Rings in Drugs

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Citation Report

#	Article	IF	CITATIONS
2	Synthesis of Heterocyclic Triads by Pd-Catalyzed Cross-Couplings and Evaluation of Their Cell-Specific Toxicity Profile. Organic Letters, 2014, 16, 2034-2037.	2.4	10
3	Rhodium atalyzed Arylative Cyclization for the Enantioselective Synthesis of (Trifluoromethyl)cyclobutanols. Chemistry - A European Journal, 2014, 20, 14194-14197.	1.7	49
4	Rapid Assembly of Functionalised Spirocyclic Indolines by Palladium atalysed Dearomatising Diallylation of Indoles with Allyl Acetate. Chemistry - A European Journal, 2014, 20, 13375-13381.	1.7	18
5	Pyridine synthesis by [4 + 2] cycloadditions of 1-azadienes: hetero-Diels Alder and transition metal-catalysed approaches. Organic Chemistry Frontiers, 2014, 1, 1010-1015.	2.3	73
6	Highly regioselective lithiation of pyridines bearing an oxetane unit by n-butyllithium. Chemical Communications, 2014, 50, 8908-8911.	2.2	23
7	Regio- and Stereospecific Synthesis of C-3 Functionalized Proline Derivatives by Palladium Catalyzed Directed C(sp ³)–H Arylation. Organic Letters, 2014, 16, 4956-4959.	2.4	134
8	Alkene Carboboration Enabled by Synergistic Catalysis. Chemistry - A European Journal, 2014, 20, 12032-12036.	1.7	154
9	Analysis of the Structural Diversity, Substitution Patterns, and Frequency of Nitrogen Heterocycles among U.S. FDA Approved Pharmaceuticals. Journal of Medicinal Chemistry, 2014, 57, 10257-10274.	2.9	3,996
10	Physicochemical Descriptors of Aromatic Character and Their Use in Drug Discovery. Journal of Medicinal Chemistry, 2014, 57, 7206-7215.	2.9	74
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12	Sultones and Sultines via a Julia–Kocienski Reaction of Epoxides. Angewandte Chemie - International Edition, 2015, 54, 15236-15240.	7.2	22
15	Diastereo―and Enantioselective Iridium Catalyzed Carbonyl (αâ€Cyclopropyl)allylation via Transfer Hydrogenation. Chemistry - A European Journal, 2015, 21, 12903-12907.	1.7	17
16	Mechanismâ€Ðriven Elaboration of an Enantioselective Bromocyclopropanation Reaction of Allylic Alcohols. Angewandte Chemie - International Edition, 2015, 54, 14108-14112.	7.2	28
17	Catalytic Synthesis of Nâ€Unprotected Piperazines, Morpholines, and Thiomorpholines from Aldehydes and SnAP Reagents. Angewandte Chemie - International Edition, 2015, 54, 10884-10888.	7.2	64
18	Method for Systematic Assessment of Chemical Changes in Molecular Scaffolds with Conserved Topology and Application to the Analysis of Scaffoldâ€Activity Relationships. Molecular Informatics, 2015, 34, 531-549.	1.4	1
20	Four-Membered Ring Systems. Progress in Heterocyclic Chemistry, 2015, 27, 87-115.	0.5	1
21	Understanding the foundations of the structural similarities between marketed drugs and endogenous human metabolites. Frontiers in Pharmacology, 2015, 6, 105.	1.6	27
	KOtBu-mediated annulation of acetonitrile with aldehyde: synthesis of substituted		

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24	Design, synthesis and biological evaluation of paralleled Aza resveratrol–chalcone compounds as potential anti-inflammatory agents for the treatment of acute lung injury. Bioorganic and Medicinal Chemistry Letters, 2015, 25, 2998-3004.	1.0	26
26	Building polyfunctional piperidines: a stereoselective strategy of a three-component Mannich reaction inspired by biosynthesis and applications in the synthesis of natural alkaloids (+)-241D; (â^')-241D; isosolenopsin A and (â^')-epimyrtine. RSC Advances, 2015, 5, 18894-18908.	1.7	16
27	Seven-membered ring scaffolds for drug discovery: Access to functionalised azepanes and oxepanes through diazocarbonyl chemistry. Bioorganic and Medicinal Chemistry, 2015, 23, 2730-2735.	1.4	37
28	Efficient Synthesis of 5â€Chalcogenylâ€1,3â€oxazinâ€2â€ones by Chalcogenâ€Mediated Yne–Carbamate Cyc An Experimental and Theoretical Study. European Journal of Organic Chemistry, 2015, 2015, 1020-1027.	isation: 1.2	16
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30	Novel Scaffold Fingerprint (SFP): Applications in Scaffold Hopping and Scaffold-Based Selection of Diverse Compounds. Journal of Chemical Information and Modeling, 2015, 55, 1-18.	2.5	26
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32	Structural diversity and potency range distribution of scaffolds from compounds active against current pharmaceutical targets. Future Medicinal Chemistry, 2015, 7, 111-122.	1.1	3
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43	Highly efficient regioselective synthesis of pyrroles via a tandem enamine formation–Michael addition–cyclization sequence under catalyst- and solvent-free conditions. Green Chemistry, 2015, 17, 3415-3423.	4.6	36
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63	Dienamine Activation of Diazoenals: Application to the Direct Synthesis of Functionalized 1,4â€Oxazines. Angewandte Chemie - International Edition, 2016, 55, 7831-7835.	7.2	37
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