Philippe Diaz

List of Publications by Year in descending order

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Version: 2024-02-01

361296 330025 1,392 40 20 citations h-index papers

g-index 48 48 48 1948 docs citations times ranked citing authors all docs

37

| # | Article | IF | CITATIONS |
|----|---|-----|-----------|
| 1 | Neuroinflammation, Hyperphosphorylated Tau, Diffuse Amyloid Plaques, and Down-Regulation of the Cellular Prion Protein in Air Pollution Exposed Children and Young Adults. Journal of Alzheimer's Disease, 2012, 28, 93-107. | 1.2 | 234 |
| 2 | Solution-Phase Synthesis of Diaryl Selenides Using Polymer-Supported Borohydride. Organic Letters, 2000, 2, 1705-1708. | 2.4 | 148 |
| 3 | 6-Methoxy- <i>N</i> -alkyl Isatin Acylhydrazone Derivatives as a Novel Series of Potent Selective Cannabinoid Receptor 2 Inverse Agonists: Design, Synthesis, and Binding Mode Prediction. Journal of Medicinal Chemistry, 2009, 52, 433-444. | 2.9 | 74 |
| 4 | 2,3â€Dihydroâ€1â€Benzofuran Derivatives as a Series of Potent Selective Cannabinoid Receptorâ€2 Agonists: Design, Synthesis, and Binding Mode Prediction through Ligandâ€Steered Modeling. ChemMedChem, 2009, 4, 1615-1629. | 1.6 | 71 |
| 5 | New synthetic retinoids obtained by palladium-catalyzed tandem cyclisation-hydride capture process. Tetrahedron, 1998, 54, 4579-4590. | 1.0 | 70 |
| 6 | MDA7: a novel selective agonist for CB ₂ receptors that prevents allodynia in rat neuropathic pain models. British Journal of Pharmacology, 2008, 155, 1104-1116. | 2.7 | 64 |
| 7 | Prevention of Paclitaxel-Induced Neuropathy Through Activation of the Central Cannabinoid Type 2 Receptor System. Anesthesia and Analgesia, 2012, 114, 1104-1120. | 1.1 | 63 |
| 8 | Design and Synthesis of a Novel Series of $\langle i \rangle N \langle i \rangle$ -Alkyl Isatin Acylhydrazone Derivatives that Act as Selective Cannabinoid Receptor 2 Agonists for the Treatment of Neuropathic Pain. Journal of Medicinal Chemistry, 2008, 51, 4932-4947. | 2.9 | 59 |
| 9 | Efficient synthetic approach to heterocycles possessing the 3,3-disubstituted-2,3-dihydrobenzofuran skeleton via diverse palladium-catalyzed tandem reactions. Tetrahedron, 2007, 63, 3340-3349. | 1.0 | 53 |
| 10 | Solid-Phase Synthesis of Diaryl Sulfides: Direct Coupling of Solid-Supported Aryl Halides with Thiols Using an Insoluble Polymer-Supported Reagent. Organic Letters, 2005, 7, 2719-2722. | 2.4 | 43 |
| 11 | Characterization of Novel Cannabinoid Based T-Type Calcium Channel Blockers with Analgesic Effects. ACS Chemical Neuroscience, 2015, 6, 277-287. | 1.7 | 42 |
| 12 | Pharmacological Characterization of a Novel Cannabinoid Ligand, MDA19, for Treatment of Neuropathic Pain. Anesthesia and Analgesia, 2010, 111, 99-109. | 1.1 | 41 |
| 13 | Palladium-catalyzed cascade allylation/carbopalladation/cross coupling: a novel three-component reaction for the synthesis of 3,3-disubstituted-2,3-dihydrobenzofurans. Tetrahedron Letters, 2003, 44, 8657-8659. | 0.7 | 39 |
| 14 | Mastering tricyclic ring systems for desirable functional cannabinoid activity. European Journal of Medicinal Chemistry, 2013, 69, 881-907. | 2.6 | 39 |
| 15 | Analgesic Effect of a Mixed T-Type Channel Inhibitor/CB ₂ Receptor Agonist. Molecular Pain, 2013, 9, 1744-8069-9-32. | 1.0 | 36 |
| 16 | NMP-7 Inhibits Chronic Inflammatory and Neuropathic Pain via Block of Cav3.2 T-type Calcium Channels and Activation of CB2 Receptors. Molecular Pain, 2014, 10, 1744-8069-10-77. | 1.0 | 32 |
| 17 | Functional Characterization and Analgesic Effects of Mixed Cannabinoid Receptor/T-Type Channel Ligands. Molecular Pain, 2011, 7, 1744-8069-7-89. | 1.0 | 31 |
| 18 | Design and evaluation of a novel fluorescent CB2 ligand as probe for receptor visualization in immune cells. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 5859-5862. | 1.0 | 25 |

