Doris M Benbrook

List of Publications by Year in descending order

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159585 102487 4,863 110 30 citations h-index papers

66 g-index 113 113 113 5578 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Guidelines for the use and interpretation of assays for monitoring autophagy (4th) Tj ETQq1 1 0.784314 rgBT /Ov	verlock 10	Tf 50 742 To
2	A new retinoic acid receptor identified from a hepatocellular carcinoma. Nature, 1988, 333, 669-672.	27.8	619
3	Retinoid activation of retinoic acid receptor but not retinoid X receptor is sufficient to rescue lethal defect in retinoic acid synthesis. Proceedings of the National Academy of Sciences of the United States of America, 2003, 100, 7135-7140.	7.1	203
4	Cyclin D1 Degradation Is Sufficient to Induce G1 Cell Cycle Arrest despite Constitutive Expression of Cyclin E2 in Ovarian Cancer Cells. Cancer Research, 2009, 69, 6565-6572.	0.9	164
5	Different binding specificities and transactivation of variant CRE's by CREB complexes. Nucleic Acids Research, 1994, 22, 1463-1469.	14.5	144
6	Therapeutic options for management of endometrial hyperplasia. Journal of Gynecologic Oncology, 2016, 27, e8.	2.2	140
7	Targeting autophagy in cancer management & Samp; ndash; strategies and developments. Cancer Management and Research, 2015, 7, 291.	1.9	96
8	A phase II trial of thalidomide in patients with refractory endometrial cancer and correlation with angiogenesis biomarkers: A Gynecologic Oncology Group study. Gynecologic Oncology, 2007, 105, 508-516.	1.4	90
9	Flex-Hets differentially induce apoptosis in cancer over normal cells by directly targeting mitochondria. Molecular Cancer Therapeutics, 2007, 6, 1814-1822.	4.1	88
10	A phase II trial of brivanib in recurrent or persistent endometrial cancer: An NRG Oncology/Gynecologic Oncology Group Study. Gynecologic Oncology, 2014, 135, 38-43.	1.4	82
11	Randomized phase III trial of tamoxifen versus thalidomide in women with biochemical-recurrent-only epithelial ovarian, fallopian tube or primary peritoneal carcinoma after a complete response to first-line platinum/taxane chemotherapy with an evaluation of serum vascular endothelial growth factor (VEGF): A Gynecologic Oncology Group Study. Gynecologic Oncology, 2010, 119, 444-450.	1.4	72
12	Synthesis of Flexible Sulfur-Containing Heteroarotinoids That Induce Apoptosis and Reactive Oxygen Species with Discrimination between Malignant and Benign Cells. Journal of Medicinal Chemistry, 2004, 47, 999-1007.	6.4	68
13	Epidermal Growth Factor Receptor in Vulvar Malignancies and Its Relationship to Metastasis and Patient Survival. Gynecologic Oncology, 1997, 65, 425-429.	1.4	66
14	Antitumor activity of SS(dsFv)PE38 and SS1(dsFv)PE38, recombinant antimesothelin immunotoxins against human gynecologic cancers grown in organotypic culture in vitro. Clinical Cancer Research, 2002, 8, 3520-6.	7.0	60
15	Biologically Active Heteroarotinoids Exhibiting Anticancer Activity and Decreased Toxicity. Journal of Medicinal Chemistry, 1997, 40, 3567-3583.	6.4	57
16	Loss of natural killer T cells promotes pancreatic cancer in <scp>LSL</scp> â€Kras ^{G12D/+} mice. Immunology, 2017, 152, 36-51.	4.4	57
17	Flexible heteroarotinoids (Flex-Hets) exhibit improved therapeutic ratios as anti-cancer agents over retinoic acid receptor agonists. Investigational New Drugs, 2005, 23, 417-428.	2.6	53
18	Involvement of c-FLIP and survivin down-regulation in flexible heteroarotinoid-induced apoptosis and enhancement of TRAIL-initiated apoptosis in lung cancer cells. Molecular Cancer Therapeutics, 2008, 7, 3556-3565.	4.1	48

