# Vittorio Pace

## List of Publications by Year in Descending Order



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| 111 papers | 2,976 citations | 33 h-index | 50 g-index |
| 150 ext. papers | 3,471 ext. citations | 5 avg, IF | 5.7 L-index |

| # | Paper | IF | Citations |
|---|---|---|---|
| 111 | Synthesis of stable -fluoromethyl putative carbanions via a chemoselective reduction-monofluoromethylation sequence of diselenides under sustainable conditions. *Tetrahedron*, **2021**, 85, 131921 | 2.4 | 6 |
| 110 | Carbenoid-Mediated Homologation Tactics for Assembling (Fluorinated) Epoxides and Aziridines. *Synlett*, **2021**, 32, 551-560 | 2.2 | 7 |
| 109 | (Difluoromethyl)trimethylsilane (TMSCHF2): A Useful Difluoromethylating Nucleophilic Source. *Australian Journal of Chemistry*, **2021**, 74, 623 | 1.2 | 2 |
| 108 | Taking advantage of lithium monohalocarbenoid intrinsic -elimination in 2-MeTHF: controlled epoxide ring-opening to halohydrins. *Organic and Biomolecular Chemistry*, **2021**, 19, 2038-2043 | 3.9 | 5 |
| 107 | Consecutive and Selective Double Methylene Insertion of Lithium Carbenoids to Isothiocyanates: A Direct Assembly of Four-Membered Sulfur-Containing Cycles. *Angewandte Chemie - International Edition*, **2021**, 60, 24854-24858 | 16.4 | 3 |
| 106 | Direct and straightforward transfer of C1 functionalized synthons to phosphorous electrophiles for accessing gem-P-containing methanes. *Organic and Biomolecular Chemistry*, **2021**, 19, 2425-2429 | 3.9 | 0 |
| 105 | Pseudo-Dipeptide Bearing -Difluoromethyl Ketone Moiety as Electrophilic Warhead with Activity against Coronaviruses. *International Journal of Molecular Sciences*, **2021**, 22, | 6.3 | 6 |
| 104 | Direct and Chemoselective Electrophilic Monofluoromethylation of Heteroatoms (-, , -, -, -) with Fluoroiodomethane. *Organic Letters*, **2020**, 22, 1345-1349 | 6.2 | 15 |
| 103 | Straightforward and direct access to -seleno- amines and sulfonylamides via the controlled addition of phenylselenomethyllithium (LiCH2SePh) to imines. *Tetrahedron*, **2020**, 76, 131220 | 2.4 | 2 |
| 102 | A Combination of Pharmacophore and Docking-based Virtual Screening to Discover new Tyrosinase Inhibitors. *Molecular Informatics*, **2020**, 39, e1900054 | 3.8 | 7 |
| 101 | Electrophilicity Scale of Activated Amides: O NMR and N NMR Chemical Shifts of Acyclic Twisted Amides in N-C(O) Cross-Coupling. *Chemistry - A European Journal*, **2020**, 26, 16246-16250 | 4.8 | 5 |
| 100 | Halogen-Imparted Reactivity in Lithium Carbenoid Mediated Homologations of Imine Surrogates: Direct Assembly of bis-Trifluoromethyl--Diketiminates and the Dual Role of LiCHI. *Angewandte Chemie - International Edition*, **2020**, 59, 20852-20857 | 16.4 | 8 |
| 99 | Halogen-Imparted Reactivity in Lithium Carbenoid Mediated Homologations of Imine Surrogates: Direct Assembly of bis-Trifluoromethyl--Diketiminates and the Dual Role of LiCH2I. *Angewandte Chemie*, **2020**, 132, 21038-21043 | 3.6 | 3 |
| 98 | Straightforward chemoselective access to unsymmetrical dithioacetals through a thiosulfonate homologation-nucleophilic substitution sequence. *Chemical Communications*, **2020**, 56, 12395-12398 | 5.8 | 6 |
| 97 | Consecutive C1-Homologation / Displacement Strategy for Converting Thiosulfonates into O,S-Oxothioacetals. *Advanced Synthesis and Catalysis*, **2020**, 362, 5444-5449 | 5.6 | 2 |
| 96 | Chemoselective Homologation-Deoxygenation Strategy Enabling the Direct Conversion of Carbonyls into ()-Halomethyl-Alkanes. *Organic Letters*, **2020**, 22, 7629-7634 | 6.2 | 12 |
