

Michel Baltas

List of Publications by Year in descending order

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77
papers

1,940
citations

236925

25
h-index

289244

40
g-index

84
all docs

84
docs citations

84
times ranked

2742
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|--|-----|-----------|
| 1 | Elucidation of the Diels-Alder Reaction Kinetics between Diphenylfulvene and Maleimide by Mechanochemistry and in Solution. <i>ACS Sustainable Chemistry and Engineering</i> , 2021, 9, 4453-4462. | 6.7 | 13 |
| 2 | Synthesis of Biologically Relevant 1,2,3- and 1,3,4-Triazoles: From Classical Pathway to Green Chemistry. <i>Molecules</i> , 2021, 26, 5667. | 3.8 | 18 |
| 3 | Synthesis and Antiplasmodial Activity of Novel Fosmidomycin Derivatives and Conjugates with Artemisinin and Aminochloroquine. <i>Molecules</i> , 2020, 25, 4858. | 3.8 | 10 |
| 4 | Synthesis of Novel G Factor or Chloroquine-Artemisinin Hybrids and Conjugates with Potent Antiplasmodial Activity. <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 921-927. | 2.8 | 23 |
| 5 | Study of the Two Steps and One-Pot Two-Step Mechanochemical Synthesis of Annulated 1,2,4-Triazoles. <i>ACS Sustainable Chemistry and Engineering</i> , 2020, 8, 3114-3125. | 6.7 | 10 |
| 6 | In Silico Repositioning of Cannabigerol as a Novel Inhibitor of the Enoyl Acyl Carrier Protein (ACP) Reductase (InhA). <i>Molecules</i> , 2019, 24, 2567. | 3.8 | 22 |
| 7 | Synthesis, In Silico, and In Vitro Evaluation of Anti-Leishmanial Activity of Oxadiazoles and Indolizine Containing Compounds Flagged against Anti-Targets. <i>Molecules</i> , 2019, 24, 1282. | 3.8 | 15 |
| 8 | Synthesis and biological evaluation of diarylheptanoids as potential antioxidant and anti-inflammatory agents. <i>European Journal of Medicinal Chemistry</i> , 2018, 144, 289-299. | 5.5 | 24 |
| 9 | Effect of the Nature of Surfactant on the Reactivity of C,N-diphenylnitrone towards Acrylonitrile in Different Microemulsions Systems. <i>Chemistry Journal of Moldova</i> , 2018, 13, 82-88. | 0.6 | 1 |
| 10 | Comprehensive experimental investigation of mechanically induced 1,4-diazines synthesis in solid state. <i>Tetrahedron</i> , 2017, 73, 2305-2310. | 1.9 | 14 |
| 11 | Total Synthesis of Tedarene A. <i>Journal of Natural Products</i> , 2017, 80, 1623-1630. | 3.0 | 8 |
| 12 | Mechanochemical Synthesis and Biological Evaluation of Novel Isoniazid Derivatives with Potent Antitubercular Activity. <i>Molecules</i> , 2017, 22, 1457. | 3.8 | 71 |
| 13 | 4-Hydroxynonenal Contributes to Angiogenesis through a Redox-Dependent Sphingolipid Pathway: Prevention by Hydralazine Derivatives. <i>Oxidative Medicine and Cellular Longevity</i> , 2017, 2017, 1-11. | 4.0 | 12 |
| 14 | Structure-Based Virtual Ligand Screening on the XRCC4/DNA Ligase IV Interface. <i>Scientific Reports</i> , 2016, 6, 22878. | 3.3 | 17 |
| 15 | Lowering the Activation Energy under Mechanochemical Conditions: The Case of 2,3-diphenylquinoxaline. <i>ChemistrySelect</i> , 2016, 1, 984-988. | 1.5 | 13 |
| 16 | Pyrrolidinone and pyrrolidine derivatives: Evaluation as inhibitors of InhA and Mycobacterium tuberculosis. <i>European Journal of Medicinal Chemistry</i> , 2016, 123, 462-475. | 5.5 | 33 |
| 17 | Synthesis and evaluation of antioxidant phenolic diaryl hydrazones as potent antiangiogenic agents in atherosclerosis. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 3571-3578. | 3.0 | 14 |
| 18 | Triazolophthalazines: Easily Accessible Compounds with Potent Antitubercular Activity. <i>ChemMedChem</i> , 2016, 11, 1078-1089. | 3.2 | 12 |

