

# Mariel Marder

## List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

57  
papers

2,336  
citations

25  
h-index

48  
g-index

61  
ext. papers

2,550  
ext. citations

4.7  
avg, IF

4.4  
L-index

| #  | Paper  | IF   | Citations |
|----|--|------|-----------|
| 57 | Multitarget 2'-hydroxychalcones as potential drugs for the treatment of neurodegenerative disorders and their comorbidities. <i>Neuropharmacology</i> , <b>2021</b> , 201, 108837  | 5.5  | 1         |
| 56 | 2-Hydroxy-4,5,6-trimethyl-4-dimethylaminochalcone, a novel fluorescent flavonoid with capacity to detect aluminium in cells and modulate Alzheimer's disease targets. <i>Journal of Photochemistry and Photobiology A: Chemistry</i> , <b>2021</b> , 409, 113137 | 4.7  | 4         |
| 55 | Longitudinal evaluation of a novel BChE PET tracer as an early biomarker in the brain of a mouse model for Alzheimer disease. <i>Theranostics</i> , <b>2021</b> , 11, 6542-6559  | 12.1 | 4         |
| 54 | Neurobehavioral evaluation and phytochemical characterization of a series of argentine valerian species. <i>Heliyon</i> , <b>2020</b> , 6, e05691  | 3.6  | 1         |
| 53 | Stereoselective Activity of 1-Propargyl-4-styrylpiperidine-like Analogues That Can Discriminate between Monoamine Oxidase Isoforms A and B. <i>Journal of Medicinal Chemistry</i> , <b>2020</b> , 63, 1361-1387  | 8.3  | 20        |
| 52 | Natural flavonoids inhibit the plasma membrane Ca-ATPase. <i>Biochemical Pharmacology</i> , <b>2019</b> , 166, 1-11  | 6    | 9         |
| 51 | Chalcone derivatives: synthesis, and evaluation of their anti-anxiety, anti-depression and analgesic effects. <i>Heliyon</i> , <b>2019</b> , 5, e01376   | 3.6  | 15        |
| 50 | N-propyl-2,2-diphenyl-2-hydroxyacetamide, a novel $\alpha$ -hydroxyamide with anticonvulsant, anxiolytic and antidepressant-like effects that inhibits voltage-gated sodium channels. <i>European Journal of Pharmacology</i> , <b>2018</b> , 819, 270-280       | 5.3  | 5         |
| 49 | Novel sulfamides and sulfamates derived from amino esters: Synthetic studies and anticonvulsant activity. <i>European Journal of Pharmacology</i> , <b>2016</b> , 774, 55-63   | 5.3  | 16        |
| 48 | The Role of Galectin-3: From Oligodendroglial Differentiation and Myelination to Demyelination and Remyelination Processes in a Cuprizone-Induced Demyelination Model. <i>Advances in Experimental Medicine and Biology</i> , <b>2016</b> , 949, 311-332         | 3.6  | 11        |
| 47 | Galectin-3 controls the response of microglial cells to limit cuprizone-induced demyelination. <i>Neurobiology of Disease</i> , <b>2014</b> , 62, 441-55   | 7.5  | 49        |
| 46 | Anticonvulsant profile of 2-ethylthio-7-methyl-4-(4-methylphenyl)pyrazolo[1,5-a][1,3,5]triazine. <i>Brazilian Journal of Pharmaceutical Sciences</i> , <b>2014</b> , 50, 73-81   | 1.8  | 3         |
| 45 | A synthetic bioisoster of trimethadione and phenytoin elicits anticonvulsant effect, protects the brain oxidative damage produced by seizures and exerts antidepressant action in mice. <i>European Neuropsychopharmacology</i> , <b>2014</b> , 24, 1405-14      | 1.2  | 7         |
| 44 | Synthesis and anticonvulsant activity of bioisosteres of trimethadione, N-derivative-1,2,3-oxathiazolidine-4-one-2,2-dioxides from $\alpha$ -hydroxyamides. <i>Bioorganic and Medicinal Chemistry</i> , <b>2013</b> , 21, 841-6                                  | 3.4  | 12        |
| 43 | In vitro binding affinities of a series of flavonoids for $\mu$ -opioid receptors. Antinociceptive effect of the synthetic flavonoid 3,3-dibromoflavanone in mice. <i>Neuropharmacology</i> , <b>2013</b> , 72, 9-19   | 5.5  | 29        |
| 42 | N,N'-dicyclohexylsulfamide and N,N'-diphenethylsulfamide are anticonvulsant sulfamides with affinity for the benzodiazepine binding site of the GABA(A) receptor and anxiolytic activity in mice. <i>Biochemical Pharmacology</i> , <b>2012</b> , 83, 253-9      | 6    | 14        |
| 41 | Flavonoids as GABAA receptor ligands: the whole story?. <i>Journal of Experimental Pharmacology</i> , <b>2012</b> , 4, 9-24  | 3    | 45        |

