List of Publications by Year in descending order

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WENDONG HUANG

#	Article	IF	CITATIONS
1	Pleiotropic roles of FXR in liver and colorectal cancers. Molecular and Cellular Endocrinology, 2022, 543, 111543.	3.2	5
2	A narrative review of molecular mechanism and therapeutic effect of cannabidiol (CBD). Basic and Clinical Pharmacology and Toxicology, 2022, 130, 439-456.	2.5	93
3	Bile Acid–Mediated Activation of Brown Fat Protects From Alcohol-Induced Steatosis and Liver Injury in Mice. Cellular and Molecular Gastroenterology and Hepatology, 2022, 13, 809-826.	4.5	19
4	METTL16 exerts an m6A-independent function to facilitate translation and tumorigenesis. Nature Cell Biology, 2022, 24, 205-216.	10.3	143
5	Intestinal AMPK modulation of microbiota mediates crosstalk with brown fat to control thermogenesis. Nature Communications, 2022, 13, 1135.	12.8	28
6	Midnolin Regulates Liver Cancer Cell Growth In Vitro and In Vivo. Cancers, 2022, 14, 1421.	3.7	4
7	Adipocyte-derived PGE2 is required for intermittent fasting–induced Treg proliferation and improvement of insulin sensitivity. JCI Insight, 2022, 7, .	5.0	13
8	MAP3K1 Variant Causes Hyperactivation of Wnt4/ \hat{l}^2 -Catenin/FOXL2 Signaling Contributing to 46,XY Disorders/Differences of Sex Development. Frontiers in Genetics, 2022, 13, 736988.	2.3	7
9	Targeting MYC and BCL2 by a natural compound for "doubleâ€hit―lymphoma. Hematological Oncology, 2022, 40, 356-369.	1.7	2
10	Inhibition of the CDK2 and Cyclin A complex leads to autophagic degradation of CDK2 in cancer cells. Nature Communications, 2022, 13, .	12.8	31
11	Genetic characterization and drug sensitivity study of newly derived HGBL double/triple-hit lymphoma cell lines. Blood Advances, 2022, 6, 5067-5071.	5.2	2
12	PPARα alleviates iron overloadâ€induced ferroptosis in mouse liver. EMBO Reports, 2022, 23, .	4.5	34
13	Improving glucose and lipids metabolism: drug development based on bile acid related targets. Cell Stress, 2021, 5, 1-18.	3.2	8
14	Vertical sleeve gastrectomy confers metabolic improvements by reducing intestinal bile acids and lipid absorption in mice. Proceedings of the National Academy of Sciences of the United States of America, 2021, 118, .	7.1	27
15	Alternative approaches to target Myc for cancer treatment. Signal Transduction and Targeted Therapy, 2021, 6, 117.	17.1	86
16	Notoginsenoside Ft1 acts as a TGR5 agonist but FXR antagonist to alleviate high fat diet-induced obesity and insulin resistance in mice. Acta Pharmaceutica Sinica B, 2021, 11, 1541-1554.	12.0	46
17	miR-26a attenuates colitis and colitis-associated cancer by targeting the multiple intestinal inflammatory pathways. Molecular Therapy - Nucleic Acids, 2021, 24, 264-273.	5.1	19
18	Metabolic nuclear receptors coordinate energy metabolism to regulate Sox9+ hepatocyte fate. IScience, 2021, 24, 103003.	4.1	3

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19	Bile acids and metabolic surgery. Liver Research, 2021, 5, 164-170.	1.4	4
20	Danning tablets alleviate high fat diet-induced obesity and fatty liver in mice via modulating SREBP pathway. Journal of Ethnopharmacology, 2021, 279, 114320.	4.1	5
21	STAT3 Activation-Induced Fatty Acid Oxidation in CD8+ T Effector Cells Is Critical for Obesity-Promoted Breast Tumor Growth. Cell Metabolism, 2020, 31, 148-161.e5.	16.2	201
22	Sweroside ameliorates NAFLD in high-fat diet induced obese mice through the regulation of lipid metabolism and inflammatory response. Journal of Ethnopharmacology, 2020, 255, 112556.	4.1	28
23	Bile Acid Composition Contributes to Metabolic Improvements after Sleeve Gastrectomy in Mice. FASEB Journal, 2020, 34, 1-1.	0.5	Ο
24	Identification of the novel Np17 oncogene in human leukemia. Aging, 2020, 12, 23647-23667.	3.1	3
25	A Novel Compound Heterozygous CYP17A1 Variant Causes 17α-Hydroxylase/17, 20-Lyase Deficiency. Frontiers in Genetics, 2019, 10, 996.	2.3	10
26	Deletion of miR-126a Promotes Hepatic Aging and Inflammation in a Mouse Model of Cholestasis. Molecular Therapy - Nucleic Acids, 2019, 16, 494-504.	5.1	19
27	H19 potentiates let-7 family expression through reducing PTBP1 binding to their precursors in cholestasis. Cell Death and Disease, 2019, 10, 168.	6.3	34
28	Mitochondrial dysfunction caused by m.2336T>C mutation with hypertrophic cardiomyopathy in cybrid cell lines. Mitochondrion, 2019, 46, 313-320.	3.4	14
29	Mitochondrial Dysfunctions Contribute to Hypertrophic Cardiomyopathy in Patient iPSC-Derived Cardiomyocytes with MT-RNR2 Mutation. Stem Cell Reports, 2018, 10, 808-821.	4.8	74
30	Stabilization of the c-Myc Protein by CAMKIIÎ ³ Promotes T Cell Lymphoma. Cancer Cell, 2017, 32, 115-128.e7.	16.8	68
31	Myeloid adrenergic signaling via CaMKII forms a feedforward loop of catecholamine biosynthesis. Journal of Molecular Cell Biology, 2017, 9, 422-434.	3.3	15
32	Bile acid signaling and bariatric surgery. Liver Research, 2017, 1, 208-213.	1.4	14
33	The G-protein-coupled bile acid receptor Gpbar1 (TGR5) protects against renal inflammation and renal cancer cell proliferation and migration through antagonizing NF-κB and STAT3 signaling pathways. Oncotarget, 2017, 8, 54378-54387.	1.8	33
34	Vertical sleeve gastrectomy activates GPBARâ€4/TGR5 to sustain weight loss, improve fatty liver, and remit insulin resistance in mice. Hepatology, 2016, 64, 760-773.	7.3	143
35	Curcumin rescues high fat diet-induced obesity and insulin sensitivity in mice through regulating SREBP pathway. Toxicology and Applied Pharmacology, 2016, 304, 99-109.	2.8	101
36	Novel FXR (farnesoid X receptor) modulators: Potential therapies for cholesterol gallstone disease. Bioorganic and Medicinal Chemistry, 2016, 24, 3986-3993.	3.0	22