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|----|--|-----|-----------|
| 19 | Identification and optimization of hydrazone-gallate derivatives as specific inhibitors of DNA methyltransferase 3A. <i>Future Medicinal Chemistry</i> , 2016, 8, 373-380. | 2.3 | 12 |
| 20 | Peptide Synthesis in Ionic Liquids (PEPSIL): All You Need is in the Toolbox!. <i>French-Ukrainian Journal of Chemistry</i> , 2016, 4, 3-13. | 0.4 | 0 |
| 21 | Structure of adducts of isoindolo[2,1-a]benzimidazole derivatives with maleimides. <i>Journal of Molecular Structure</i> , 2015, 1084, 177-181. | 3.6 | 1 |
| 22 | Small molecules inhibitors of plasminogen activator inhibitor-1 – An overview. <i>European Journal of Medicinal Chemistry</i> , 2015, 92, 619-636. | 5.5 | 56 |
| 23 | SnCl ₂ /EtOH-Mediated Synthesis of Novel 4-Ethoxy- and 4-Chloroindazoles Bearing Sulfonamide Moieties. <i>Synthetic Communications</i> , 2015, 45, 2005-2013. | 2.1 | 0 |
| 24 | Design, synthesis and evaluation of new GEQ derivatives as inhibitors of InhA enzyme and Mycobacterium tuberculosis growth. <i>European Journal of Medicinal Chemistry</i> , 2015, 101, 218-235. | 5.5 | 43 |
| 25 | Crystal structure of the enoyl-ACP reductase of Mycobacterium tuberculosis (InhA) in the apo-form and in complex with the active metabolite of isoniazid pre-formed by a biomimetic approach. <i>Journal of Structural Biology</i> , 2015, 190, 328-337. | 2.8 | 31 |
| 26 | Synthesis and evaluation of 1,2-hydroxytriazoles and related compounds as antitubercular agents. <i>French-Ukrainian Journal of Chemistry</i> , 2015, 3, 82-96. | 0.4 | 1 |
| 27 | Unexpected copper mediated benzyl O→O migration during an Ullmann ether coupling. <i>Tetrahedron Letters</i> , 2014, 55, 528-530. | 1.4 | 4 |
| 28 | Synthesis in ionic liquids only: access to 1,2-oxo-1,3-thio-esters via Mukaiyama coupling. <i>Tetrahedron Letters</i> , 2014, 55, 1353-1356. | 1.4 | 6 |
| 29 | Synthesis of 1,2,3-triazoles Through Copper Iodide Catalyzed Oxygenation. <i>European Journal of Organic Chemistry</i> , 2014, 2014, 654-659. | 2.4 | 17 |
| 30 | Synthesis of 3-heteryl substituted pyrrolidine-2,5-diones via catalytic Michael reaction and evaluation of their inhibitory activity against InhA and Mycobacterium tuberculosis. <i>European Journal of Medicinal Chemistry</i> , 2014, 71, 46-52. | 5.5 | 38 |
| 31 | LiAlH ₄ -Promoted Tandem Reduction/Oxidation of Fluorenyl Derivatives under Air. <i>European Journal of Organic Chemistry</i> , 2014, 2014, 6538-6546. | 2.4 | 4 |
| 32 | Synthesis, antioxidant and cytoprotective evaluation of potential antiatherogenic phenolic hydrazones. A structure-activity relationship insight. <i>Bioorganic and Medicinal Chemistry</i> , 2014, 22, 4269-4276. | 3.0 | 25 |
| 33 | Antimalarial Bicyclic Peroxides Belonging to the G-Factor Family: Mechanistic Aspects of their Formation and Iron (II) Induced Reduction. <i>Current Topics in Medicinal Chemistry</i> , 2014, 14, 1668-1683. | 2.1 | 5 |
| 34 | Synthesis and evaluation of 1,2-ketotriazoles and 1,2-diketotriazoles as inhibitors of Mycobacterium tuberculosis. <i>European Journal of Medicinal Chemistry</i> , 2013, 69, 167-173. | 5.5 | 35 |
| 35 | Design, chemical synthesis of 3-(9H-fluoren-9-yl)pyrrolidine-2,5-dione derivatives and biological activity against enoyl-ACP reductase (InhA) and Mycobacterium tuberculosis. <i>European Journal of Medicinal Chemistry</i> , 2013, 70, 37-48. | 5.5 | 39 |
| 36 | Cinnamic Acid Derivatives in Tuberculosis, Malaria and Cardiovascular Diseases - A Review. <i>Current Organic Chemistry</i> , 2012, 16, 747-768. | 1.6 | 32 |