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19	A Gynecologic Oncology Group phase II trial of the protein kinase C-beta inhibitor, enzastaurin and evaluation of markers with potential predictive and prognostic value in persistent or recurrent epithelial ovarian and primary peritoneal malignancies. Gynecologic Oncology, 2011, 121, 455-461.	1.4	48
20	Heteroarotinoids Inhibit Head and Neck Cancer Cell Lines in Vitro and in Vivo Through Both RAR and RXR Retinoic Acid Receptors. Journal of Medicinal Chemistry, 1999, 42, 4434-4445.	6.4	46
21	Novel Heteroarotinoids as Potential Antagonists of Mycobacteriumbovis BCG. Journal of Medicinal Chemistry, 2004, 47, 1008-1017.	6.4	45
22	The synthetic heteroarotinoid SHetA2 induces apoptosis in squamous carcinoma cells through a receptor-independent and mitochondria-dependent pathway. Cancer Research, 2003, 63, 3826-32.	0.9	45
23	CAAT/Enhancer Binding Protein Homologous Protein–Dependent Death Receptor 5 Induction Is a Major Component of SHetA2-Induced Apoptosis in Lung Cancer Cells. Cancer Research, 2008, 68, 5335-5344.	0.9	44
24	Measurements of adiposity as clinical biomarkers for first-line bevacizumab-based chemotherapy in epithelial ovarian cancer. Gynecologic Oncology, 2014, 133, 11-15.	1.4	44
25	A phase II trial of thalidomide in patients with refractory leiomyosarcoma of the uterus and correlation with biomarkers of angiogenesis: A gynecologic oncology group study. Gynecologic Oncology, 2007, 106, 596-603.	1.4	39
26	Synthesis, Structureâ^'Activity Relationships, and RARγâ^'Ligand Interactions of Nitrogen Heteroarotinoids. Journal of Medicinal Chemistry, 1999, 42, 3602-3614.	6.4	38
27	The Pro-Survival Function of Akt Kinase can be Overridden or Altered to Contribute to Induction of Apoptosis. Current Cancer Drug Targets, 2011, 11, 586-599.	1.6	38
28	Development of flexible-heteroarotinoids for kidney cancer. Molecular Cancer Therapeutics, 2009, 8, 1227-1238.	4.1	35
29	History of Retinoic Acid Receptors. Sub-Cellular Biochemistry, 2014, 70, 1-20.	2.4	35
30	SHetA2 interference with mortalin binding to p66shc and p53 identified using drug-conjugated magnetic microspheres. Investigational New Drugs, 2014, 32, 412-423.	2.6	33
31	Oral toxicity and pharmacokinetic studies of SHetA2, a new chemopreventive agent, in rats and dogs. Drug and Chemical Toxicology, 2013, 36, 284-295.	2.3	32
32	Flexible heteroarotinoid (Flex-Het) SHetA2 inhibits angiogenesis in vitro and in vivo. Investigational New Drugs, 2009, 27, 304-318.	2.6	29
33	Chemoprevention of Colon and Small Intestinal Tumorigenesis in <i>APCmin/+</i> Mice By SHetA2 (NSC721689) without Toxicity. Cancer Prevention Research, 2013, 6, 908-916.	1.5	27
34	Novel ovarian cancer maintenance therapy targeted at mortalin and mutant p53. International Journal of Cancer, 2020, 147, 1086-1097.	5.1	27
35	High performance liquid chromatographic analysis and preclinical pharmacokinetics of the heteroarotinoid antitumor agent, SHetA2. Cancer Chemotherapy and Pharmacology, 2006, 58, 561-569.	2.3	26
36	Gene Expression Analysis of Biological Systems Driving an Organotypic Model of Endometrial Carcinogenesis and Chemoprevention. Gene Regulation and Systems Biology, 2008, 2, GRSB.S344.	2.3	26

