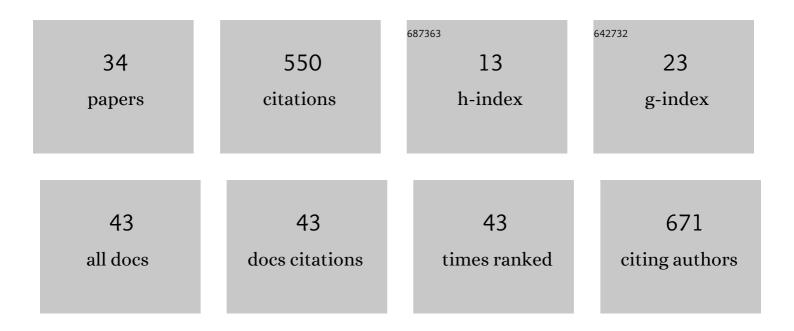
## Jae Hong Seo

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Total Synthesis of Cyclopiamide A Using Palladium-Catalyzed Domino Cyclization. Molecules, 2020, 25, 4903.	3.8	0
2	The phosphodiesterase 5 inhibitor, KJHâ€1002, reverses a mouse model of amnesia by activating a cGMP/cAMP response element binding protein pathway and decreasing oxidative damage. British Journal of Pharmacology, 2018, 175, 3347-3360.	5.4	21
3	Palladium-Catalyzed One-Pot Approach to 3-(1,3-Diarylprop-2-yn-1-ylidene)oxindoles. Heterocycles, 2018, 96, 1795.	0.7	3
4	Stereoselective Synthesis of 3-(1,3-Diarylallylidene)oxindoles via a Palladium-Catalyzed Tandem Reaction. Journal of Organic Chemistry, 2017, 82, 1864-1871.	3.2	19
5	Total Synthesis of $(\hat{A}_{\pm})$ -Decytospolides A and B. Synlett, 2017, 28, 249-252.	1.8	2
6	Consecutive One-Pot versus Domino Multicomponent Approaches to 3-(Diarylmethylene)oxindoles. Molecules, 2017, 22, 503.	3.8	12
7	Application of physiologically based pharmacokinetic modeling in predicting drug–drug interactions for sarpogrelate hydrochloride in humans. Drug Design, Development and Therapy, 2016, Volume 10, 2959-2972.	4.3	3
8	Aspirination of $\hat{l}$ ±-Aminoalcohol (Sarpogrelate M1). Molecules, 2016, 21, 1126.	3.8	0
9	Hendricksonâ€Reagentâ€Mediated Conversion of <i>N</i> â€Boc Carbamates to Isocyanates: Applications for the Synthesis of 3,4â€Dihydroisoquinolinâ€1â€ones and Ureas. Asian Journal of Organic Chemistry, 2016, 5, 287-292.	2.7	9
10	ldentification and characterization of potent, selective and metabolically stable IKKβ inhibitor. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 1120-1123.	2.2	1
11	Concise and Scalable Synthesis of Xanthylentin. Bulletin of the Korean Chemical Society, 2015, 36, 1957-1958.	1.9	3
12	Palladium-Catalyzed One-Pot Approach to 3-(Diarylmethylene)oxindoles from Propiolamidoaryl Triflate. Molecules, 2015, 20, 14022-14032.	3.8	6
13	Improved oral absorption of cilostazol via sulfonate salt formation with mesylate and besylate. Drug Design, Development and Therapy, 2015, 9, 3961.	4.3	12
14	Synthesis of 2-(Arylmethylene)-1,4-benzoxazin-3-one by One-Pot Sonogashira and 6-exo-Dig Cyclization. Heterocycles, 2015, 91, 1660.	0.7	1
15	Palladium-Catalyzed Tandem Approach to 3-(Diarylmethylene)oxindoles Using Microwave Irradiation. Synlett, 2015, 26, 2296-2300.	1.8	10
16	Discovery of a potent enoyl-acyl carrier protein reductase (Fabl) inhibitor suitable for antistaphylococcal agent. Bioorganic and Medicinal Chemistry Letters, 2015, 25, 4481-4486.	2.2	9
17	Successful reduction of off-target hERG toxicity by structural modification of a T-type calcium channel blocker. Bioorganic and Medicinal Chemistry Letters, 2014, 24, 880-883.	2.2	6