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37	FXR blocks the growth of liver cancer cells through inhibiting mTOR-s6K pathway. Biochemical and Biophysical Research Communications, 2016, 474, 351-356.	2.1	18
38	Identification of a novel RNA giant nuclear body in cancer cells. Oncotarget, 2016, 7, 4724-4734.	1.8	3
39	The G-Protein-Coupled Bile Acid Receptor Gpbar1 (TGR5) Inhibits Gastric Inflammation Through Antagonizing NF-κB Signaling Pathway. Frontiers in Pharmacology, 2015, 6, 287.	3.5	81
40	Activating CAR and \hat{l}^2 -catenin induces uncontrolled liver growth and tumorigenesis. Nature Communications, 2015, 6, 5944.	12.8	79
41	Stereoselective synthesis, biological evaluation, and modeling of novel bile acid-derived G-protein coupled Bile acid receptor 1 (GP-BAR1, TGR5) agonists. Bioorganic and Medicinal Chemistry, 2015, 23, 1613-1628.	3.0	30
42	Farnesoid X Receptor Antagonizes JNK Signaling Pathway in Liver Carcinogenesis by Activating SOD3. Molecular Endocrinology, 2015, 29, 322-331.	3.7	38
43	miR-26a enhances autophagy to protect against ethanol-induced acute liver injury. Journal of Molecular Medicine, 2015, 93, 1045-1055.	3.9	52
44	Bile acid nuclear receptor FXR and digestive system diseases. Acta Pharmaceutica Sinica B, 2015, 5, 135-144.	12.0	264
45	Bile acid signaling and liver regeneration. Biochimica Et Biophysica Acta - Gene Regulatory Mechanisms, 2015, 1849, 196-200.	1.9	82
46	Identification of miR-26a as a Target Gene of Bile Acid Receptor GPBAR-1/TGR5. PLoS ONE, 2015, 10, e0131294.	2.5	13
47	Small-molecule induction of phospho-elF4E sumoylation and degradation via targeting its phosphorylated serine 209 residue. Oncotarget, 2015, 6, 15111-15121.	1.8	14
48	The G-protein-coupled bile acid receptor Gpbar1 (TGR5) suppresses gastric cancer cell proliferation and migration through antagonizing STAT3 signaling pathway. Oncotarget, 2015, 6, 34402-34413.	1.8	47
49	Crizotinib induces autophagy through inhibition of the STAT3 pathway in multiple lung cancer cell lines. Oncotarget, 2015, 6, 40268-40282.	1.8	47
50	Autophagy inhibition sensitizes hepatocellular carcinoma to the multikinase inhibitor linifanib. Scientific Reports, 2014, 4, 6683.	3.3	56
51	GPBAR1/TGR5 Mediates Bile Acid-Induced Cytokine Expression in Murine Kupffer Cells. PLoS ONE, 2014, 9, e93567.	2.5	61
52	Hepatocarcinogenesis in FXRâ^'/â^' Mice Mimics Human HCC Progression That Operates through HNF1α Regulation of FXR Expression. Molecular Endocrinology, 2012, 26, 775-785.	3.7	97
53	Downregulation of nuclear receptor FXR is associated with multiple malignant clinicopathological characteristics in human hepatocellular carcinoma. American Journal of Physiology - Renal Physiology, 2012, 303, G1245-G1253.	3.4	80
54	CaMKII Î ³ , a critical regulator of CML stem/progenitor cells, is a target of the natural product berbamine. Blood, 2012, 120, 4829-4839.	1.4	86

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55	Promotion of liver regeneration/repair by farnesoid X receptor in both liver and intestine in mice. Hepatology, 2012, 56, 2336-2343.	7.3	121
56	Deletion of IFNÎ ³ enhances hepatocarcinogenesis in FXR knockout mice. Journal of Hepatology, 2012, 57, 1004-1012.	3.7	25
57	Farnesoid X receptor antagonizes nuclear factor l̂ºB in hepatic inflammatory response. Hepatology, 2008, 48, 1632-1643.	7.3	498
58	Spontaneous Development of Liver Tumors in the Absence of the Bile Acid Receptor Farnesoid X Receptor. Cancer Research, 2007, 67, 863-867.	0.9	397
59	Guarding the gate against hyperbilirubinaemia. Gut, 0, , gutjnl-2022-327532.	12.1	0