Pascal Pigeon

List of Publications by Year in descending order

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86 2,948 34 52
papers citations h-index g-index

95 95 95 2347 all docs docs citations times ranked citing authors

| # | Article | IF | CITATIONS |
|----|---|-----|-----------|
| 1 | Diversity-oriented synthesis and bioactivity evaluation of N-substituted ferrocifen compounds as novel antiproliferative agents against TNBC cancer cells. European Journal of Medicinal Chemistry, 2022, 234, 114202. | 2.6 | 8 |
| 2 | \hat{l}_{\pm} -Hydroxylactams as Efficient Entries to Diversely Functionalized Ferrociphenols: Synthesis and Antiproliferative Activity Studies. Molecules, 2022, 27, 4549. | 1.7 | 3 |
| 3 | Heterogeneity of Response to Iron-Based Metallodrugs in Glioblastoma Is Associated with Differences in Chemical Structures and Driven by FAS Expression Dynamics and Transcriptomic Subtypes. International Journal of Molecular Sciences, 2021, 22, 10404. | 1.8 | 11 |
| 4 | Antimicrobial, Antitumor and Side Effects Assessment of a Newly Synthesized Tamoxifen Analog. Current Topics in Medicinal Chemistry, 2020, 20, 2281-2288. | 1.0 | 4 |
| 5 | Importance of Combining Advanced Particle Size Analysis Techniques To Characterize Cell-Penetrating Peptide–Ferrocifen Self-Assemblies. Journal of Physical Chemistry Letters, 2019, 10, 6613-6620. | 2.1 | 7 |
| 6 | Small Structural Differences between Two Ferrocenyl Diphenols Determine Large Discrepancies of Reactivity and Biological Effects. ChemMedChem, 2019, 14, 1717-1726. | 1.6 | 17 |
| 7 | Atypical Lone Pair–π Interaction with Quinone Methides in a Series of Imidoâ€Ferrociphenol Anticancer Drug Candidates. Angewandte Chemie, 2019, 131, 8509-8513. | 1.6 | 6 |
| 8 | Atypical Lone Pair–π Interaction with Quinone Methides in a Series of Imidoâ€Ferrociphenol Anticancer Drug Candidates. Angewandte Chemie - International Edition, 2019, 58, 8421-8425. | 7.2 | 30 |
| 9 | Selective cytotoxicity of arene tricarbonylchromium towards tumour cell lines. Journal of Organometallic Chemistry, 2018, 862, 7-12. | 0.8 | 5 |
| 10 | A new generation of ferrociphenols leads to a great diversity of reactive metabolites, and exhibits remarkable antiproliferative properties. Chemical Science, 2018, 9, 70-78. | 3.7 | 44 |
| 11 | Anticancer properties of lipid and poly($\hat{l}\mu$ -caprolactone) nanocapsules loaded with ferrocenyl-tamoxifen derivatives. Journal of Pharmacy and Pharmacology, 2018, 70, 1474-1484. | 1.2 | 8 |
| 12 | Enhanced and preferential internalization of lipid nanocapsules into human glioblastoma cells: effect of a surface-functionalizing NFL peptide. Nanoscale, 2018, 10, 13485-13501. | 2.8 | 26 |
| 13 | Aryl Butenes Active against K562 Cells and Lacking Tyrosinase Inhibitory Activity as New Leads in the Treatment of Leukemia. Mini-Reviews in Medicinal Chemistry, 2018, 18, 1294-1301. | 1.1 | 2 |
| 14 | Tamoxifen-like metallocifens target the thioredoxin system determining mitochondrial impairment leading to apoptosis in Jurkat cells. Metallomics, 2017, 9, 949-959. | 1.0 | 30 |
| 15 | A New Series of Succinimido-ferrociphenols and Related Heterocyclic Species Induce Strong Antiproliferative Effects, Especially against Ovarian Cancer Cells Resistant to Cisplatin. Journal of Medicinal Chemistry, 2017, 60, 8358-8368. | 2.9 | 40 |