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|----|---|-----|-----------|
| 37 | Cinnamic Derivatives in Tuberculosis. , 2012, , . | | 6 |
| 38 | Recent advances in the development of cinnamic-like derivatives as antituberculosis agents. Expert Opinion on Therapeutic Patents, 2012, 22, 155-168. | 5.0 | 21 |
| 39 | Diaryl ether derivatives as anticancer agents "a review. MedChemComm, 2012, 3, 1356. | 3.4 | 59 |
| 40 | Chemical synthesis and biological evaluation of triazole derivatives as inhibitors of InhA and antituberculosis agents. European Journal of Medicinal Chemistry, 2012, 52, 275-283. | 5.5 | 81 |
| 41 | Synthesis of 1,2-diketotriazoles by Aerobic Copper-Catalyzed Oxygenation with Triazole as an Intramolecular Assisting Group. European Journal of Organic Chemistry, 2012, 2012, 409-416. | 2.4 | 25 |
| 42 | Antiatherogenic Effect of Bisvanillyl-Hydralazone, a New Hydralazine Derivative with Antioxidant, Carbonyl Scavenger, and Antiapoptotic Properties. Antioxidants and Redox Signaling, 2011, 14, 2093-2106. | 5.4 | 23 |
| 43 | Design, Synthesis, and Biological Evaluation of New Cinnamic Derivatives as Antituberculosis Agents. Journal of Medicinal Chemistry, 2011, 54, 1449-1461. | 6.4 | 100 |
| 44 | Synthesis and biological activities of triazole derivatives as inhibitors of InhA and antituberculosis agents. European Journal of Medicinal Chemistry, 2011, 46, 5524-5531. | 5.5 | 84 |
| 45 | Synthesis and anticancer activity evaluation of 2-(4-alkoxyphenyl)cyclopropyl hydrazides and triazolo phthalazines. Bioorganic and Medicinal Chemistry, 2010, 18, 2537-2548. | 3.0 | 34 |
| 46 | Revisiting the aldol reaction of cis-1,2-epoxyaldehyde promoted by BF ₃ ·Et ₂ O: direct access to 2-deoxy-2-fluoro heptulosonic ester analogues. Carbohydrate Research, 2010, 345, 2421-2426. | 2.3 | 3 |
| 47 | Synthesis and antioxidant activity evaluation of a syringic hydrazones family. European Journal of Medicinal Chemistry, 2010, 45, 3019-3026. | 5.5 | 116 |
| 48 | Synthesis and evaluation of a novel series of pseudo-cinnamic derivatives as antituberculosis agents. Bioorganic and Medicinal Chemistry Letters, 2009, 19, 341-343. | 2.2 | 48 |
| 49 | Carbonyl scavenger and antiatherogenic effects of hydrazine derivatives. Free Radical Biology and Medicine, 2008, 45, 1457-1467. | 2.9 | 92 |
| 50 | Development of Novel Antiatherogenic Biaryls: Design, Synthesis, and Reactivity. Journal of Medicinal Chemistry, 2008, 51, 3171-3181. | 6.4 | 58 |
| 51 | New approach to carbamoyl-polyoxamic acid derivatives through an oxazolidinone synthon. Tetrahedron: Asymmetry, 2007, 18, 1320-1329. | 1.8 | 7 |
| 52 | Synthesis of ferulic ester dimers, functionalisation and biological evaluation as potential antiatherogenic and antiplasmodial agents. Bioorganic and Medicinal Chemistry, 2007, 15, 6018-6026. | 3.0 | 26 |
| 53 | Mukaiyama aldolisation reactions of 1,2-epoxyaldehydes in aqueous media. Tetrahedron, 2005, 61, 8895-8903. | 1.9 | 10 |
| 54 | Design, Synthesis, and Evaluation of Pharmacological Properties of Cinnamic Derivatives as Antiatherogenic Agents. Journal of Medicinal Chemistry, 2005, 48, 8115-8124. | 6.4 | 37 |