| 95 | Modular and Chemoselective Strategy for Accessing (Distinct) -Dihaloketones from Weinreb Amides and Dihalomethyllithiums. *Advanced Synthesis and Catalysis*, **2020**, 362, 5056-5061 | 5.6 | 4 |

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| 94 | Novel Dual Ligands Targeting Sigma1 Receptor and Acetylcholinesterase Endowed with Antioxidant Properties. *Proceedings (mdpi)*, **2019**, 22, 49 | 0.3 | |
| 93 | Chemoselective reduction of isothiocyanates to thioformamides mediated by the Schwartz reagent. *Organic and Biomolecular Chemistry*, **2019**, 17, 1970-1978 | 3.9 | 14 |
| 92 | Weinreb Amides as Privileged Acylating Agents for Accessing -Substituted Ketones. *Synthesis*, **2019**, 51, 2792-2808 | 2.9 | 24 |
| 91 | Palladium-Catalyzed Regioselective Syn-Chloropalladation-Olefin Insertion-Oxidative Chlorination Cascade: Synthesis of Dichlorinated Tetrahydroquinolines. *Organic Letters*, **2019**, 21, 3465-3469 | 6.2 | 13 |
| 90 | O NMR and N NMR chemical shifts of sterically-hindered amides: ground-state destabilization in amide electrophilicity. *Chemical Communications*, **2019**, 55, 4423-4426 | 5.8 | 5 |
| 89 | Biocatalyzed Synthesis of Statins: A Sustainable Strategy for the Preparation of Valuable Drugs. *Catalysts*, **2019**, 9, 260 | 4 | 23 |
| 88 | Direct Access to 9-Chloro-1-benzo[]furo[3,4-]azepin-1-ones via Palladium(II)-Catalyzed Intramolecular -Oxypalladation/Olefin Insertion/sp-C-H Bond Activation Cascade. *Organic Letters*, **2019**, 21, 5784-5788 | 6.2 | 12 |
| 87 | Direct and Chemoselective Synthesis of Tertiary Difluoroketones via Weinreb Amide Homologation with a CHF-Carbene Equivalent. *Organic Letters*, **2019**, 21, 8261-8265 | 6.2 | 30 |
| 86 | The synthetic versatility of the TiffeneauDemjanov chemistry in homologation tactics. *Monatshefte F Chemie*, **2019**, 150, 2011-2019 | 1.4 | 7 |
| 85 | Highly chemoselective difluoromethylative homologation of iso(thio)cyanates: expeditious access to unprecedented difluoro(thio)amides. *Chemical Communications*, **2019**, 55, 12960-12963 | 5.8 | 13 |
| 84 | A Straightforward Homologation of Carbon Dioxide with Magnesium Carbenoids en Route to -Halocarboxylic Acids. *Advanced Synthesis and Catalysis*, **2019**, 361, 1001-1006 | 5.6 | 7 |
| 83 | Modular and Chemoselective Strategy for the Direct Access to -Fluoroepoxides and Aziridines via the Addition of Fluoroiodomethyllithium to Carbonyl-Like Compounds. *Organic Letters*, **2019**, 21, 584-588 | 6.2 | 43 |
| 82 | Sustainable Asymmetric Organolithium Chemistry: Enantio- and Chemoselective Acylations through Recycling of Solvent, Sparteine, and Weinreb "Amine". *ChemSusChem*, **2019**, 12, 1147-1154 | 8.3 | 11 |
| 81 | Telescoped, Divergent, Chemoselective C1 and C1-C1 Homologation of Imine Surrogates: Access to Quaternary Chloro- and Halomethyl-Trifluoromethyl Aziridines. *Angewandte Chemie - International Edition*, **2019**, 58, 2479-2484 | 16.4 | 37 |
| 80 | Cyclopentyl Methyl Ether: An Elective Ecofriendly Ethereal Solvent in Classical and Modern Organic Chemistry. *ChemSusChem*, **2019**, 12, 40-70 | 8.3 | 73 |
| 79 | Design, Synthesis, and Pharmacological Evaluation of Novel 2/3 Subunit-Selective -Aminobutyric Acid Type A (GABA) Receptor Modulators. *Journal of Medicinal Chemistry*, **2019**, 62, 317-341 | 8.3 | 6 |