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37	Genotoxicity of the cancer chemopreventive drug candidates CP-31398, SHetA2, and phospho-ibuprofen. Mutation Research - Genetic Toxicology and Environmental Mutagenesis, 2012, 746, 78-88.	1.7	26
38	Metabolism of a sulfur ontaining heteroarotionoid antitumor agent, SHetA2, using liquid chromatography/tandem mass spectrometry. Rapid Communications in Mass Spectrometry, 2008, 22, 3371-3381.	1.5	25
39	Label-Free Real-Time Microarray Imaging of Cancer Protein–Protein Interactions and Their Inhibition by Small Molecules. Analytical Chemistry, 2016, 88, 3130-3135.	6.5	25
40	Potential and mechanism of mebendazole for treatment and maintenance of ovarian cancer. Gynecologic Oncology, 2021, 160, 302-311.	1.4	25
41	Synthesis and Characterization of Heteroarotinoids Demonstrate Structure Specificity Relationships. Journal of Medicinal Chemistry, 1998, 41, 3753-3757.	6.4	22
42	Retinoids and steroids regulate menstrual phase histological features in human endometrial organotypic cultures. Fertility and Sterility, 2002, 78, 596-602.	1.0	21
43	Anti-CD73 and anti-OX40 immunotherapy coupled with a novel biocompatible enzyme prodrug system for the treatment of recurrent, metastatic ovarian cancer. Cancer Letters, 2018, 425, 174-182.	7.2	21
44	Preclinical Efficacy and Involvement of AKT, mTOR, and ERK Kinases in the Mechanism of Sulforaphane against Endometrial Cancer. Cancers, 2020, 12, 1273.	3.7	21
45	Accelerated vascular aging and persistent cognitive impairment in older female breast cancer survivors. GeroScience, 2018, 40, 325-336.	4.6	20
46	A phase II trial of thalidomide in patients with refractory uterine carcinosarcoma and correlation with biomarkers of angiogenesis: A Gynecologic Oncology Group study. Gynecologic Oncology, 2012, 127, 356-361.	1.4	19
47	Refining Retinoids with Heteroatoms. Mini-Reviews in Medicinal Chemistry, 2002, 2, 277-283.	2.4	19
48	Nucleotide sequence of cDNA encoding a novel human thyroid hormone receptor. Nucleic Acids Research, 1987, 15, 9613-9613.	14.5	18
49	Induction of death receptor ligand-mediated apoptosis in epithelial ovarian carcinoma: The search for sensitizing agents. Gynecologic Oncology, 2009, 115, 438-442.	1.4	17
50	Silencing BMI1 radiosensitizes human breast cancer cells by inducing DNA damage and autophagy. Oncology Reports, 2017, 37, 2382-2390.	2.6	17
51	Similarities and Differences of Hsp70, hsc70, Grp78 and Mortalin as Cancer Biomarkers and Drug Targets. Cells, 2021, 10, 2996.	4.1	17
52	Biological Assay for Activity and Molecular Mechanism of Retinoids in Cervical Tumor Cells. Gynecologic Oncology, 1997, 66, 114-121.	1.4	16
53	Chemically induced carcinogenesis in rodent models of aging: assessing organismal resilience to genotoxic stressors in geroscience research. GeroScience, 2019, 41, 209-227.	4.6	16
54	NF- \hat{l}^2B is involved in SHetA2 circumvention of TNF- \hat{l}^\pm resistance, but not induction of intrinsic apoptosis. Anti-Cancer Drugs, 2010, 21, 297-305.	1.4	15

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55	A stratified randomized double-blind phase II trial of celecoxib for treating patients with cervical intraepithelial neoplasia: The potential predictive value of VEGF serum levels: An NRG Oncology/Gynecologic Oncology Group study. Gynecologic Oncology, 2017, 145, 291-297.	1.4	15
56	Patient-Derived Xenografts of High-Grade Serous Ovarian Cancer Subtype as a Powerful Tool in Pre-Clinical Research. Cancers, 2021, 13, 6288.	3.7	15
57	Internal standard-based analysis of microarray data2â€"Analysis of functional associations between HVE-genes. Nucleic Acids Research, 2011, 39, 7881-7899.	14.5	14
58	Optimization of a Vaginal Suppository Formulation to Deliver SHetA2 as a Novel Treatment for Cervical Dysplasia. Journal of Pharmaceutical Sciences, 2018, 107, 638-646.	3.3	14
59	Retinoids Enhance Cisplatin-Based Chemoradiation in Cervical Cancer Cells in Vitro. Gynecologic Oncology, 2002, 85, 223-225.	1.4	13
60	Histopathologic, Genetic and Molecular Characterization of Endometrial Cancer Racial Disparity. Cancers, 2021, 13, 1900.	3.7	13
61	Complexity, Retinoid-Responsive Gene Networks, and Bladder Carcinogenesis. Advances in Experimental Medicine and Biology, 1999, 462, 449-467.	1.6	13
62	Heteroarotinoids with Anti-Cancer Activity Against Ovarian Cancer Cells. Open Medicinal Chemistry Journal, 2007, 1, 11-23.	2.4	13
63	Insulin Exerts Direct Effects on Carcinogenic Transformation of Human Endometrial Organotypic Cultures. Cancer Investigation, 2014, 32, 63-70.	1.3	12
64	Synthesis and evaluation of second generation Flex-Het scaffolds against the human ovarian cancer A2780 cell line. European Journal of Medicinal Chemistry, 2015, 96, 209-217.	5.5	12
65	Activity of oxygen-versus sulfur-containing analogs of the Flex-Het anticancer agent SHetA2. European Journal of Medicinal Chemistry, 2018, 158, 720-732.	5. 5	12
66	Pharmacokinetics and interspecies scaling of a novel, orally-bioavailable anti-cancer drug, SHetA2. PLoS ONE, 2018, 13, e0194046.	2.5	12
67	Utility and Mechanism of SHetA2 and Paclitaxel for Treatment of Endometrial Cancer. Cancers, 2021, 13, 2322.	3.7	11
68	Immunohistochemical analysis of proliferation and differentiation in organotypic cultures of cervical tumor cell lines. Tissue and Cell, 1995, 27, 269-274.	2.2	10
69	An ELISA method for detection of human antibodies to an immunotoxin. Journal of Pharmacological and Toxicological Methods, 2002, 47, 169-175.	0.7	10
70	Development of a dietary formulation of the SHetA2 chemoprevention drug for mice. Investigational New Drugs, 2018, 36, 561-570.	2.6	7
71	Synthesis and biological evaluation of SHetA2 (NSC-721689) analogs against the ovarian cancer cell line A2780. European Journal of Medicinal Chemistry, 2019, 170, 16-27.	5.5	7
72	Complementary Targeting of Rb Phosphorylation and Growth in Cervical Cancer Cell Cultures and a Xenograft Mouse Model by SHetA2 and Palbociclib. Cancers, 2020, 12, 1269.	3.7	7