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|----|--|------|-----|
| 40 | Chronic intraperitoneal and oral treatments with hesperidin induce central nervous system effects in mice. <i>Phytotherapy Research</i> , <b>2012</b> , 26, 308-12   | 6.7  | 12  |
| 39 | Adult CNP::EGFP transgenic mouse shows pronounced hypomyelination and an increased vulnerability to cuprizone-induced demyelination. <i>Experimental Neurology</i> , <b>2012</b> , 233, 490-504  | 5.7  | 6   |
| 38 | 2'-Nitroflavone induces apoptosis and modulates mitogen-activated protein kinase pathways in human leukaemia cells. <i>Anti-Cancer Drugs</i> , <b>2012</b> , 23, 815-26  | 2.4  | 7   |
| 37 | Galectin-3 drives oligodendrocyte differentiation to control myelin integrity and function. <i>Cell Death and Differentiation</i> , <b>2011</b> , 18, 1746-56  | 12.7 | 102 |
| 36 | Hesperidin induces antinociceptive effect in mice and its aglycone, hesperetin, binds to $\mu$ opioid receptor and inhibits GIRK1/2 currents. <i>Pharmacology Biochemistry and Behavior</i> , <b>2011</b> , 99, 333-41                               | 3.9  | 20  |
| 35 | Central nervous system activities of two diterpenes isolated from <i>Aloysia virgata</i> . <i>Phytomedicine</i> , <b>2011</b> , 18, 393-401  | 6.5  | 16  |
| 34 | Isovaleramida, principio anticonvulsivo aislado de <i>Valeriana pavonii</i> . <i>Biomedica</i> , <b>2010</b> , 30, 245   | 0.9  | 8   |
| 33 | Hesperidin, a flavonoid glycoside with sedative effect, decreases brain pERK1/2 levels in mice. <i>Pharmacology Biochemistry and Behavior</i> , <b>2009</b> , 92, 291-6  | 3.9  | 21  |
| 32 | In vitro induction of apoptosis and in vivo effects of a flavone nitroderivative in murine mammary adenocarcinoma cells. <i>International Journal of Cancer</i> , <b>2009</b> , 125, 222-8   | 7.5  | 3   |
| 31 | Neuroactive flavonoid glycosides from <i>Tilia petiolaris</i> DC. extracts. <i>Phytotherapy Research</i> , <b>2009</b> , 23, 1453-7  | 6.7  | 16  |
| 30 | Opioid receptors are involved in the sedative and antinociceptive effects of hesperidin as well as in its potentiation with benzodiazepines. <i>European Journal of Pharmacology</i> , <b>2008</b> , 580, 306-13                                     | 5.3  | 48  |
| 29 | 6,3'-Dinitroflavone is a low efficacy modulator of GABA(A) receptors. <i>European Journal of Pharmacology</i> , <b>2008</b> , 591, 142-6   | 5.3  | 2   |
| 28 | 2'-Nitroflavone induces cell cycle arrest and apoptosis in HeLa human cervical carcinoma cells. <i>Cancer Letters</i> , <b>2008</b> , 268, 146-57  | 9.9  | 12  |
| 27 | Anticonvulsant and anxiolytic-like effects of compounds isolated from <i>Polygala sabulosa</i> (Polygalaceae) in rodents: in vitro and in vivo interactions with benzodiazepine binding sites. <i>Psychopharmacology</i> , <b>2008</b> , 197, 351-60 | 4.7  | 26  |
| 26 | Central nervous system depressant action of flavonoid glycosides. <i>European Journal of Pharmacology</i> , <b>2006</b> , 539, 168-76  | 5.3  | 173 |
| 25 | The anxiolytic-like effects of <i>Aloysia polystachya</i> (Griseb.) Moldenke (Verbenaceae) in mice. <i>Journal of Ethnopharmacology</i> , <b>2006</b> , 105, 400-8   | 5    | 45  |
| 24 | Antitumor activity of some natural flavonoids and synthetic derivatives on various human and murine cancer cell lines. <i>Bioorganic and Medicinal Chemistry</i> , <b>2006</b> , 14, 2966-71   | 3.4  | 190 |
| 23 | Synergistic interaction between hesperidin, a natural flavonoid, and diazepam. <i>European Journal of Pharmacology</i> , <b>2005</b> , 512, 189-98   | 5.3  | 53  |