| 78 | Substituted -Sulfur Methyl Carbanions: Effective Homologating Agents for the Chemoselective Preparation of -Oxo Thioethers from Weinreb Amides. *European Journal of Organic Chemistry*, **2018**, 2018, 2466-2470 | 3.2 | 15 |
| 77 | Expeditious and Chemoselective Synthesis of -Aryl and -Alkyl Selenomethylketones via Homologation Chemistry. *Organic Letters*, **2018**, 20, 2685-2688 | 6.2 | 28 |

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| 76 | Merging lithium carbenoid homologation and enzymatic reduction: A combinative approach to the HIV-protease inhibitor Nelfinavir. *Tetrahedron*, **2018**, 74, 2211-2217 | 2.4 | 18 |
| 75 | □Arylamino Diazoketones: Diazomethane-Loading Controlled Synthesis, Spectroscopic Investigations, and Structural X-ray Analysis. *Journal of Organic Chemistry*, **2018**, 83, 4336-4347 | 4.2 | 11 |
| 74 | Fluoroiodomethane: A versatile CH2F Source. *Australian Journal of Chemistry*, **2018**, 71, 473 | 1.2 | 11 |
| 73 | An unusual thionyl chloride-promoted C-C bond formation to obtain 4,4Sbipyrazolones. *Beilstein Journal of Organic Chemistry*, **2018**, 14, 1287-1292 | 2.5 | 6 |
| 72 | Homologation of halostannanes with carbenoids: a convenient and straightforward one-step access to □functionalized organotin reagents. *Chemical Communications*, **2018**, 54, 10112-10115 | 5.8 | 13 |
| 71 | Recent advances in the synthesis and reactivity of spiro-epoxyoxindoles. *Chemistry of Heterocyclic Compounds*, **2018**, 54, 389-393 | 1.4 | 6 |
| 70 | A practical guide for using lithium halocarbenoids in homologation reactions. *Monatshefte F□ Chemie*, **2018**, 149, 1285-1291 | 1.4 | 4 |
| 69 | Telescoped, Divergent, Chemoselective C1 and C1-C1 Homologation of Imine Surrogates: Access to Quaternary Chloro- and Halomethyl-Trifluoromethyl Aziridines. *Angewandte Chemie*, **2018**, 131, 2501 | 3.6 | |
| 68 | Identification of dual Sigma1 receptor modulators/acetylcholinesterase inhibitors with antioxidant and neurotrophic properties, as neuroprotective agents. *European Journal of Medicinal Chemistry*, **2018**, 158, 353-370 | 6.8 | 10 |
| 67 | Easy as one, two, three. *Nature Chemistry*, **2018**, 10, 1081-1082 | 17.6 | 2 |
| 66 | Homologation chemistry with nucleophilic □substituted organometallic reagents: chemocontrol, new concepts and (solved) challenges. *Chemical Communications*, **2018**, 54, 6692-6704 | 5.8 | 35 |
| 65 | New Perspectives in Lithium Carbenoid Mediated Homologations. *Synlett*, **2017**, 28, 879-888 | 2.2 | 38 |
| 64 | A greener and efficient access to substituted four- and six-membered sulfur-bearing heterocycles. *Organic and Biomolecular Chemistry*, **2017**, 15, 5000-5015 | 3.9 | 14 |
| 63 | Recent advancements on the use of 2-methyltetrahydrofuran in organometallic chemistry. *Monatshefte F□ Chemie*, **2017**, 148, 37-48 | 1.4 | 55 |
| 62 | Efficient Access to All-Carbon Quaternary and Tertiary □Functionalized Homoallyl-type Aldehydes from Ketones. *Angewandte Chemie*, **2017**, 129, 12851-12856 | 3.6 | 18 |
| 61 | Synthesis of tetrasubstituted pyrazoles containing pyridinyl substituents. *Beilstein Journal of Organic Chemistry*, **2017**, 13, 895-902 | 2.5 | 4 |
| 60 | Exploiting a "Beast" in Carbenoid Chemistry: Development of a Straightforward Direct Nucleophilic Fluoromethylation Strategy. *Journal of the American Chemical Society*, **2017**, 139, 13648-13651 | 16.4 | 79 |
| 59 | Efficient Access to All-Carbon Quaternary and Tertiary □Functionalized Homoallyl-type Aldehydes from Ketones. *Angewandte Chemie - International Edition*, **2017**, 56, 12677-12682 | 16.4 | 50 |