| 16 | The inhibition of tyrosinase by some aryl butenes: A desired activity or a side effect to avoid. Journal of Organometallic Chemistry, 2017, 848, 133-141. | 0.8 | 4 |
| 17 | Side-Chain Effects on the 1-(Bis-aryl-methylidene)-[3] ferrocenophane Skeleton: Antiproliferative Activity against TNBC Cancer Cells and Comparison with the Acyclic Ferrocifen Series. European Journal of Inorganic Chemistry, 2017, 2017, 454-465. | 1.0 | 6 |
| 18 | Synthesis and antiproliferative evaluation of novel hydroxypropyl-ferrociphenol derivatives, resulting from the modification of hydroxyl groups. Journal of Organometallic Chemistry, 2017, 829, 108-115. | 0.8 | 11 |

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| 19 | Ferrocenyl Quinone Methide–Thiol Adducts as New Antiproliferative Agents: Synthesis, Metabolic Formation from Ferrociphenols, and Oxidative Transformation. Angewandte Chemie, 2016, 128, 10587-10590. | 1.6 | 10 |
| 20 | Ferrocenyl Quinone Methide–Thiol Adducts as New Antiproliferative Agents: Synthesis, Metabolic Formation from Ferrociphenols, and Oxidative Transformation. Angewandte Chemie - International Edition, 2016, 55, 10431-10434. | 7.2 | 33 |
| 21 | Enzymatic oxidation of ansa-ferrocifen leads to strong and selective thioredoxin reductase inhibition in vitro. Journal of Inorganic Biochemistry, 2016, 165, 146-151. | 1.5 | 19 |
| 22 | The length of the bridging chain in ansa-metallocenes influences their antiproliferative activity against triple negative breast cancer cells (TNBC). Dalton Transactions, 2016, 45, 13126-13134. | 1.6 | 8 |
| 23 | Efficacy of a novel ferrocenyl diaryl butene citrate compound as a biocide for preventing healthcare-associated infections. MedChemComm, 2016, 7, 948-954. | 3.5 | 2 |
| 24 | Organometallic Antitumor Compounds: Ferrocifens as Precursors to Quinone Methides. Angewandte Chemie - International Edition, 2015, 54, 10230-10233. | 7.2 | 68 |
| 25 | Oxidative Metabolism of Ferrocene Analogues of Tamoxifen: Characterization and Antiproliferative Activities of the Metabolites. ChemMedChem, 2015, 10, 981-990. | 1.6 | 33 |
| 26 | Antiplasmodial activity of iron(II) and ruthenium(II) organometallic complexes against Plasmodium falciparum blood parasites. Memorias Do Instituto Oswaldo Cruz, 2015, 110, 981-988. | 0.8 | 12 |
| 27 | Phthalimido–ferrocidiphenol cyclodextrin complexes: Characterization and anticancer activity. International Journal of Pharmaceutics, 2015, 491, 323-334. | 2.6 | 14 |
| 28 | Antibacterial properties and mode of action of new triaryl butene citrate compounds. European Journal of Medicinal Chemistry, 2014, 76, 408-413. | 2.6 | 10 |
| 29 | Evidence for Targeting Thioredoxin Reductases with Ferrocenyl Quinone Methides. A Possible Molecular Basis for the Antiproliferative Effect of Hydroxyferrocifens on Cancer Cells. Journal of Medicinal Chemistry, 2014, 57, 8849-8859. | 2.9 | 102 |
| 30 | Ferrocifen derivatives that induce senescence in cancer cells: selected examples. Journal of Inorganic Biochemistry, 2014, 141, 144-151. | 1.5 | 56 |
| 31 | Oxidative Sequence of a Ruthenocene-Based Anticancer Drug Candidate in a Basic Environment. Organometallics, 2014, 33, 4940-4946. | 1.1 | 18 |
| 32 | Molecular Mechanism of Action of 2â€Ferrocenylâ€1,1â€diphenylbutâ€1â€ene on HLâ€60 Leukemia Cells. ChemMedChem, 2014, 9, 2580-2586. | 1.6 | 14 |
