

# JÃ©rÃ©me Blanchet

## List of Publications by Year in descending order

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44  
papers

1,470  
citations

331670

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315739

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63  
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63  
docs citations

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times ranked

1679  
citing authors

#	ARTICLE	IF	CITATIONS
1	Phenylsilane and Silicon Tetraacetate: Versatile Promotors for Amide Synthesis. <i>European Journal of Organic Chemistry</i> , 2020, 2020, 388-392.	2.4	21
2	H-bonding vs Protonation of Alkynes in Regioselective Hydroamination Reactions: A Glimpse into the Reactivity of Arylogous Ynolethers and Ynamines. <i>Journal of Organic Chemistry</i> , 2019, 84, 15448-15475.	3.2	5
3	Catalytic and metal-free intramolecular hydroalkoxylation of alkynes. <i>Tetrahedron Letters</i> , 2019, 60, 534-537.	1.4	4
4	Borinic Acid Mediated Hydrosilylations: Reductions of Carbonyl Derivatives. <i>European Journal of Organic Chemistry</i> , 2019, 2019, 995-998.	2.4	11
5	Recent Advances in Amide Reductions. <i>Synthesis</i> , 2018, 50, 984-997.	2.3	66
6	Domino Ring Expansion: Regioselective Access to 9-Membered Lactones with a Fused Indole Unit from 2-Nitrophenyl-1,3-cyclohexanediones. <i>Chemistry - A European Journal</i> , 2018, 24, 2080-2084.	3.3	27
7	Borinic Acid Catalysed Reduction of Tertiary Amides with Hydrosilanes: A Mild and Chemoselective Synthesis of Amines. <i>Chemistry - A European Journal</i> , 2017, 23, 2005-2009.	3.3	46
8	Metal-Free Reduction of Phosphine Oxides, Sulfoxides, and <i>N</i> -Oxides with Hydrosilanes using a Borinic Acid Precatalyst. <i>ChemCatChem</i> , 2017, 9, 4460-4464.	3.7	21
9	Formamide Synthesis through Borinic Acid Catalysed Transamidation under Mild Conditions. <i>Chemistry - A European Journal</i> , 2016, 22, 5894-5898.	3.3	72
10	Catalytic Chemical Amide Synthesis at Room Temperature: One More Step Toward Peptide Synthesis. <i>Journal of Organic Chemistry</i> , 2015, 80, 4532-4544.	3.2	114
11	An Organocatalytic Access to Spiro[4.5]decanes and Spiro[4.6]undecanes Containing Aminolactones and 3-Aminopyrrolidines. <i>Synthesis</i> , 2015, 47, 2549-2553.	2.3	2
12	Borinic acid catalysed peptide synthesis. <i>Chemical Communications</i> , 2015, 51, 16084-16087.	4.1	76
13	An Easy Route to (Hetero)arylboronic Acids. <i>Chemistry - A European Journal</i> , 2014, 20, 6608-6612.	3.3	62
14	( $\beta$ )-Cytisine and Derivatives: Synthesis, Reactivity, and Applications. <i>Chemical Reviews</i> , 2014, 114, 712-778.	47.7	113
15	Expedient BINOL derivative arylations. <i>Tetrahedron Letters</i> , 2014, 55, 6420-6422.	1.4	7
16	Sequential One-Pot Access to Molecular Diversity through Aniline Aqueous Borylation. <i>Journal of Organic Chemistry</i> , 2014, 79, 10568-10580.	3.2	33
17	Diastereoselective organocatalytic Mannich access to azacyclic system en route to lyconadin A. <i>Tetrahedron Letters</i> , 2014, 55, 5074-5077.	1.4	3
18	Stereoselective access to heteroarylmethylene-substituted pyrrolidines: fully organocatalytic Mannich-hydroamination reactions. <i>Chemical Communications</i> , 2013, 49, 1651.	4.1	14