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| 58 | Evidence and isolation of tetrahedral intermediates formed upon the addition of lithium carbenoids to Weinreb amides and N-acylpyrroles. *Chemical Communications*, **2017**, 53, 9498-9501 | 5.8 | 39 |
| 57 | Lithium Halomethylcarbenoids: Preparation and Use in the Homologation of Carbon Electrophiles. *Chemical Record*, **2016**, 16, 2061-76 | 6.6 | 39 |
| 56 | Structures of Highly Twisted Amides Relevant to Amide N-C Cross-Coupling: Evidence for Ground-State Amide Destabilization. *Chemistry - A European Journal*, **2016**, 22, 14494-8 | 4.8 | 87 |
| 55 | Isocyanates and isothiocyanates as versatile platforms for accessing (thio)amide-type compounds. *Organic and Biomolecular Chemistry*, **2016**, 14, 7848-54 | 3.9 | 39 |
| 54 | Palladium-Catalyzed Internal Nucleophile-Assisted Hydration-Olefin Insertion Cascade: Diastereoselective Synthesis of 2,3-Dihydro-1H-inden-1-ones. *Organic Letters*, **2016**, 18, 3442-5 | 6.2 | 21 |
| 53 | Potassium-Exchanged Zirconium Hydrogen Phosphate [□Zr(KPO4)2]-Catalyzed Synthesis of 2-Amino-4H-pyran Derivatives under Solvent-Free Conditions. *Synthesis*, **2016**, 48, 1533-1540 | 2.9 | 12 |
| 52 | Bromomethyllithium-mediated chemoselective homologation of disulfides to dithioacetals. *Chemical Communications*, **2016**, 52, 2639-42 | 5.8 | 47 |
| 51 | Novel Enantiopure Sigma Receptor Modulators: Quick (Semi-)Preparative Chiral Resolution via HPLC and Absolute Configuration Assignment. *Molecules*, **2016**, 21, | 4.8 | 6 |
| 50 | The use of the Comins-Meyers Amide in Synthetic Chemistry: An Overview. *Natural Product Communications*, **2016**, 11, 1934578X1601101 | 0.9 | 2 |
| 49 | Dynamic Kinetic Resolution via Hydrolase□Metal Combo Catalysis **2016**, 373-396 | | 2 |
| 48 | Chemoselective Addition of Halomethyllithiums to Functionalized Isatins:A Straightforward Access to Spiro-Epoxyoxindoles. *Advanced Synthesis and Catalysis*, **2016**, 358, 172-177 | 5.6 | 40 |
| 47 | Chemoselective Schwartz Reagent Mediated Reduction of Isocyanates to Formamides. *Organic Letters*, **2016**, 18, 2750-3 | 6.2 | 52 |
| 46 | Highly efficient synthesis of functionalized □oxyketones via Weinreb amides homologation with □oxygenated organolithiums. *Chemical Communications*, **2016**, 52, 7584-7 | 5.8 | 35 |
| 45 | Synthesis and biological evaluation of new aryl-alkyl(alkenyl)-4-benzylpiperidines, novel Sigma Receptor (SR) modulators, as potential anticancer-agents. *European Journal of Medicinal Chemistry*, **2016**, 124, 649-665 | 6.8 | 21 |
| 44 | Synthesis of 6,12-Epiminodibenzo[b,f][1,5]diazocines via an Ytterbium Triflate-Catalyzed, AB Three-Component Reaction. *Journal of Organic Chemistry*, **2016**, 81, 9687-9694 | 4.2 | 15 |
| 43 | Eco-friendly chemoselective N-functionalization of isatins mediated by supported KF in 2-MeTHF. *Green Chemistry*, **2015**, 17, 4194-4197 | 10 | 19 |
| 42 | Diethylaluminium Azide: A Versatile Reagent in Organic Synthesis. *Australian Journal of Chemistry*, **2015**, 68, 703 | 1.2 | 3 |
| 41 | Chemoselective efficient synthesis of functionalized □oxonitriles through cyanomethylation of Weinreb amides. *Organic and Biomolecular Chemistry*, **2015**, 13, 1969-73 | 3.9 | 31 |

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| 40 | A Robust, Eco-Friendly Access to Secondary Thioamides through the Addition of Organolithium Reagents to Isothiocyanates in Cyclopentyl Methyl Ether (CPME). *Chemistry - A European Journal*, **2015**, 21, 18966-70 | 4.8 | 34 |