| 33 | Atypical McMurry Cross-Coupling Reactions Leading to a New Series of Potent Antiproliferative Compounds Bearing the Key [Ferrocenyl-Ene-Phenol] Motif. Molecules, 2014, 19, 10350-10369. | 1.7 | 18 |
| 34 | The inÂvivo performance of ferrocenyl tamoxifen lipid nanocapsules in xenografted triple negative breast cancer. Biomaterials, 2013, 34, 6949-6956. | 5.7 | 43 |
| 35 | Ferrocenyl flavonoid-induced morphological modifications of endothelial cells and cytotoxicity against B16 murine melanoma cells. Journal of Organometallic Chemistry, 2013, 734, 78-85. | 0.8 | 28 |
| 36 | Effect of the amino chain length and the transformation into citric acid salts of aryl-diphenyl-butenes and ferrocenyl-diphenyl-butenes bearing two dimethylaminoalkyl chains on their antimicrobial activities. SpringerPlus, 2013, 2, 508. | 1.2 | 4 |

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| 37 | Selection of a suitable disc bioassay for the screening of anti-tumor molecules. International Journal of Biomedical Science, 2013, 9, 230-6. | 0.5 | 3 |
| 38 | Ferrocenyl catechols: synthesis, oxidation chemistry and anti-proliferative effects on MDA-MB-231 breast cancer cells. Dalton Transactions, 2012, 41, 7537. | 1.6 | 45 |
| 39 | Synthesis and Antiproliferative Effects of [3]Ferrocenophane Transposition Products and Pinacols Obtained from McMurry Cross-Coupling Reactions. Organometallics, 2012, 31, 5856-5866. | 1.1 | 20 |
| 40 | A new series of ferrocifen derivatives, bearing two aminoalkyl chains, with strong antiproliferative effects on breast cancer cells. New Journal of Chemistry, 2011, 35, 2212. | 1.4 | 38 |
| 41 | Biological evaluation of twenty-eight ferrocenyl tetrasubstituted olefins: Cancer cell growth inhibition, ROS production and hemolytic activity. European Journal of Medicinal Chemistry, 2011, 46, 3778-3787. | 2.6 | 38 |
| 42 | Evaluation of bactericidal and fungicidal activity of ferrocenyl or phenyl derivatives in the diphenyl butene series. Journal of Organometallic Chemistry, 2011, 696, 1038-1048. | 0.8 | 45 |
| 43 | Antiparasitic and immunomodulatory activities of 1,1â€bis(4â€hydroxyphenyl)â€2â€phenylâ€butâ€1â€ene and its protected and free 2â€ferrocenyl derivatives. Drug Development Research, 2010, 71, 69-75. | ⁵ 1.4 | 6 |
| 44 | Synthesis, Cytotoxicity, and COMPARE Analysis of Ferrocene and [3]Ferrocenophane Tetrasubstituted Olefin Derivatives against Human Cancer Cells. ChemMedChem, 2010, 5, 2039-2050. | 1.6 | 76 |
| 45 | Comparative toxicity of [3]ferrocenophane and ferrocene moieties on breast cancer cells. Tetrahedron Letters, 2010, 51, 118-120. | 0.7 | 54 |
| 46 | Facile synthesis and strong antiproliferative activity of disubstituted diphenylmethylidenyl-[3]ferrocenophanes on breast and prostate cancer cell lines. MedChemComm, 2010, 1, 149. | 3.5 | 36 |
| 47 | Synthesis and Structure–Activity Relationships of Ferrocenyl Tamoxifen Derivatives with Modified Side Chains. Chemistry - A European Journal, 2009, 15, 684-696. | 1.7 | 58 |
| 48 | Dose effect activity of ferrocifen-loaded lipid nanocapsules on a 9L-glioma model. International Journal of Pharmaceutics, 2009, 379, 317-323. | 2.6 | 55 |
| 49 | The replacement of a phenol group by an aniline or acetanilide group enhances the cytotoxicity of 2-ferrocenyl-1,1-diphenyl-but-l-ene compounds against breast cancer cells. Journal of Organometallic Chemistry, 2009, 694, 895-901. | 0.8 | 65 |
