## Malin Wickström

List of Publications by Year in descending order

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| #  | Article  | IF   | CITATIONS |
|----|--|------|-----------|
| 1  | Aminopeptidase N (CD13) as a target for cancer chemotherapy. Cancer Science, 2011, 102, 501-508.   | 3.9  | 293       |
| 2  | $Wnt/\hat{l}^2$ -catenin pathway regulates MGMT gene expression in cancer and inhibition of Wnt signalling prevents chemoresistance. Nature Communications, 2015, 6, 8904.   | 12.8 | 177       |
| 3  | Neuroblastoma—A Neural Crest Derived Embryonal Malignancy. Frontiers in Molecular Neuroscience,<br>2019, 12, 9.  | 2.9  | 166       |
| 4  | Pharmacological profiling of disulfiram using human tumor cell lines and human tumor cells from patients. Biochemical Pharmacology, 2007, 73, 25-33.   | 4.4  | 99        |
| 5  | Targeting the hedgehog signal transduction pathway at the level of GLI inhibits neuroblastoma cell growth <i>in vitro</i> and <i>in vivo</i> . International Journal of Cancer, 2013, 132, 1516-1524.              | 5.1  | 99        |
| 6  | Cytotoxic Effects of Cardiac Glycosides in Colon Cancer Cells, Alone and in Combination with Standard Chemotherapeutic Drugs. Journal of Natural Products, 2009, 72, 1969-1974.                                    | 3.0  | 91        |
| 7  | Melflufen - a peptidase-potentiated alkylating agent in clinical trials. Oncotarget, 2017, 8, 66641-66655.   | 1.8  | 65        |
| 8  | Inhibitors and promoters of tubulin polymerization: Synthesis and biological evaluation of chalcones and related dienones as potential anticancer agents. Bioorganic and Medicinal Chemistry, 2011, 19, 2659-2665. | 3.0  | 61        |
| 9  | Celecoxib Prevents Neuroblastoma Tumor Development and Potentiates the Effect of<br>Chemotherapeutic Drugs In vitro and In vivo. Clinical Cancer Research, 2007, 13, 1036-1044.                                    | 7.0  | 56        |
| 10 | Effects of small molecule inhibitors of PI3K/Akt/mTOR signaling on neuroblastoma growth <i>in vitro</i> and <i>in vivo</i> . International Journal of Cancer, 2011, 129, 2958-2965.                                | 5.1  | 54        |
| 11 | Molecular mechanisms and therapeutic targets in neuroblastoma. Pharmacological Research, 2018, 131, 164-176.   | 7.1  | 53        |
| 12 | Rho-associated kinase is a therapeutic target in neuroblastoma. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, E6603-E6612.   | 7.1  | 52        |
| 13 | The alkylating prodrug J1 can be activated by aminopeptidase N, leading to a possible target directed release of melphalan. Biochemical Pharmacology, 2010, 79, 1281-1290.   | 4.4  | 48        |
| 14 | Image-Based Screening for the Identification of Novel Proteasome Inhibitors. Journal of Biomolecular Screening, 2007, 12, 203-210.   | 2.6  | 43        |
| 15 | Activity of Hydrolytic Enzymes in Tumour Cells is a Determinant for Anti-tumour Efficacy of the Melphalan Containing ProdrugJ1. Journal of Drug Targeting, 2003, 11, 355-363.                                      | 4.4  | 41        |
| 16 | Protective Role of Humanin on Bortezomib-Induced Bone Growth Impairment in Anticancer Treatment.<br>Journal of the National Cancer Institute, 2014, 106, djt459.   | 6.3  | 41        |
| 17 | Inhibition of chemerin/CMKLR1 axis in neuroblastoma cells reduces clonogenicity and cell viability <i>in vitro</i> and impairs tumor growth <i>in vivo</i> . Oncotarget, 2017, 8, 95135-95151.                     | 1.8  | 40        |
| 18 | The novel melphalan prodrug J1 inhibits neuroblastoma growth in vitro and in vivo. Molecular Cancer Therapeutics, 2007, 6, 2409-2417.  | 4.1  | 38        |

