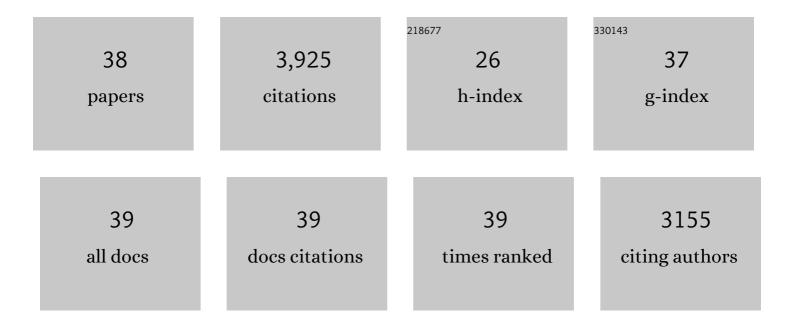
Kurt Jarnagin

List of Publications by Year in descending order

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KIIDT IADNACIN

| # | Article | lF | CITATIONS |
|----|---|------|-----------|
| 1 | Macrofilaricidal Benzimidazole–Benzoxaborole Hybrids as an Approach to the Treatment of River Blindness: Part 1. Amide Linked Analogs. ACS Infectious Diseases, 2020, 6, 173-179. | 3.8 | 11 |
| 2 | Phenotypic, chemical and functional characterization of cyclic nucleotide phosphodiesterase 4 (PDE4) as a potential anthelmintic drug target. PLoS Neglected Tropical Diseases, 2017, 11, e0005680. | 3.0 | 36 |
| 3 | Treatment of Skin Inflammation with Benzoxaborole Phosphodiesterase Inhibitors: Selectivity, Cellular Activity, and Effect on Cytokines Associated with Skin Inflammation and Skin Architecture Changes. Journal of Pharmacology and Experimental Therapeutics, 2016, 358, 413-422. | 2.5 | 55 |
| 4 | Crisaborole Topical Ointment, 2%: A Nonsteroidal, Topical, Anti-Inflammatory Phosphodiesterase 4 Inhibitor in Clinical Development for the Treatment of Atopic Dermatitis. Journal of Drugs in Dermatology, 2016, 15, 390-6. | 0.8 | 52 |
| 5 | Discovery and structure–activity relationships of 6-(benzoylamino)benzoxaboroles as orally active anti-inflammatory agents. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 5870-5873. | 2.2 | 21 |
| 6 | Structure–activity relationships of 6-(aminomethylphenoxy)-benzoxaborole derivatives as anti-inflammatory agent. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 1680-1683. | 2.2 | 25 |
| 7 | Inhibition of Toll-Like Receptor-Mediated Inflammation In Vitro and In Vivo by a Novel Benzoxaborole. Journal of Pharmacology and Experimental Therapeutics, 2013, 344, 436-446. | 2.5 | 12 |
| 8 | Linking Phenotype to Kinase: Identification of a Novel Benzoxaborole Hinge-Binding Motif for Kinase Inhibition and Development of High-Potency Rho Kinase Inhibitors. Journal of Pharmacology and Experimental Therapeutics, 2013, 347, 615-625. | 2.5 | 47 |
| 9 | Boronâ€based phosphodiesterase inhibitors show novel binding of boron to PDE4 bimetal center. FEBS Letters, 2012, 586, 3410-3414. | 2.8 | 88 |
| 10 | Early rapid identification of in vivo rat metabolites of AN6414, a novel boron-containing PDE4 inhibitor by QTRAP LC/MS/MS to support drug discovery. Journal of Pharmaceutical and Biomedical Analysis, 2012, 70, 344-353. | 2.8 | 24 |
| 11 | Discovery of Novel Orally Bioavailable Oxaborole 6-Carboxamides That Demonstrate Cure in a Murine Model of Late-Stage Central Nervous System African Trypanosomiasis. Antimicrobial Agents and Chemotherapy, 2010, 54, 4379-4388. | 3.2 | 95 |
| 12 | The liver pharmacological and xenobiotic gene response repertoire. Molecular Systems Biology, 2008, 4, 175. | 7.2 | 71 |
| 13 | NSAID-Induced Acute Phase Response is Due to Increased Intestinal Permeability and Characterized by Early and Consistent Alterations in Hepatic Gene Expression. Toxicologic Pathology, 2006, 34, 168-179. | 1.8 | 51 |
| 14 | Classification of a large microarray data set: Algorithm comparison and analysis of drug signatures. Genome Research, 2005, 15, 724-736. | 5.5 | 104 |
| 15 | A Gene Expression Signature that Predicts the Future Onset of Drug-Induced Renal Tubular Toxicity. Toxicologic Pathology, 2005, 33, 675-683. | 1.8 | 112 |
| 16 | Use of a mixed tissue RNA design for performance assessments on multiple microarray formats. Nucleic Acids Research, 2005, 33, e187-e187. | 14.5 | 30 |
| 17 | Development of a large-scale chemogenomics database to improve drug candidate selection and to understand mechanisms of chemical toxicity and action. Journal of Biotechnology, 2005, 119, 219-244. | 3.8 | 282 |
| 18 | Differential Distribution of Bradykinin B ₂ Receptors in the Rat and Human Cardiovascular System. Hypertension, 2001, 37, 110-120. | 2.7 | 49 |