| 50 | Synthesis, oxidation chemistry and cytotoxicity studies on ferrocene derivatives of diethylstilbestrol. Dalton Transactions, 2009, , 10871. | 1.6 | 36 |
| 51 | A [3]Ferrocenophane Polyphenol Showing a Remarkable Antiproliferative Activity on Breast and Prostate Cancer Cell Lines. Journal of Medicinal Chemistry, 2009, 52, 4964-4967. | 2.9 | 125 |
| 52 | Role of aromatic substituents on the antiproliferative effects of diphenyl ferrocenyl butene compounds. Dalton Transactions, 2009, , 4318. | 1.6 | 28 |
| 53 | Ferrocenyl compounds possessing protected phenol and thiophenol groups: Synthesis, X-ray structure, and in vitro biological effects against breast cancer. Journal of Organometallic Chemistry, 2008, 693, 1716-1722. | 0.8 | 40 |
| 54 | Electrochemical attachment of a conjugated amino–ferrocifen complex onto carbon and metal surfaces. Journal of Electroanalytical Chemistry, 2008, 619-620, 169-175. | 1.9 | 43 |

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| 55 | Nanoparticles loaded with ferrocenyl tamoxifen derivatives for breast cancer treatment. International Journal of Pharmaceutics, 2008, 347, 128-135. | 2.6 | 61 |
| 56 | Lipid nanocapsules loaded with an organometallic tamoxifen derivative as a novel drug-carrier system for experimental malignant gliomas. Journal of Controlled Release, 2008, 130, 146-153. | 4.8 | 113 |
| 57 | Ferrocifens and Ferrocifenols as New Potential Weapons against Breast Cancer. Chimia, 2007, 61, 716. | 0.3 | 152 |
| 58 | The influence of phenolic hydroxy substitution on the electron transfer and anti-cancer properties of compounds based on the 2-ferrocenyl-1-phenyl-but-1-ene motif. Dalton Transactions, 2007, , 5073. | 1.6 | 83 |
| 59 | Organometallic diphenols: The importance of the organometallic moiety on the expression of a cytotoxic effect on breast cancer cells. Journal of Organometallic Chemistry, 2007, 692, 1315-1326. | 0.8 | 66 |
| 60 | Organometallic analogues of tamoxifen: Effect of the amino side-chain replacement by a carbonyl ferrocenyl moiety in hydroxytamoxifen. Journal of Organometallic Chemistry, 2007, 692, 1219-1225. | 0.8 | 46 |
| 61 | Modification of the Estrogenic Properties of Diphenols by the Incorporation of Ferrocene. Generation of Antiproliferative Effects in Vitro. Journal of Medicinal Chemistry, 2005, 48, 3937-3940. | 2.9 | 200 |
| 62 | Selective Estrogen Receptor Modulators in the Ruthenocene Series. Synthesis and Biological Behavior. Journal of Medicinal Chemistry, 2005, 48, 2814-2821. | 2.9 | 109 |
| 63 | Selective Estrogen-Receptor Modulators (SERMs) in the Cyclopentadienylrhenium Tricarbonyl Series: Synthesis and Biological Behaviour. ChemBioChem, 2004, 5, 1104-1113. | 1.3 | 66 |
| 64 | A short route to cyclopentadienyltricarbonylrhenium substituted derivatives. Journal of Organometallic Chemistry, 2003, 668, 140-144. | 0.8 | 10 |
| 65 | Intramolecular Addition of a Hydroxyl to an N-Acyliminium System. Application to the Synthesis of Isoindolo[2,1-a][3,1]benzoxazine and Isoindolo[1,2-c][2,4]benzoxazepine Derivatives. Heterocycles, 2002, 56, 129. | 0.4 | 13 |
| 66 | First anti-oestrogen in the cyclopentadienyl rhenium tricarbonyl series. Synthesis and study of antiproliferative effects. Chemical Communications, 2001, , 383-384. | 2.2 | 67 |
| 67 | Study of a 1,6-hydride shift in an open chain of hydroxylactam-triarylcarbinols. Tetrahedron, 2001, 57, 4939-4943. | 1.0 | 8 |