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|----|---|-----|-----------|
| 19 | The novel alkylating prodrug J1: diagnosis directed activity profile ex vivo and combination analyses in vitro. Investigational New Drugs, 2008, 26, 195-204.   | 2.6 | 36        |
| 20 | Simvastatin induces apoptosis in human breast cancer cells in a NFκB-dependent manner and abolishes the anti-apoptotic signaling of TF/FVIIa and TF/FVIIa/FXa. Thrombosis Research, 2008, 122, 191-202.   | 1.7 | 34        |
| 21 | In vitro evaluation of clinical activity and toxicity of anticancer drugs using tumor cells from<br>patients and cells representing normal tissues. Cancer Chemotherapy and Pharmacology, 2012, 69,<br>697-707.   | 2.3 | 33        |
| 22 | The novel alkylating prodrug melflufen (J1) inhibits angiogenesis in vitro and in vivo. Biochemical<br>Pharmacology, 2013, 86, 888-895.   | 4.4 | 33        |
| 23 | Metronomic scheduling of imatinib abrogates clonogenicity of neuroblastoma cells and enhances their susceptibility to selected chemotherapeutic drugs <i>in vitro</i> and <i>in vivo</i> . International Journal of Cancer, 2009, 124, 1227-1234.                 | 5.1 | 30        |
| 24 | Lithium protects hippocampal progenitors, cognitive performance and hypothalamus-pituitary function after irradiation to the juvenile rat brain. Oncotarget, 2017, 8, 34111-34127.  | 1.8 | 27        |
| 25 | Significant cytotoxic activity <i>in vitro</i> of the EGFR tyrosine kinase inhibitor gefitinib in acute<br>myeloblastic leukaemia. European Journal of Haematology, 2008, 81, 344-353.  | 2.2 | 26        |
| 26 | Targeting Fibroblast Growth Factor Receptor (FGFR) and Phosphoinositide 3-kinase (PI3K) Signaling<br>Pathways in Medulloblastoma Cell Lines. Anticancer Research, 2020, 40, 53-66.  | 1.1 | 25        |
| 27 | Synthesis and Characterization of a Multi Ringâ€Fused 2â€Pyridoneâ€Based Fluorescent Scaffold. European<br>Journal of Organic Chemistry, 2010, 2010, 6171-6178.   | 2.4 | 20        |
| 28 | Planar cell polarity gene expression correlates with tumor cell viability and prognostic outcome in neuroblastoma. BMC Cancer, 2016, 16, 259.   | 2.6 | 20        |
| 29 | Phenotype-Based Screening of Mechanistically Annotated Compounds in Combination with Gene<br>Expression and Pathway Analysis Identifies Candidate Drug Targets in a Human Squamous Carcinoma<br>Cell Model. Journal of Biomolecular Screening, 2006, 11, 457-468. | 2.6 | 18        |
| 30 | In vitro and in vivo activity of melflufen (J1) in lymphoma. BMC Cancer, 2016, 16, 263.   | 2.6 | 18        |
| 31 | Phenotype-based drug screening in primary ovarian carcinoma cultures identifies intracellular iron depletion as a promising strategy for cancer treatment. Biochemical Pharmacology, 2011, 82, 139-147.   | 4.4 | 16        |
| 32 | Effects of PI3K and FGFR inhibitors alone and in combination, and with/without cytostatics in childhood neuroblastoma cell lines. International Journal of Oncology, 2021, 58, 211-225.   | 3.3 | 16        |
| 33 | Alternative Cytotoxic Effects of the Postulated IGF-IR Inhibitor Picropodophyllin <i>In Vitro</i> .<br>Molecular Cancer Therapeutics, 2013, 12, 1526-1536.  | 4.1 | 15        |
| 34 | Inhibition of Rho-Associated Kinase Suppresses Medulloblastoma Growth. Cancers, 2020, 12, 73.   | 3.7 | 10        |
| 35 | The impact of S6K1 kinase on neuroblastoma cell proliferation is independent of GLI1 signaling. BMC Cancer, 2014, 14, 600.  | 2.6 | 9         |
| 36 | In�vitro antitumor effects of FGFR and PI3K inhibitors on human papillomavirus positive and negative<br>tonsillar and base of tongue cancer cell lines. Oncology Letters, 2019, 18, 6249-6260.  | 1.8 | 9         |

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| 37 | Screening for phenotype selective activity in multidrug resistant cells identifies a novel tubulin active agent insensitive to common forms of cancer drug resistance. BMC Cancer, 2013, 13, 374. | 2.6  | 7         |
| 38 | Wingless/β-catenin signaling as a modulator of chemoresistance in cancer. Molecular and Cellular<br>Oncology, 2016, 3, e1131356.  | 0.7  | 7         |
| 39 | MYCMI-7: A Small MYC-Binding Compound that Inhibits MYC: MAX Interaction and Tumor Growth in a MYC-Dependent Manner. Cancer Research Communications, 2022, 2, 182-201.                            | 1.7  | 6         |
| 40 | SYK Inhibition Potentiates the Effect of Chemotherapeutic Drugs on Neuroblastoma Cells in Vitro.<br>Cancers, 2019, 11, 202.   | 3.7  | 5         |
| 41 | GIT1 protects against breast cancer growth through negative regulation of Notch. Nature<br>Communications, 2022, 13, 1537.  | 12.8 | 5         |
| 42 | Inhibition of the ubiquitin-proteasome system by an NQO1-activatable compound. Cell Death and Disease, 2021, 12, 914.   | 6.3  | 3         |