Jie Tu

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/9936637/publications.pdf

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16	389	11	17
papers	citations	h-index	g-index
17	17	17	372 citing authors
all docs	docs citations	times ranked	

#	Article	IF	Citations
1	Heat shock protein 90 (Hsp90)/Histone deacetylase (HDAC) dual inhibitors for the treatment of azoles-resistant Candida albicans. European Journal of Medicinal Chemistry, 2022, 227, 113961.	2.6	22
2	Structure-Guided Discovery of the Novel Covalent Allosteric Site and Covalent Inhibitors of Fructose-1,6-Bisphosphate Aldolase to Overcome the Azole Resistance of Candidiasis. Journal of Medicinal Chemistry, 2022, 65, 2656-2674.	2.9	3
3	Discovery of Novel Sertraline Derivatives as Potent Anti- <i>Cryptococcus</i> Agents. Journal of Medicinal Chemistry, 2022, 65, 6541-6554.	2.9	8
4	Discovery of Piperidol Derivatives for Combinational Treatment of Azole-Resistant Candidiasis. ACS Infectious Diseases, 2021, 7, 650-660.	1.8	13
5	Effects of Hsp90 Inhibitor Ganetespib on Inhibition of Azole-Resistant Candida albicans. Frontiers in Microbiology, 2021, 12, 680382.	1.5	11
6	Lanosterol 14î±-demethylase (CYP51)/histone deacetylase (HDAC) dual inhibitors for treatment of Candida tropicalis and Cryptococcus neoformans infections. European Journal of Medicinal Chemistry, 2021, 221, 113524.	2.6	24
7	Novel Carboline Fungal Histone Deacetylase (HDAC) Inhibitors for Combinational Treatment of Azole-Resistant Candidiasis. Journal of Medicinal Chemistry, 2021, 64, 1116-1126.	2.9	35
8	Drug Repurposing of Haloperidol: Discovery of New Benzocyclane Derivatives as Potent Antifungal Agents against Cryptococcosis and Candidiasis. ACS Infectious Diseases, 2020, 6, 768-786.	1.8	35
9	Targeting fungal virulence factor by small molecules: Structure-based discovery of novel secreted aspartic protease 2 (SAP2) inhibitors. European Journal of Medicinal Chemistry, 2020, 201, 112515.	2.6	14
10	Discovery of Novel Fungal Lanosterol 14α-Demethylase (CYP51)/Histone Deacetylase Dual Inhibitors to Treat Azole-Resistant Candidiasis. Journal of Medicinal Chemistry, 2020, 63, 5341-5359.	2.9	45
11	Discovery of novel simplified isoxazole derivatives of sampangine as potent anti-cryptococcal agents. Bioorganic and Medicinal Chemistry, 2019, 27, 832-840.	1.4	12
12	Discovery of Simplified Sampangine Derivatives with Potent Antifungal Activities against Cryptococcal Meningitis. ACS Infectious Diseases, 2019, 5, 1376-1384.	1.8	21
13	Discovery of Carboline Derivatives as Potent Antifungal Agents for the Treatment of Cryptococcal Meningitis. Journal of Medicinal Chemistry, 2019, 62, 2376-2389.	2.9	24
14	Emerging New Targets for the Treatment of Resistant Fungal Infections. Journal of Medicinal Chemistry, 2018, 61, 5484-5511.	2.9	90
15	Discovery of simplified sampangine derivatives as novel fungal biofilm inhibitors. European Journal of Medicinal Chemistry, 2018, 143, 1510-1523.	2.6	18
16	Novel non-peptidic small molecule inhibitors of secreted aspartic protease 2 (SAP2) for the treatment of resistant fungal infections. Chemical Communications, 2018, 54, 13535-13538.	2.2	13