| 68 | Thieno[2′,3′:5,6]azepino[2,1â€∢i>a]isoindolones from hydroxylactamâ€alcohols <i>via N</i> â€acylimin ion olefin cyclization. Journal of Heterocyclic Chemistry, 2001, 38, 35-39. | ium 1.4 | 3 |
| 69 | Quinoxalines, Bezodiazepines and Bezodiazocines Fused to Pyrrole and Isoindole via N-Acyliminium Ion Aromatic Cyclization. Heterocycles, 2000, 52, 273. | 0.4 | 9 |
| 70 | Acyliminium ionâ€olefin cyclization leading to isoindolo[2,1â€ <i>a</i>)]quinoline derivatives. Journal of Heterocyclic Chemistry, 1999, 36, 691-695. | 1.4 | 15 |
| 71 | Polycyclic systems: Synthesis of isoindolo[2,1â€ <i>b</i>]â€pyrrolo[1,2â€ <i>d</i>][2,4]benzodiazocine and isoindoloâ€{1,2â€ <i>d</i>][1,5]benzodiazocine. Journal of Heterocyclic Chemistry, 1999, 36, 735-738. | 1.4 | 6 |
| 72 | Selective access to N-aryl or N-alkyl derivatives of isoindolo [2,1-b] [2,4] benzo (or thieno) diazepines. Tetrahedron, 1998, 54, 1497-1506. | 1.0 | 20 |

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| 73 | Diisoindolothieno[2,4]diazepines via a diastereoselective N-acyliminium ion cyclization. Tetrahedron Letters, 1998, 39, 8659-8662. | 0.7 | 9 |
| 74 | Synthesis of benzo(or furo)[5,6]azepino[2,1-a]isoindolone derivatives: π-cyclisations of N-acyliminium ions. Tetrahedron Letters, 1998, 39, 9187-9190. | 0.7 | 36 |
| 75 | Synthesis and reduction of thieno[2′,3′(3′,2′ or 3′,4′):5,6]-azocino[2,1-a]isoindole-7, 13-diones. Heterocyclic Chemistry, 1998, 35, 1429-1433. | Journal of | f ₂ |
| 76 | New fused lactones from indolizinediones via N-acyliminium ions. Tetrahedron, 1998, 54, 8737-8744. | 1.0 | 20 |
| 77 | Novel Approach to Isoindolo[2,1-a]quinolines. Synthetic Communications, 1998, 28, 2507-2516. | 1.1 | 18 |
| 78 | Introduction of a Carboxymethylamino(or oxy Or thio) Group in the 3 Position of 2-Aryl(or) Tj ETQq0 0 0 rgBT /Ov | erlock 10 | Tf 50 542 Td |
| 79 | Acyliminium ion cyclizations: Synthesis of thieno[2′,3′:3,4]pyrrolo[2,1-a] isoindolone and benzo[a]thieno[2,3(3,2 or 3,4)-g]indolizinones. Tetrahedron, 1997, 53, 2495-2504. | 1.0 | 43 |
| 80 | Synthesis of dibenz[c,e]azepine and benzo[e]thieno[c]azepine via, N-acyliminium ion cyclization. Tetrahedron Letters, 1997, 38, 1041-1042. | 0.7 | 25 |
| 81 | A New Access to Isoindolo[2,1-b][2,4]benzodiazepines through an N-Acyliminium Ion - Amide Cyclization. Tetrahedron Letters, 1997, 38, 2985-2988. | 0.7 | 45 |
| 82 | Tetracyclic systems: Synthesis of isoindolo[1,2â€ <i>b</i>]thienoâ€[2,3(3,2 or 3,4)â€ <i>e</i>][1,3]thiazocines and Isoindolo[2,1â€ <i>a</i>]thienoâ€[2,3(3,2 or 3,4)â€ <i>f</i>][1,4] and [1,5]diazocines. Journal of Heterocyclic Chemistry, 1997, 34, 375-380. | 1.4 | 12 |
| 83 | Intramolecular amidoalkylation cyclizations in synthesis of novel pyrrolo(or) Tj ETQq1 1 0.784314 rgBT /Overlock | 10 Tf 50 3 | 342 Td (isc <mark>in</mark> |
| 84 | Synthesis of thieno[2′,3′(3′,4′ or 3′,2′):5,6]azepino[2,1â€ <i>a</i>]isoindolediones from <i>N</i> â€Thienyl′(3)â€ylmethylphthalimides. Journal of Heterocyclic Chemistry, 1996, 33, 129-135. | 1.4 | 21 |
| 85 | Benzothienoindolizidines via intramolecular aryl radical cyclization or palladium catalyzed cyclization. Tetrahedron Letters, 1996, 37, 7707-7710. | 0.7 | 61 |
| 86 | Inhibition of Cathepsin B by Ferrocenyl Indenes Highlights a new Pharmacological Facet of Ferrocifens. European Journal of Inorganic Chemistry, 0, , . | 1.0 | 3 |