

# Matti Lepistö

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/11475755/publications.pdf>

Version: 2024-02-01

12  
papers

398  
citations

933447

10  
h-index

1199594

12  
g-index

12  
all docs

12  
docs citations

12  
times ranked

630  
citing authors

#	ARTICLE	IF	CITATIONS
1	Ligand Binding Mechanism in Steroid Receptors: From Conserved Plasticity to Differential Evolutionary Constraints. <i>Structure</i> , 2015, 23, 2280-2290.	3.3	96
2	Discovery of a Novel Oral Glucocorticoid Receptor Modulator (AZD9567) with Improved Side Effect Profile. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 1785-1799.	6.4	54
3	Synthesis and Functionalization of Cyclic Sulfonimidamides: A Novel Chiral Heterocyclic Carboxylic Acid Bioisostere. <i>ACS Medicinal Chemistry Letters</i> , 2012, 3, 574-578.	2.8	46
4	Selective Nonsteroidal Glucocorticoid Receptor Modulators for the Inhaled Treatment of Pulmonary Diseases. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 8591-8605.	6.4	41
5	HTS followed by NMR based counterscreening. Discovery and optimization of pyrimidones as reversible and competitive inhibitors of xanthine oxidase. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014, 24, 1315-1321.	2.2	32
6	The discovery of potent and selective non-steroidal glucocorticoid receptor modulators, suitable for inhalation. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2014, 24, 2571-2577.	2.2	32
7	Potent and Orally Bioavailable Inverse Agonists of ROR $\beta$ Resulting from Structure-Based Design. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 7796-7813.	6.4	31
8	Saccharin Aza Bioisosteres—Synthesis and Preclinical Property Comparisons. <i>ACS Medicinal Chemistry Letters</i> , 2017, 8, 672-677.	2.8	23
9	Discovery of indazole ethers as novel, potent, non-steroidal glucocorticoid receptor modulators. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2016, 26, 5741-5748.	2.2	17
10	Discovery of Potent and Orally Bioavailable Inverse Agonists of the Retinoic Acid Receptor-Related Orphan Receptor C2. <i>ACS Medicinal Chemistry Letters</i> , 2019, 10, 972-977.	2.8	16
11	AZD0284, a Potent, Selective, and Orally Bioavailable Inverse Agonist of Retinoic Acid Receptor-Related Orphan Receptor C2. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 13807-13829.	6.4	7
12	Backbone $^1\text{H}$ , $^{13}\text{C}$ , and $^{15}\text{N}$ resonance assignments of the ligand binding domain of the human wildtype glucocorticoid receptor and the F602S mutant variant. <i>Biomolecular NMR Assignments</i> , 2018, 12, 263-268.	0.8	3