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73	Sensitization of cervical cancer cell lines to low-dose radiation by retinoic acid does not require functional p53. Gynecologic Oncology, 2005, 97, 142-150.	1.4	6
74	Organotypic cultures represent tumor microenvironment for drug testing. Drug Discovery Today: Disease Models, 2006, 3, 143-148.	1.2	6
75	Selective Growth Inhibition of Cancer Cells by <i>L</i> -Methioninase-Containing Fusion Protein Targeted to the Urokinase Receptor. Pharmacology, 2009, 84, 271-275.	2.2	6
76	Influence of the estrus cycle of the mouse on the disposition of SHetA2 after vaginal administration. European Journal of Pharmaceutics and Biopharmaceutics, 2018, 130, 272-280.	4.3	6
77	Bioanalytical method development and validation of HPLCUV assay for the quantification of SHetA2 in mouse and human plasma: Application to pharmacokinetics study. Journal of Pharmaceutical Synthesisgof& Drug Research, 2017, 6, 2.	1.0	6
78	N-[3,4-Dihydro-4-(acetoxymethyl)-2,2,4-trimethyl-2H-1-benzothiopyran-6-yl]-N′-(4-nitrophenyl)thiourea and N-[3,4-dihydro-4-(hydroxymethyl)-2,2,4-trimethyl-2H-1-benzothiopyran-6-yl]-N′-(4-nitrophenyl)thiourea, a Major Metabolite of	1.6	5
79	N-(3,4-Dihydro-2,2,4,4-tetramethyl-2H-1-benzothiopyran-6-YL)-N′-(4-nitrophenyl)thiourea. Phosphorus, Biotechnology and Biopharmaceuticals: Transforming Proteins and Genes Into Drugs, 2nd Edition. Clinical Infectious Diseases, 2015, 60, 331-332.	5.8	5
80	The pro-inflammatory effect of obesity on high grade serous ovarian cancer. Gynecologic Oncology, 2016, 143, 40-45.	1.4	5
81	Correlation of clinical data with fallopian tube specimen immune cells and tissue culture capacity. Tissue and Cell, 2018, 52, 57-64.	2.2	5
82	Tetrahydroquinoline units in flexible heteroarotinoids (Flex-Hets) convey anti-cancer properties in A2780 ovarian cancer cells. Bioorganic and Medicinal Chemistry, 2020, 28, 115244.	3.0	5
83	Physiologically Based Pharmacokinetic Modeling and Tissue Distribution Characteristics of SHetA2 in Tumor-Bearing Mice. AAPS Journal, 2020, 22, 51.	4.4	5
84	SHetA2 Attack on Mortalin and Colleagues in Cancer Therapy and Prevention. Frontiers in Cell and Developmental Biology, 2022, 10, 848682.	3.7	5
85	The Mechanism of Retinoic Acid Radiosensitization Is Independent of AP-1 Repression in a Cervical Carcinoma Cell Line. Gynecologic Oncology, 1999, 73, 253-256.	1.4	4
86	Sensitivities of Uterine Adenocarcinoma, Mixed Mullerian Tumor (MMT) and Sarcoma Cell Lines to Chemotherapeutic Agents and a Flex-Het Drug. American Journal of Pharmacology and Toxicology, 2006, 1, 83-86.	0.7	4
87	Pharmacokinetics and Pharmacodynamics of Escalating Doses of SHetA2 After Vaginal Administration to Mice. Journal of Pharmaceutical Sciences, 2018, 107, 3179-3186.	3.3	3
88	Anti-Cancer Activities and Interaction of Imiquimod and Flex-Het, SHetA2, in Melanoma and Ovarian Cancer. Journal of Cancer Therapy, 2013, 04, 7-19.	0.4	3
89	Retinoids Chemosensitize Ovarian Cancer Cell Lines to Cisplatin Independent of Nuclear Receptors and p53. American Journal of Pharmacology and Toxicology, 2006, 1, 87-93.	0.7	3
90	Identification of Candidate Biomarker and Drug Targets for Improving Endometrial Cancer Racial Disparities. International Journal of Molecular Sciences, 2022, 23, 7779.	4.1	3