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| 22 | Sedative and sleep-enhancing properties of linarin, a flavonoid-isolated from <i>Valeriana officinalis</i> . <i>Pharmacology Biochemistry and Behavior</i> , <b>2004</b> , 77, 399-404   | 3.9 | 160 |
| 21 | Antiproliferative activity of various flavonoids and related compounds: additive effect of interferon-alpha2b. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2004</b> , 14, 133-6   | 2.9 | 27  |
| 20 | 6-methylapigenin and hesperidin: new valeriana flavonoids with activity on the CNS. <i>Pharmacology Biochemistry and Behavior</i> , <b>2003</b> , 75, 537-45   | 3.9 | 146 |
| 19 | Isolation and identification of 6-methylapigenin, a competitive ligand for the brain GABA(A) receptors, from <i>Valeriana wallichii</i> . <i>Planta Medica</i> , <b>2002</b> , 68, 934-6   | 3.1 | 72  |
| 18 | GABA(A)-receptor ligands of flavonoid structure. <i>Current Topics in Medicinal Chemistry</i> , <b>2002</b> , 2, 853-67  | 3   | 135 |
| 17 | Molecular modeling and QSAR analysis of the interaction of flavone derivatives with the benzodiazepine binding site of the GABA(A) receptor complex. <i>Bioorganic and Medicinal Chemistry</i> , <b>2001</b> , 9, 323-35   | 3.4 | 77  |
| 16 | 6-Chloro-3'-nitroflavone is a potent ligand for the benzodiazepine binding site of the GABA(A) receptor devoid of intrinsic activity. <i>Pharmacology Biochemistry and Behavior</i> , <b>2000</b> , 65, 313-20   | 3.9 | 17  |
| 15 | 6,3'-dibromoflavone and 6-nitro-3'-bromoflavone: new additions to the 6,3'-disubstituted flavone family of high-affinity ligands of the brain benzodiazepine binding site with agonistic properties. <i>Biochemical and Biophysical Research Communications</i> , <b>2000</b> , 273, 694-8 | 3.4 | 14  |
| 14 | Flavonoids and the central nervous system: from forgotten factors to potent anxiolytic compounds. <i>Journal of Pharmacy and Pharmacology</i> , <b>1999</b> , 51, 519-26   | 4.8 | 111 |
| 13 | 6-Methyl-3'-bromoflavone, a high-affinity ligand for the benzodiazepine binding site of the GABA(A) receptor with some antagonistic properties. <i>Biochemical and Biophysical Research Communications</i> , <b>1999</b> , 262, 643-6  | 3.4 | 8   |
| 12 | Pharmacological characterization of 6-bromo-3'-nitroflavone, a synthetic flavonoid with high affinity for the benzodiazepine receptors. <i>Pharmacology Biochemistry and Behavior</i> , <b>1998</b> , 61, 239-46   | 3.9 | 17  |
| 11 | Neuroactive flavonoids: new ligands for the Benzodiazepine receptors. <i>Phytomedicine</i> , <b>1998</b> , 5, 235-43   | 6.5 | 63  |
| 10 | Detection of benzodiazepine receptor ligands in small libraries of flavone derivatives synthesized by solution phase combinatorial chemistry. <i>Biochemical and Biophysical Research Communications</i> , <b>1998</b> , 249, 481-5  | 3.4 | 60  |
| 9  | Sedative and hypnotic properties of <i>Salvia guaranitica</i> St. Hil. and of its active principle, Cirsilinol. <i>Phytomedicine</i> , <b>1997</b> , 4, 47-52  | 6.5 | 24  |
| 8  | Overview--flavonoids: a new family of benzodiazepine receptor ligands. <i>Neurochemical Research</i> , <b>1997</b> , 22, 419-25  | 4.6 | 160 |
| 7  | 6-Bromo-3'-nitroflavone, a new high affinity benzodiazepine receptor agonist recognizes two populations of cerebral cortical binding sites. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>1997</b> , 7, 373-378   | 2.9 | 17  |
| 6  | Synthesis of halogenated/nitrated flavone derivatives and evaluation of their affinity for the central benzodiazepine receptor. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>1997</b> , 7, 2003-2008   | 2.9 | 30  |
| 5  | 6-Bromoflavone, a high affinity ligand for the central benzodiazepine receptors is a member of a family of active flavonoids. <i>Biochemical and Biophysical Research Communications</i> , <b>1996</b> , 223, 384-9  | 3.4 | 53  |

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|---|--|-----|----|
| 4 | Anxiolytic properties of 6,3'-dinitroflavone, a high-affinity benzodiazepine receptor ligand. <i>European Journal of Pharmacology</i> , <b>1996</b> , 318, 23-30   | 5.3 | 63 |
| 3 | Cirsiliol and caffeic acid ethyl ester, isolated from <i>Salvia guaranitica</i> , are competitive ligands for the central benzodiazepine receptors. <i>Phytomedicine</i> , <b>1996</b> , 3, 29-31  | 6.5 | 33 |
| 2 | 6,3'-Dinitroflavone, a novel high affinity ligand for the benzodiazepine receptor with potent anxiolytic properties. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>1995</b> , 5, 2717-2720  | 2.9 | 30 |
| 1 | Reaction of 2,6-dinitroanisole with cyclohexylamine in toluene/ethanol binary solvents. Further support for the dimer nucleophile mechanism in aromatic nucleophilic substitution. <i>Journal of the Chemical Society Perkin Transactions II</i> , <b>1993</b> , 229-233 |     | 12 |