related properties. MedChemComm, 2018, 9, 951-962. | 3.5 | 9 |
| 139 | Interaction of Approved Drugs with Synaptic Vesicle Protein 2A. Archiv Der Pharmazie, 2017, 350, 1700003. | 2.1 | 9 |
| 140 | Chronic and acute adenosine A2A receptor blockade prevents long-term episodic memory disruption caused by acute cannabinoid CB1 receptor activation. Neuropharmacology, 2017, 117, 316-327. | 2.0 | 37 |
| 141 | Prasugrel suppresses development of lithium-induced nephrogenic diabetes insipidus in mice. Purinergic Signalling, 2017, 13, 239-248. | 1.1 | 10 |
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