| 39 | Chemoselective Additions of Chloromethyllithium Carbenoid to Cyclic Enones: A Direct Access to Chloromethyl Allylic Alcohols. *Advanced Synthesis and Catalysis*, **2014**, 356, 1761-1766 | 5.6 | 25 |
| 38 | Cu(I)-NHC catalyzed asymmetric silyl transfer to unsaturated lactams and amides. *Organic Letters*, **2014**, 16, 476-9 | 6.2 | 78 |
| 37 | 2-Methyltetrahydrofuran **2014**, 1-6 | | 1 |
| 36 | Halomethyllithium Carbenoids: Versatile Reagents for the Homologation of Electrophilic Carbon Units. *Australian Journal of Chemistry*, **2014**, 67, 311 | 1.2 | 23 |
| 35 | Homologation of Isocyanates with Lithium Carbenoids: A Straightforward Access to α-Halomethyl- and α,α-Dihalomethylamides. *Synthesis*, **2014**, 46, 2897-2909 | 2.9 | 37 |
| 34 | Increasing the Reactivity of Amides towards Organometallic Reagents: An Overview. *Advanced Synthesis and Catalysis*, **2014**, 356, 3697-3736 | 5.6 | 168 |
| 33 | Expanding the Synthetic Portfolio of Organolithiums: Direct Use in Catalytic Cross-Coupling Reactions. *ChemCatChem*, **2014**, 6, 1516-1519 | 5.2 | 24 |
| 32 | Chemoenzymatic synthesis of carbohydrates as antidiabetic and anticancer drugs. *Current Topics in Medicinal Chemistry*, **2014**, 14, 2694-711 | 3 | 6 |
| 31 | Synthesis of α,β-unsaturated α-haloketones through the chemoselective addition of halomethyllithiums to Weinreb amides. *Journal of Organic Chemistry*, **2013**, 78, 7764-70 | 4.2 | 45 |
| 30 | NHC-Cu(I) catalysed asymmetric conjugate silyl transfer to unsaturated lactones: application in kinetic resolution. *Chemical Communications*, **2013**, 49, 5150-2 | 5.8 | 52 |
| 29 | Addition of lithium carbenoids to isocyanates: a direct access to synthetically useful N-substituted 2-haloacetamides. *Chemical Communications*, **2013**, 49, 8383-5 | 5.8 | 65 |
| 28 | Highly efficient and environmentally benign preparation of Weinreb amides in the biphasic system 2-MeTHF/water. *RSC Advances*, **2013**, 3, 10158 | 3.7 | 20 |
| 27 | Highly efficient and chemoselective α-iodination of acrylate esters through Morita-Baylis-Hillman-type chemistry. *Organic and Biomolecular Chemistry*, **2013**, 11, 1085-8 | 3.9 | 13 |
| 26 | Chemoselective Activation Strategies of Amidic Carbonyls towards Nucleophilic Reagents. *Australian Journal of Chemistry*, **2013**, 66, 507 | 1.2 | 72 |
| 25 | Chemoselective Synthesis of N-Substituted α-Amino-α′-chloro Ketones via Chloromethylation of Glycine-Derived Weinreb Amides. *Advanced Synthesis and Catalysis*, **2013**, 355, 919-926 | 5.6 | 33 |
| 24 | Chemoselective oxidative hydrolysis of EWG protected α-arylamino vinyl bromides to α-arylamino-α′-bromoacetones. *Tetrahedron Letters*, **2013**, 54, 4369-4372 | 2 | 8 |
| 23 | Chemoselective CaO-mediated acylation of alcohols and amines in 2-methyltetrahydrofuran. *ChemSusChem*, **2013**, 6, 905-10 | 8.3 | 16 |

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| 22 | Biocatalyzed On Water Synthesis of Chiral Building Blocks for the Preparation of Anti-Cancer Drugs: a GreenerApproach. *Current Organic Chemistry*, **2013**, 17, 1132-1157 | 1.7 | 5 |
| 21 | Amino-halomethylketones: synthetic methodologies and pharmaceutical applications as serine and cysteine protease inhibitors. *Mini-Reviews in Medicinal Chemistry*, **2013**, 13, 988-96 | 3.2 | 9 |
| 20 | Highly chemoselective synthesis of aryl allylic sulfoxides through calcium hypobromite oxidation of aryl allylic sulfides. *Tetrahedron Letters*, **2012**, 53, 967-972 | 2 | 18 |