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|----|--|-----|-----------|
| 19 | Spinal gene expression profiling and pathways analysis of a CB2 agonist (MDA7)-targeted prevention of paclitaxel-induced neuropathy. Neuroscience, 2014, 260, 185-194. | 1.1 | 25 |
| 20 | Synthesis of New Quinolinequinone Derivatives and Preliminary Exploration of their Cytotoxic Properties. Journal of Medicinal Chemistry, 2013, 56, 3806-3819. | 2.9 | 22 |
| 21 | Definition of functionally and structurally distinct repressive states in the nuclear receptor PPAR \hat{I}^3 . Nature Communications, 2019, 10, 5825. | 5.8 | 20 |
| 22 | Modified carbazoles destabilize microtubules and kill glioblastoma multiform cells. European Journal of Medicinal Chemistry, 2018, 159, 74-89. | 2.6 | 19 |
| 23 | New selenium-containing acetylenic retinoids by direct coupling of alkynylsilanes with selenylhalides. Tetrahedron Letters, 1998, 39, 9003-9006. | 0.7 | 16 |
| 24 | Suzuki–Miyaura cross-coupling of benzylic bromides under microwave conditions. Tetrahedron Letters, 2011, 52, 5656-5658. | 0.7 | 15 |
| 25 | Up-Regulation of mRNA Ventricular PRNP Prion Protein Gene Expression in Air Pollution Highly Exposed Young Urbanites: Endoplasmic Reticulum Stress, Glucose Regulated Protein 78, and Nanosized Particles. International Journal of Molecular Sciences, 2013, 14, 23471-23491. | 1.8 | 14 |
| 26 | Cav3.2 T-type calcium channels control acute itch in mice. Molecular Brain, 2020, 13, 119. | 1.3 | 13 |
| 27 | In Vivo Efficacy of Enabling Formulations Based on Hydroxypropyl-Î ² -Cyclodextrins, Micellar Preparation, and Liposomes for the Lipophilic Cannabinoid CB2 Agonist, MDA7. Journal of Pharmaceutical Sciences, 2013, 102, 352-364. | 1.6 | 12 |
| 28 | Identification of Tazarotenic Acid as the First Xenobiotic Substrate of Human Retinoic Acid Hydroxylase CYP26A1 and CYP26B1. Journal of Pharmacology and Experimental Therapeutics, 2016, 357, 281-292. | 1.3 | 11 |
| 29 | Development and Characterization of Novel and Selective Inhibitors of Cytochrome P450 CYP26A1, the Human Liver Retinoic Acid Hydroxylase. Journal of Medicinal Chemistry, 2016, 59, 2579-2595. | 2.9 | 11 |
| 30 | A brain-penetrant microtubule-targeting agent that disrupts hallmarks of glioma tumorigenesis. Neuro-Oncology Advances, 2021, 3, vdaa165. | 0.4 | 10 |
| 31 | Comparison of the ligand binding site of CYP2C8 with CYP26A1 and CYP26B1: a structural basis for the identification of new inhibitors of the retinoic acid hydroxylases. Journal of Enzyme Inhibition and Medicinal Chemistry, 2016, 31, 148-161. | 2.5 | 9 |
| 32 | Characterization of CYP26B1-Selective Inhibitor, DX314, as a Potential Therapeutic for Keratinization Disorders. Journal of Investigative Dermatology, 2021, 141, 72-83.e6. | 0.3 | 9 |
| 33 | Synthesis and biological activities of new heterocyclic aromatic retinoids. Bioorganic and Medicinal Chemistry Letters, 1997, 7, 2289-2294. | 1.0 | 8 |
| 34 | Coupling reaction of chalcogenyl halides with alkynes on a solid support. Synthesis of new selenium-containing retinoids. Tetrahedron Letters, 2000, 41, 5193-5197. | 0.7 | 6 |
| 35 | Synthesis of Bridged Bicyclic β-Trifluoroacetoxy β-Trifluoromethyl α-Amino Acid Derivatives by an Original Dakin-West/Diels-Alder Tandem Sequence. Synlett, 1995, 1995, 101-102. | 1.0 | 5 |
| 36 | Preclinical assessment of dual CYP26[A1/B1] inhibitor, DX308, as an improved treatment for keratinization disorders. Skin Health and Disease, 2021, 1, e22. | 0.7 | 2 |

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|----|--|-----|-----------|
| 37 | Chemoenzymatic synthesis of enantiomers of a new retinoid to investigate the role of chirality in the biological response. Bioorganic and Medicinal Chemistry Letters, 1995, 5, 2801-2804. | 1.0 | 1 |
| 38 | Palladium-Catalyzed Cascade Allylation/Carbopalladation/Cross Coupling: A Novel Three-Component Reaction for the Synthesis of 3,3-Disubstituted-2,3-dihydrobenzofurans ChemInform, 2004, 35, no. | 0.1 | 0 |
| 39 | 529 Novel CYP26 inhibitors potentiate the effects of all- trans -retinoic acid on phenotype of normal and Darier disease keratinocytes in reconstructed human epidermis. Journal of Investigative Dermatology, 2017, 137, S91. | 0.3 | O |
| 40 | DDIS-29. BRAIN-PENETRANT MICROTUBULE-TARGETING AGENT, ST-401, KILLS GLIOBLASTOMA THROUGH A NOVEL MECHANISM. Neuro-Oncology, 2019, 21, vi69-vi69. | 0.6 | 0 |