18	2-Phenylbenzofuran derivatives alleviate mitochondrial damage via the inhibition of β-amyloid aggregation. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 5882-5886.	2.2	5

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19	Stereoselective determination of ginsenosides <scp>R</scp> g3 and <scp>R</scp> h2 epimers in rat plasma by <scp>LC</scp> â€ <scp>MS</scp> / <scp>MS</scp> : Application to a pharmacokinetic study. Journal of Separation Science, 2013, 36, 1904-1912.	2.5	22
20	Synthesis of 3,4â€Ðihydroisoquinolinâ€1â€ones from <i>N</i> à€Bocâ€(βâ€Arylethyl)carbamates via Isocyanate Intermediates. European Journal of Organic Chemistry, 2013, 2013, 965-971.	2.4	21
21	Batten disease is linked to altered expression of mitochondria-related metabolic molecules. Neurochemistry International, 2013, 62, 931-935.	3.8	19
22	Synthesis of 3-(Diarylmethylene)oxindoles via a Palladium-Catalyzed One-Pot Reaction: Sonogashira-Heck-Suzuki-Miyaura Combined Reaction. Synlett, 2013, 24, 1993-1997.	1.8	14
23	Discovery of potent and selective rhodanine type IKKβ inhibitors by hit-to-lead strategy. Bioorganic and Medicinal Chemistry Letters, 2012, 22, 5668-5674.	2.2	19
24	Small molecules that protect against β-amyloid-induced cytotoxicity by inhibiting aggregation of β-amyloid. Bioorganic and Medicinal Chemistry, 2012, 20, 4921-4935.	3.0	20
25	Synthesis of the C′D′E′F′ Domain of Maitotoxin. Journal of the American Chemical Society, 2011, 133, 214-219.	13.7	30
26	A Practical and Cost-Effective Synthesis of d-erythro-Sphingosine from d-ribo-Phytosphingosine via a Cyclic Sulfate Intermediate. Synthesis, 2011, 2011, 867-872.	2.3	1
27	Total Synthesis of the Polycyclic Fungal Metabolite (±)â€Communesinâ€F. Angewandte Chemie - International Edition, 2010, 49, 2000-2003.	13.8	103
28	Synthesis of the QRSTU Domain of Maitotoxin and Its 85-‹i>epi‹/i>- and 86-‹i>epi‹/i>-Diastereoisomers. Journal of the American Chemical Society, 2010, 132, 9900-9907.	13.7	35
29	Evolution of a Strategy for Total Synthesis of the Marine Fungal Alkaloid (±)-Communesin F. Journal of Organic Chemistry, 2010, 75, 2667-2680.	3.2	47
30	Synthetic Studies on Perophoramidine and the Communesins:Â Construction of the Vicinal Quaternary Stereocenters. Journal of Organic Chemistry, 2006, 71, 8891-8900.	3.2	60
31	New Cephalosporin Antibiotics with 3-Triazolylpyridiniummethyl Substituents Journal of Antibiotics, 2001, 54, 460-462.	2.0	7
32	1,3-Dihydro-1,3-diacetyl-2H-benzimidazol-2-one: A New Versatile and Selective Acetylating Agent. Heterocycles, 2000, 53, 529.	0.7	11
33	Synthesis and structure–activity relationships of quaternary ammonium cephalosporins with 3-pyrazolylpyridinium derivatives. Bioorganic and Medicinal Chemistry Letters, 2000, 10, 1211-1214.	2.2	8
34	Synthesis andin vitro antibacterial activity of quaternary ammonium cephalosporin derivatives bearing oxazolidinone moiety. Archives of Pharmacal Research, 1999, 22, 579-584.	6.3	4