

# Onintza Sagredo

## List of Publications by Year in descending order

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Version: 2024-02-01

30  
papers

2,351  
citations

331259

21  
h-index

476904

29  
g-index

30  
all docs

30  
docs citations

30  
times ranked

2628  
citing authors

#	ARTICLE	IF	CITATIONS
1	Preclinical investigation of Î²-caryophyllene as a therapeutic agent in an experimental murine model of Dravet syndrome. <i>Neuropharmacology</i> , 2022, 205, 108914.	2.0	5
2	Possible therapeutic applications of cannabis in the neuropsychopharmacology field. <i>European Neuropsychopharmacology</i> , 2020, 36, 217-234.	0.3	24
3	Neuropathological Characterization of a Dravet Syndrome Knock-In Mouse Model Useful for Investigating Cannabinoid Treatments. <i>Frontiers in Molecular Neuroscience</i> , 2020, 13, 602801.	1.4	13
4	Chapter 2. Phytocannabinoids Versus Endocannabinoids. A Modern View of the Endocannabinoid System. <i>RSC Drug Discovery Series</i> , 2020, , 10-47.	0.2	0
5	Cannabinoid signalling in the immature brain: Encephalopathies and neurodevelopmental disorders. <i>Biochemical Pharmacology</i> , 2018, 157, 85-96.	2.0	16
6	Effects of a Sativex-Like Combination of Phytocannabinoids on Disease Progression in R6/2 Mice, an Experimental Model of Huntingtonâ€™s Disease. <i>International Journal of Molecular Sciences</i> , 2017, 18, 684.	1.8	20
7	A double-blind, randomized, cross-over, placebo-controlled, pilot trial with Sativex in Huntingtonâ€™s disease. <i>Journal of Neurology</i> , 2016, 263, 1390-1400.	1.8	105
8	Analysis of endocannabinoid signaling elements and related proteins in lymphocytes of patients with Dravet syndrome. <i>Pharmacology Research and Perspectives</i> , 2016, 4, e00220.	1.1	13
9	Neuroprotective Properties of Cannabigerol in Huntington's Disease: Studies in R6/2 Mice and 3-Nitropropionate-lesioned Mice. <i>Neurotherapeutics</i> , 2015, 12, 185-199.	2.1	92
10	A restricted population of CB <sub>1</sub> cannabinoid receptors with neuroprotective activity. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014, 111, 8257-8262.	3.3	136
11	Cannabidiol for neurodegenerative disorders: important new clinical applications for this phytocannabinoid?. <i>British Journal of Clinical Pharmacology</i> , 2013, 75, 323-333.	1.1	254
12	The inhibition of 2-arachidonoyl-glycerol (2-AG) biosynthesis, rather than enhancing striatal damage, protects striatal neurons from malonate-induced death: a potential role of cyclooxygenase-2-dependent metabolism of 2-AG. <i>Cell Death and Disease</i> , 2013, 4, e862-e862.	2.7	69
13	Natural Cannabinoids Improve Dopamine Neurotransmission and Tau and Amyloid Pathology in a Mouse Model of Tauopathy. <i>Journal of Alzheimer's Disease</i> , 2013, 35, 525-539.	1.2	98
14	Cannabinoids: Novel Medicines for the Treatment of Huntingtons Disease. <i>Recent Patents on CNS Drug Discovery</i> , 2012, 7, 41-48.	0.9	64
15	Q23â€¦A double blind, cross over, placebo-controlled, phase II trial of Sativex in Huntington's Disease. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2012, 83, A62.2-A62.	0.9	3
16	Sativex-like Combination of Phytocannabinoids is Neuroprotective in Malonate-Lesioned Rats, an Inflammatory Model of Huntingtonâ€™s Disease: Role of CB <sub>1</sub> and CB <sub>2</sub> Receptors. <i>ACS Chemical Neuroscience</i> , 2012, 3, 400-406.	1.7	81
17	New Serotonin 5-HT <sub>1A</sub> Receptor Agonists with Neuroprotective Effect against Ischemic Cell Damage. <i>Journal of Medicinal Chemistry</i> , 2011, 54, 7986-7999.	2.9	36
18	Neuroprotective effects of phytocannabinoid-based medicines in experimental models of Huntington's disease. <i>Journal of Neuroscience Research</i> , 2011, 89, 1509-1518.	1.3	84

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19	Loss of striatal type 1 cannabinoid receptors is a key pathogenic factor in Huntington's disease. <i>Brain</i> , 2011, 134, 119-136.	3.7	178
20	The endocannabinoid system as a target for the treatment of neuronal damage. <i>Expert Opinion on Therapeutic Targets</i> , 2010, 14, 387-404.	1.5	78
21	Cannabinoid CB <sub>2</sub> receptor agonists protect the striatum against malonate toxicity: Relevance for Huntington's disease. <i>Glia</i> , 2009, 57, 1154-1167.	2.5	165
22	Microglial CB2 cannabinoid receptors are neuroprotective in Huntington's disease excitotoxicity. <i>Brain</i> , 2009, 132, 3152-3164.	3.7	323
23	Lack of association between polymorphisms in cannabinoid receptor gene (CNR1) and fatty acid amide hydroxylase gene (FAAH) and eating disorders in a preliminary study. <i>Psychiatric Genetics</i> , 2009, 19, 336.	0.6	5
24	Role of CB2 receptors in neuroprotective effects of cannabinoids. <i>Molecular and Cellular Endocrinology</i> , 2008, 286, S91-S96.	1.6	105
25	The Endocannabinoid System in Huntingtons Disease. <i>Current Pharmaceutical Design</i> , 2008, 14, 2317-2325.	0.9	61
26	Cannabidiol reduced the striatal atrophy caused by 3-nitropropionic acid <i>in vivo</i> by mechanisms independent of the activation of cannabinoid, vanilloid TRPV <sub>1</sub> and adenosine A <sub>2A</sub> receptors. <i>European Journal of Neuroscience</i> , 2007, 26, 843-851.	1.2	120
27	Cannabinoids and Neuroprotection in Basal Ganglia Disorders. <i>Molecular Neurobiology</i> , 2007, 36, 82-91.	1.9	79
28	Chronic $\delta^9$ -tetrahydrocannabinol administration affects serotonin levels in the rat frontal cortex. <i>Naunyn-Schmiedeberg's Archives of Pharmacology</i> , 2006, 372, 313-317.	1.4	37
29	Therapeutic Potential of the Endocannabinoid System in the Brain. <i>Mini-Reviews in Medicinal Chemistry</i> , 2005, 5, 609-617.	1.1	13
30	Antinociceptive, behavioural and neuroendocrine effects of CP 55,940 in young rats. <i>Developmental Brain Research</i> , 2002, 136, 85-92.	2.1	74