

# Matteo Santucci

## List of Publications by Year in descending order

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25  
papers

544  
citations

687363

13  
h-index

642732

23  
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all docs

25  
docs citations

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times ranked

979  
citing authors

#	ARTICLE	IF	CITATIONS
1	The Hippo Pathway and YAP/TAZ-TEAD Protein-Protein Interaction as Targets for Regenerative Medicine and Cancer Treatment. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 4857-4873.	6.4	141
2	Chroman-4-One Derivatives Targeting Pteridine Reductase 1 and Showing Anti-Parasitic Activity. <i>Molecules</i> , 2017, 22, 426.	3.8	39
3	Structure-Based Virtual Screening for the Discovery of Novel Inhibitors of New Delhi Metallo- $\beta$ -lactamase-1. <i>ACS Medicinal Chemistry Letters</i> , 2018, 9, 45-50.	2.8	38
4	Long period fiber grating working in reflection mode as valuable biosensing platform for the detection of drug resistant bacteria. <i>Sensors and Actuators B: Chemical</i> , 2016, 230, 510-520.	7.8	35
5	Computational and biological profile of boronic acids for the detection of bacterial serine- and metallo- $\beta$ -lactamases. <i>Scientific Reports</i> , 2017, 7, 17716.	3.3	35
6	Repurposing of Drugs Targeting YAP-TEAD Functions. <i>Cancers</i> , 2018, 10, 329.	3.7	33
7	Host dihydrofolate reductase (DHFR)-directed cycloguanil analogues endowed with activity against influenza virus and respiratory syncytial virus. <i>European Journal of Medicinal Chemistry</i> , 2017, 135, 467-478.	5.5	28
8	Label-free fiber optic optrode for the detection of class C $\beta$ -lactamases expressed by drug resistant bacteria. <i>Biomedical Optics Express</i> , 2017, 8, 5191.	2.9	25
9	Virtual screening identifies broad-spectrum $\beta$ -lactamase inhibitors with activity on clinically relevant serine- and metallo-carbapenemases. <i>Scientific Reports</i> , 2020, 10, 12763.	3.3	25
10	Optimization of Peptides That Target Human Thymidylate Synthase to Inhibit Ovarian Cancer Cell Growth. <i>Journal of Medicinal Chemistry</i> , 2014, 57, 1355-1367.	6.4	22
11	Virtual Screening and X-ray Crystallography Identify Non-Substrate Analog Inhibitors of Flavin-Dependent Thymidylate Synthase. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 9269-9275.	6.4	19
12	Synthesis, biological evaluation and molecular modeling of novel azaspiro dihydrotriazines as influenza virus inhibitors targeting the host factor dihydrofolate reductase (DHFR). <i>European Journal of Medicinal Chemistry</i> , 2018, 155, 229-243.	5.5	19
13	Accelerating Drug Discovery Efforts for Trypanosomatid Infections Using an Integrated Transnational Academic Drug Discovery Platform. <i>SLAS Discovery</i> , 2019, 24, 346-361.	2.7	18
14	Folic Acid-Peptide Conjugates Combine Selective Cancer Cell Internalization with Thymidylate Synthase Dimer Interface Targeting. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 3204-3221.	6.4	13
15	Intracellular quantitative detection of human thymidylate synthase engagement with an unconventional inhibitor using tetracysteine-diarsenical-probe technology. <i>Scientific Reports</i> , 2016, 6, 27198.	3.3	10
16	Alanine Mutants of the Interface Residues of Human Thymidylate Synthase Decode Key Features of the Binding Mode of Allosteric Anticancer Peptides. <i>Journal of Medicinal Chemistry</i> , 2015, 58, 1012-1018.	6.4	9
17	Identification of a 2,4-diaminopyrimidine scaffold targeting <i>Trypanosoma brucei</i> pteridine reductase 1 from the LIBRA compound library screening campaign. <i>European Journal of Medicinal Chemistry</i> , 2020, 189, 112047.	5.5	8
18	Multitarget, Selective Compound Design Yields Potent Inhibitors of a Kinetoplastid Pteridine Reductase 1. <i>Journal of Medicinal Chemistry</i> , 2022, 65, 9011-9033.	6.4	8

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19	Conformational Propensity and Biological Studies of Proline Mutated LR Peptides Inhibiting Human Thymidylate Synthase and Ovarian Cancer Cell Growth. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 7374-7380.	6.4	6
20	Cyclic Peptides Acting as Allosteric Inhibitors of Human Thymidylate Synthase and Cancer Cell Growth. <i>Molecules</i> , 2019, 24, 3493.	3.8	4
21	Evidence of Destabilization of the Human Thymidylate Synthase (hTS) Dimeric Structure Induced by the Interface Mutation Q62R. <i>Biomolecules</i> , 2019, 9, 134.	4.0	3
22	Structural Bases for the Synergistic Inhibition of Human Thymidylate Synthase and Ovarian Cancer Cell Growth by Drug Combinations. <i>Cancers</i> , 2021, 13, 2061.	3.7	2
23	Repurposing the Trypanosomatidic GSK Kinetobox for the Inhibition of Parasitic Pteridine and Dihydrofolate Reductases. <i>Pharmaceutics</i> , 2021, 14, 1246.	3.8	2
24	Intrinsic Fluorescence of the Active and the Inactive Functional Forms of Human Thymidylate Synthase. <i>ChemBioChem</i> , 2021, 22, 1800-1810.	2.6	1
25	Identification of a Quinone Derivative as a YAP/TEAD Activity Modulator from a Repurposing Library. <i>Pharmaceutics</i> , 2022, 14, 391.	4.5	1