| 19 | A straightforward and general access to phthalimido-substituted propan-2-ones. *Tetrahedron Letters*, **2012**, 53, 5106-5109 | 2 | 9 |
| 18 | 2-Methyltetrahydrofuran (2-MeTHF): a biomass-derived solvent with broad application in organic chemistry. *ChemSusChem*, **2012**, 5, 1369-79 | 8.3 | 405 |
| 17 | Dynamic Kinetic Resolution via Hydrolase-Metal Combo Catalysis in Stereoselective Synthesis of Bioactive Compounds. *Advanced Synthesis and Catalysis*, **2012**, 354, 2585-2611 | 5.6 | 63 |
| 16 | Robust eco-friendly protocol for the preparation of hydroxy-acetylenic esters by sequential one-pot elimination addition of 2-bromoacrylates to aldehydes promoted by LTMP in 2-MeTHF. *Green Chemistry*, **2012**, 14, 1859 | 10 | 26 |
| 15 | 2-Methyltetrahydrofuran: A Versatile Eco-Friendly Alternative to THF in Organometallic Chemistry. *Australian Journal of Chemistry*, **2012**, 65, 301 | 1.2 | 42 |
| 14 | Chemoenzymatic synthesis of chiral unsymmetrical benzoin esters. *Tetrahedron*, **2011**, 67, 7321-7329 | 2.4 | 23 |
| 13 | Structural bases for understanding the stereoselectivity in ketone reductions with ADH from Thermus thermophilus: A quantitative model. *Journal of Molecular Catalysis B: Enzymatic*, **2011**, 70, 23-31 | | 14 |
| 12 | Highly efficient chemoselective N-TBS protection of anilines under exceptional mild conditions in the eco-friendly solvent 2-methyltetrahydrofuran. *Green Chemistry*, **2011**, 13, 1986 | 10 | 35 |
| 11 | Highly regioselective control of 1,2-addition of organolithiums to unsaturated compounds promoted by lithium bromide in 2-methyltetrahydrofuran: a facile and eco-friendly access to allylic alcohols and amines. *Tetrahedron*, **2011**, 67, 2670-2675 | 2.4 | 49 |
| 10 | Highly Regioselective and Efficient Synthesis of Aminoepoxides by Ring Closure of Aminohalohydrins Mediated by KF-Celite. *Synlett*, **2011**, 2011, 1831-1834 | 2.2 | 10 |
| 9 | 1,3-Dichloroacetone. *Synlett*, **2010**, 2010, 2825-2826 | 2.2 | 2 |
| 8 | Improved Arndt-Eistert synthesis of alpha-diazoketones requiring minimal diazomethane in the presence of calcium oxide as acid scavenger. *Journal of Organic Chemistry*, **2010**, 75, 5760-3 | 4.2 | 54 |
| 7 | 2-Methyltetrahydrofuran as a suitable green solvent for phthalimide functionalization promoted by supported KF. *Green Chemistry*, **2010**, 12, 1380 | 10 | 56 |
| 6 | Highly Efficient Synthesis of New Arylamino-chloropropan-2-ones via Oxidative Hydrolysis of Vinyl Chlorides Promoted by Calcium Hypochlorite. *Advanced Synthesis and Catalysis*, **2009**, 351, 3199-3206 | 5.6 | 22 |
| 5 | Efficient HornerWadsworthEmmons intramolecular cyclisation of a N-substituted phthalimide promoted by KF-Alumina: a general tool for the synthesis of functionalised isoindolinones. *Tetrahedron Letters*, **2009**, 50, 3050-3053 | 2 | 27 |

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| 4 | Effective monoallylation of anilines catalyzed by supported KF. *Organic Letters*, **2007**, 9, 2661-4 | 6.2 | 37 |
| 3 | Preparation of 2-Amino-4H-chromene Derivatives from Coumarins in Basic Media. *European Journal of Organic Chemistry*, **2006**, 2006, 746-751 | 3.2 | 7 |
| 2 | (Difluoromethyl)trimethylsilane (TMSCHF2)1-17 | | |
| 1 | Consecutive and Selective Double Methylene Insertion of Lithium Carbenoids to Isothiocyanates: A Direct Assembly of Four-membered Sulfur-Containing Cycles. *Angewandte Chemie*, | 3.6 | |