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19	An organocatalytic route to 2-heteroarylmethylene decorated <i>N</i> -arylpyrroles. Beilstein Journal of Organic Chemistry, 2013, 9, 1480-1486.	2.2	8
20	Synthesis of a New Chiral Sulfonic Acid. Synthesis, 2012, 44, 1349-1352.	2.3	10
21	Expanding the Scope of the Direct Regiospecific Asymmetric Aldol Reaction to Enones and Dienones Catalyzed by a BINOL-Derived Brønsted Acid. European Journal of Organic Chemistry, 2011, 2011, 6628-6631.	2.4	14
22	Brønsted Acid Catalyzed Asymmetric Aldol Reaction: A Complementary Approach to Enamine Catalysis. Organic Letters, 2010, 12, 3582-3585.	4.6	92
23	Synthesis of P,N-2,2'-biphenyl derivatives with central chirality. Science China Chemistry, 2010, 53, 1907-1913.	8.2	2
24	One-Pot Hydroxy Group Activation/Carbon-Carbon Bond Forming Sequence Using a Brønsted Base/Brønsted Acid System. Advanced Synthesis and Catalysis, 2010, 352, 2881-2886.	4.3	40
25	Chiral 3-aminopyrrolidines as a rigid diamino scaffold for organocatalysis and organometallic chemistry. Tetrahedron: Asymmetry, 2010, 21, 1511-1521.	1.8	16
26	Synthesis of BINOL derived phosphorodithioic acids as new chiral Brønsted acids and an improved synthesis of 3,3'-disubstituted H8-BINOL derivatives. Tetrahedron, 2009, 65, 10617-10622.	1.9	35
27	Asymmetric Malonic and Acetoacetic Acid Syntheses – A Century of Enantioselective Decarboxylative Protonations. European Journal of Organic Chemistry, 2008, 2008, 5493-5506.	2.4	81
28	4-Toluenesulfonic acid: an environmentally benign catalyst for Nazarov cyclizations. Tetrahedron Letters, 2008, 49, 2541-2545.	1.4	21
29	Desymmetrization of a <i>meso</i> -Allylic Acetal by Enantioselective Conjugate Elimination. Organic Letters, 2008, 10, 729-732.	4.6	15
30	3-Trifluoromethanesulfonamido-pyrrolidine: A General Organocatalyst for <i>anti</i> -Selective Mannich Reactions. Organic Letters, 2008, 10, 1029-1032.	4.6	62
31	Directed Ortho Metalation ~ Cross Coupling Strategies. <i>N</i> -Cumyl Arylsulfonamides. Facile Deprotection and Expedient Route to 7- and 4,7-Substituted Saccharins. Journal of Organic Chemistry, 2007, 72, 3199-3206.	3.2	44
32	A rapid and convenient synthesis of $\beta^2$ -proline. Tetrahedron Letters, 2007, 48, 5727-5730.	1.4	18
33	Aziridinium from <i>N,N</i> -Dibenzyl Serine Methyl Ester: Synthesis of Enantiomerically Pure $\beta^2$ -Amino and $\beta^{\pm 2}$ -Diamino Esters. Organic Letters, 2006, 8, 2183-2186.	4.6	45
34	Reeve's Synthesis of 2-Imino-4-thiazolidinone from Alkyl (Aryl) Trichloromethylcarbinol Revisited: A Three-Component Process from Aldehyde, Chloroform and Thiourea. ChemInform, 2004, 35, no.	0.0	0
35	Synthesis and Reactivity of Mixed Alkynylalanes by Direct Triethylamine-Catalyzed Alumination of Terminal Alkynes. ChemInform, 2004, 35, no.	0.0	0
36	Reeve's synthesis of 2-imino-4-thiazolidinone from alkyl (aryl) trichloromethylcarbinol revisited, a three-component process from aldehyde, chloroform and thiourea. Tetrahedron Letters, 2004, 45, 4449-4452.	1.4	50

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37	Synthesis and Reactivity of Mixed Alkynylalanes by Direct Triethylamine-Catalyzed Aluminatation of Terminal Alkynes. <i>Organic Letters</i> , 2004, 6, 2333-2336.	4.6	59
38	RECENT PROGRESS IN THE ASYMMETRIC SYNTHESIS OF $\hat{1}\pm$ -SUBSTITUTED PROPARGYLAMINES. <i>Organic Preparations and Procedures International</i> , 2002, 34, 467-492.	1.3	16
39	Isomerization of Chiral Non-Racemic $\hat{1}\pm$ -Substituted Propargylic Amines to Terminal Acetylenes. <i>European Journal of Organic Chemistry</i> , 2002, 2002, 2598.	2.4	7
40	Asymmetric synthesis of $\hat{1}^2$ -pseudopeptides from chiral 3,4-aziridinolactams. <i>Tetrahedron: Asymmetry</i> , 2002, 13, 995-1004.	1.8	14
41	Asymmetric synthesis of $\hat{1}\pm$ -substituted propynyl amines. Application to the preparation of a polysubstituted dihydroisoindoline framework. <i>Tetrahedron Letters</i> , 2001, 42, 3171-3173.	1.4	11
42	[2,3]-Meisenheimer rearrangement of N-allyl phenylglycinol derivatives. $Ni\text{-}C$ versus $Ci\text{-}C$ chirality transfer. <i>Tetrahedron Letters</i> , 2000, 41, 8279-8283.	1.4	28
43	Synthesis of Enantiomerically Pure $\hat{1}\pm$ -Substituted Propargylic Amines by Reaction of Organoaluminum Reagents with Oxazolidines. <i>Journal of Organic Chemistry</i> , 2000, 65, 6423-6426.	3.2	48
44	Diastereoselective alkynylation of chiral non-racemic oxazolidines with mixed organoaluminum compounds. <i>Tetrahedron Letters</i> , 1999, 40, 2935-2938.	1.4	33