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|----|---|-----|-----------|
| 55 | Asymmetric syntheses of (â€“)lentiginosine and an original pyrrolizidinic analogue thereof from a versatile epoxyamine intermediate. <i>Organic and Biomolecular Chemistry</i> , 2005, 3, 2626. | 2.8 | 41 |
| 56 | Crucial role of the peroxyketal function for antimalarial activity in the G-factor series. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2004, 14, 1433-1436. | 2.2 | 24 |
| 57 | Stereoselective Access to the Versatile 4-Aminohex-5-ene-1,2,3-triol Pattern. <i>Journal of Organic Chemistry</i> , 2004, 69, 8775-8779. | 3.2 | 10 |
| 58 | A Flexible Route Towards Five-Membered Ring Imino Sugars and Their Novel 2-Deoxy-2-fluoro Analogues. <i>European Journal of Organic Chemistry</i> , 2003, 2003, 2903-2910. | 2.4 | 25 |
| 59 | Direct Access to Furanosidic Eight-Membered Ulosonic Esters from cis-Î±,Î²-Epoxy Aldehydes. <i>European Journal of Organic Chemistry</i> , 2003, 2003, 672-688. | 2.4 | 26 |
| 60 | Addition of Lithium Ethyl Fluoroacetate to cis and trans Î±,Î²-Epoxyaldehydes. Access to C2 Fluorinated Butyrolactones.. <i>ChemInform</i> , 2003, 34, no. | 0.0 | 0 |
| 61 | Addition of lithium ethyl fluoroacetate to cis and trans Î±,Î²-epoxyaldehydes. Access to C2 fluorinated butyrolactones. <i>Tetrahedron Letters</i> , 2003, 44, 1891-1894. | 1.4 | 4 |
| 62 | Synthesis of phosphonocinnamic thioesters, substrate analogues of cinnamoyl-CoA reductase, a key enzyme in the lignification process. <i>Tetrahedron Letters</i> , 2003, 44, 2445-2447. | 1.4 | 8 |
| 63 | Concise asymmetric syntheses of (â€“)lentiginosine and of its pyrrolizidinic analogue. <i>Chemical Communications</i> , 2003, , 582-583. | 4.1 | 30 |
| 64 | Stereoselective Preparation of Protected Thymine Polyoxin C and Approaches Towards Synthesis of Its C2-Modified Analogues. <i>European Journal of Organic Chemistry</i> , 2001, 2001, 1105-1113. | 2.4 | 15 |
| 65 | Stereochemistry Control in the Lewis Acid Mediated Lactonization Reaction of Î±,Î²-Epoxy-Î²-silyloxy Esters. <i>European Journal of Organic Chemistry</i> , 2001, 2001, 4247. | 2.4 | 6 |
| 66 | De Novo Asymmetric Synthesis of Protected 5-O-Carbamoylpolyoxamic Acid. <i>Synthesis</i> , 2000, 2000, 1409-1414. | 2.3 | 8 |
| 67 | Boron trifluoride as a promoter and fluoride donor in the aldol reaction of trans Î±,Î²-epoxyaldehydes. Access to 5- and 6-fluoro heptulosonic ester analogues. <i>Tetrahedron Letters</i> , 1999, 40, 7323-7327. | 1.4 | 9 |
| 68 | Lactonisation and lactone ether formation of nerol geraniol compounds. Use of ¹³ C to identify the cyclisation process. <i>Tetrahedron</i> , 1999, 55, 5129-5138. | 1.9 | 12 |
| 69 | Attempt to rationalize the diastereoselectivity in the addition of ester enolate to optically active Î±,Î²-epoxyaldehydes. <i>Tetrahedron</i> , 1999, 55, 14013-14030. | 1.9 | 33 |
| 70 | Influence of the nature and substitution of chiral 2,3-epoxy alcohol derivatives on the enantiomeric elution order on chiralcel OD column. <i>Chirality</i> , 1998, 10, 804-807. | 2.6 | 4 |
| 71 | Total Synthesis of a Thymidine 2-Deoxypolyoxin C Analogue. <i>Journal of Organic Chemistry</i> , 1998, 63, 2601-2608. | 3.2 | 24 |
| 72 | Stereoselective synthesis of five and/or six membered ring hydroxylactones obtained by Lewis acid mediated reaction of Î±,Î²-epoxy-Î²-hydroxyesters; access to 5-methylated 2-deoxysugars.. <i>Tetrahedron</i> , 1997, 53, 659-672. | 1.9 | 23 |

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|----|--|-----|-----------|
| 73 | Enhanced diastereoselectivity in the addition of ester enolate to optically active $\hat{1}\pm, \hat{1}^2$ -epoxyaldehydes obtained from nerol and geraniol. <i>Tetrahedron</i> , 1996, 52, 9047-9056. | 1.9 | 24 |
| 74 | Synthesis of a 3-Deoxy-D-arabino-2-heptulosonic Acid Derivative. <i>Journal of Organic Chemistry</i> , 1995, 60, 7343-7347. | 3.2 | 20 |
| 75 | Diastereoface differentiation in addition of lithium enolates to chiral $\hat{1}\pm, \hat{1}^2$ -epoxyaldehydes. <i>Tetrahedron</i> , 1993, 49, 5253-5266. | 1.9 | 32 |
| 76 | A short synthesis of substituted $\hat{1}^2$ -hydroxy $\hat{1}^3$ -butyrolactones and 2-deoxyhexofuranosides. <i>Tetrahedron Letters</i> , 1992, 33, 1439-1442. | 1.4 | 17 |
| 77 | Diastereoselection in the addition of enolates to chiral $\hat{1}\pm, \hat{1}^2$ -epoxyaldehydes. <i>Tetrahedron Letters</i> , 1991, 32, 5345-5348. | 1.4 | 25 |