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|----|--|-----|-----------|
| 19 | Structure-based design of six novel classes of nonpeptide antagonists of the bradykinin B2 receptor. Bioorganic and Medicinal Chemistry Letters, 2000, 10, 2421-2425. | 2.2 | 11 |
| 20 | Identification of the Binding Site for a Novel Class of CCR2b Chemokine Receptor Antagonists. Journal of Biological Chemistry, 2000, 275, 25562-25571. | 3.4 | 232 |
| 21 | Identification of Surface Residues of the Monocyte Chemotactic Protein 1 That Affect Signaling through the Receptor CCR2â€. Biochemistry, 1999, 38, 16167-16177. | 2.5 | 103 |
| 22 | Identification of Residues in the Monocyte Chemotactic Protein-1 That Contact the MCP-1 Receptor, CCR2â€. Biochemistry, 1999, 38, 13013-13025. | 2.5 | 141 |
| 23 | Purification and physical characterization of cloned human cAMP phosphodiesterases PDE-4D and-4C. Cell Biochemistry and Biophysics, 1998, 28, 187-217. | 1.8 | 6 |
| 24 | Comparison of Recombinant Human PDE4 Isoforms. Cellular Signalling, 1998, 10, 427-440. | 3.6 | 52 |
| 25 | Monomeric Monocyte Chemoattractant Protein-1 (MCP-1) Binds and Activates the MCP-1 Receptor CCR2B. Journal of Biological Chemistry, 1998, 273, 33157-33165. | 3.4 | 183 |
| 26 | Multiple splice variants of phosphodiesterase PDE4C cloned from human lung and testis. Biochimica Et Biophysica Acta Gene Regulatory Mechanisms, 1997, 1353, 287-297. | 2.4 | 38 |
| 27 | Nonpeptide bradykinin antagonist analogs based on a model of a Sterling-Winthrop nonpeptide bradykinin antagonist overlapped with cyclic hexapeptide bradykinin antagonist peptides. Bioorganic and Medicinal Chemistry Letters, 1997, 7, 1921-1926. | 2.2 | 6 |
| 28 | Na+Ions Binding to the Bradykinin B2Receptor Suppress Agonist-Independent Receptor Activationâ€. Biochemistry, 1996, 35, 13368-13377. | 2,5 | 42 |
| 29 | Mutations in the B2 Bradykinin Receptor Reveal a Different Pattern of Contacts for Peptidic Agonists and Peptidic Antagonists. Journal of Biological Chemistry, 1996, 271, 28277-28286. | 3.4 | 62 |
| 30 | The N-terminal Amino Group of [Tyr8]Bradykinin Is Bound Adjacent to Analogous Amino Acids of the Human and Rat B2 Receptor. Journal of Biological Chemistry, 1996, 271, 27382-27387. | 3.4 | 27 |
| 31 | Extracellular Domains of the Bradykinin B2 Receptor Involved in Ligand Binding and Agonist Sensing Defined by Anti-peptide Antibodies. Journal of Biological Chemistry, 1996, 271, 1748-1755. | 3.4 | 83 |
| 32 | Cyclic hexapeptide antagonists of the bradykinin B2 receptor: Receptor binding and solution backbone conformation. International Journal of Peptide Research and Therapeutics, 1995, 1, 229-234. | 0.1 | 5 |
| 33 | Cloning and functional expression of the cDNA encoding rat lanosterol 14-α demethylase. Gene, 1995, 161, 243-248. | 2.2 | 31 |
| 34 | The cDNA of a human lymphocyte cyclic-AMP phosphodiesterase (PDE IV) reveals a multigene family. Gene, 1993, 129, 239-247. | 2.2 | 78 |
| 35 | Fed-batch culture of insect cells: a method to increase the yield of recombinant human nerve growth factor (rhNGF) in the baculovirus expression system. Journal of Biotechnology, 1993, 31, 205-217. | 3.8 | 72 |
| | Claning of a P2 Bradyhinin Decentory Eveningtion of the Bradyhinin Binding Site by Site Directed | | |

Cloning of a B2 Bradykinin Receptor: Examination of the Bradykinin Binding Site by Site Directed Mutagenesis., 1992, 38 (Pt 1), 487-496.

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|----|--|------|-----------|
| 37 | Physicochemical Characterization of Recombinant Human Nerve Growth Factor Produced in Insect Cells with a Baculovirus Vector. Journal of Neurochemistry, 1991, 57, 1052-1061. | 3.9 | 19 |
| 38 | The human insulin receptor cDNA: The structural basis for hormone-activated transmembrane signalling. Cell, 1985, 40, 747-758. | 28.9 | 1,563 |