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91	Promise and problems of translational research. Gynecologic Oncology, 2006, 103, 14-17.	1.4	2
92	Vaginal Suppositories Containing SHetA2 to Treat Cervical Dysplasia: Pharmacokinetics of Daily Doses and Preliminary Safety Profile. Journal of Pharmaceutical Sciences, 2020, 109, 2000-2008.	3.3	2
93	The Dawning of the Age of Personalized Medicine in Gynecologic Oncology. Cancers, 2020, 12, 3135.	3.7	2
94	Role of AP-1 Antagonism in Growth Inhibition of Cervical Cancer Cell Lines by Retinoids. American Journal of Pharmacology and Toxicology, 2006, 1 , 40-47.	0.7	2
95	Sera Protein Signatures of Endometrial Cancer Lymph Node Metastases. International Journal of Molecular Sciences, 2022, 23, 3277.	4.1	2
96	Optimization and synthesis of (E)-4-[2-(3,4-dihydro-4,4-dimethyl-2H-1-benzopyran-6-y1)-1-propenyl]benzoic acid-11-[14C]. Journal of Labelled Compounds and Radiopharmaceuticals, 1999, 42, 789-796.	1.0	1
97	Ki-67 Expression in a Cervical Cancer Organotypic Model Correlates with Growth and EGF-R Expression. Journal of Lower Genital Tract Disease, 1999, 3, 111-115.	1.9	0
98	Reticulin Expression Demonstrates Hormonal Responsiveness in a Model of Cycling Human Endometrium. Obstetrics and Gynecology, 2001, 97, 25S.	2.4	0
99	Modeling effects of diabetes and obesity co-morbidities in endometrial cancer development and progression. BMC Proceedings, 2012, 6, .	1.6	0
100	Potential of Pharmaceutical Intervention in Platelets and Cancer Positive Feedback Loop. FASEB Journal, 2021, 35, .	0.5	0
101	SHetA2 Increases the Activity of Palbociclib in Cervical Cancer in vitro and in vivo. FASEB Journal, 2021, 35, .	0.5	0
102	Abstract 1252: The mechanism of the drug, SHetA2, in cervical cancer cells involves growth inhibition, mitochondria damage and release of AIF to cause caspase-independent cell death., 2021 ,,.		0
103	Prevention of Gynecologic Malignancies. , 2004, , 883-919.		0
104	Abstract A13: Chemoprevention agent SHetA2 induces G1 arrest through modulation of a biological system driven by Cyclin D1. , 2008, , .		0
105	Abstract A140: Chemoprevention of familial adenomatous polyposis by a flexibleâ€heteroarotinoid (Flexâ€Het), SHetA2, in APCMinmice. , 2010, , .		0
106	Abstract 1341: Insulin directly induces endometrial cell proliferation and carcinogenesis., 2011,,.		0
107	Abstract 1798: Mortalin precursor as potential marker for chemoprevention with SHetA2., 2017,,.		0
108	Abstract 27: Development of a rat model of atypical endometrial hyperplasia and a vaginal suppository formulation of SHetA2 for chemoprevention studies. , 2020, , .		0

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109	Abstract 2840: Development of a model of dihomo-gamma-linolenic acid interference with platelet promotion of ovarian cancer. , 2020, , .		0
110	Implication of integrins in eptifibatide interference with platelet stimulation of ovarian cancer. FASEB Journal, 2022, 36, .	0.5	0