

Zhiwei Song

List of Publications by Year in descending order

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Version: 2024-02-01

30
papers

1,379
citations

430874

18
h-index

454955

30
g-index

31
all docs

31
docs citations

31
times ranked

1918
citing authors

#	ARTICLE	IF	CITATIONS
1	The "less-is-more" in therapeutic antibodies: Afucosylated anti-cancer antibodies with enhanced antibody-dependent cellular cytotoxicity. <i>MABs</i> , 2018, 10, 693-711.	5.2	215
2	Challenges of glycosylation analysis and control: an integrated approach to producing optimal and consistent therapeutic drugs. <i>Drug Discovery Today</i> , 2016, 21, 740-765.	6.4	164
3	IRES-mediated Tricistronic vectors for enhancing generation of high monoclonal antibody expressing CHO cell lines. <i>Journal of Biotechnology</i> , 2012, 157, 130-139.	3.8	136
4	Roles of the nucleotide sugar transporters (SLC35 family) in health and disease. <i>Molecular Aspects of Medicine</i> , 2013, 34, 590-600.	6.4	109
5	The sweet tooth of biopharmaceuticals: Importance of recombinant protein glycosylation analysis. <i>Biotechnology Journal</i> , 2012, 7, 1462-1472.	3.5	99
6	Optimization of Heavy Chain and Light Chain Signal Peptides for High Level Expression of Therapeutic Antibodies in CHO Cells. <i>PLoS ONE</i> , 2015, 10, e0116878.	2.5	83
7	RNAi suppression of Bax and Bak enhances viability in fed-batch cultures of CHO cells. <i>Metabolic Engineering</i> , 2006, 8, 509-522.	7.0	82
8	Inactivation of GDP-fucose transporter gene (<i>Slc35c1</i>) in CHO cells by ZFNs, TALENs and CRISPR-Cas9 for production of fucose-free antibodies. <i>Biotechnology Journal</i> , 2016, 11, 399-414.	3.5	57
9	Identification of functional elements of the GDP-fucose transporter SLC35C1 using a novel Chinese hamster ovary mutant. <i>Glycobiology</i> , 2012, 22, 897-911.	2.5	44
10	The Golgi CMP-sialic acid transporter: A new CHO mutant provides functional insights. <i>Glycobiology</i> , 2008, 18, 851-860.	2.5	42
11	Golgi Phosphoprotein 3 Mediates the Golgi Localization and Function of Protein O-Linked Mannose β -1,2-N-Acetylglucosaminyltransferase 1. <i>Journal of Biological Chemistry</i> , 2014, 289, 14762-14770.	3.4	42
12	A functional analysis of N-glycosylation-related genes on sialylation of recombinant erythropoietin in six commonly used mammalian cell lines. <i>Metabolic Engineering</i> , 2010, 12, 526-536.	7.0	40
13	Caspase activation, sialidase release and changes in sialylation pattern of recombinant human erythropoietin produced by CHO cells in batch and fed-batch cultures. <i>Cytotechnology</i> , 2006, 51, 67-79.	1.6	29
14	Specific inhibition of caspase-8 and -9 in CHO cells enhances cell viability in batch and fed-batch cultures. <i>Metabolic Engineering</i> , 2007, 9, 406-418.	7.0	27
15	CHO-gmt5, a novel CHO glycosylation mutant for producing afucosylated and asialylated recombinant antibodies. <i>Bioengineered</i> , 2013, 4, 90-94.	3.2	25
16	RCA-I-resistant CHO mutant cells have dysfunctional GnT I and expression of normal GnT I in these mutants enhances sialylation of recombinant erythropoietin. <i>Metabolic Engineering</i> , 2010, 12, 360-368.	7.0	24
17	Attenuated glutamine synthetase as a selection marker in CHO cells to efficiently isolate highly productive stable cells for the production of antibodies and other biologics. <i>MABs</i> , 2019, 11, 965-976.	5.2	23
18	Highly sialylated recombinant human erythropoietin production in large-scale perfusion bioreactor utilizing CHO-gmt4 (JW152) with restored GnT I function. <i>Biotechnology Journal</i> , 2014, 9, 100-109.	3.5	21

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19	Identification of essential amino acid residues in the hydrophilic loop regions of the CMP-sialic acid transporter and UDP-galactose transporter. <i>Glycobiology</i> , 2010, 20, 689-701.	2.5	19
20	CHO Glycosylation Mutants as Potential Host Cells to Produce Therapeutic Proteins with Enhanced Efficacy. <i>Advances in Biochemical Engineering/Biotechnology</i> , 2012, 131, 63-87.	1.1	18
21	Targeting of embryonic annexin A2 expressed on ovarian and breast cancer by the novel monoclonal antibody 2448. <i>Oncotarget</i> , 2018, 9, 13206-13221.	1.8	17
22	Characterization of a GDP-Fucose Transporter and a Fucosyltransferase Involved in the Fucosylation of Glycoproteins in the Diatom <i>Phaeodactylum tricornutum</i> . <i>Frontiers in Plant Science</i> , 2019, 10, 610.	3.6	14
23	Producing recombinant therapeutic glycoproteins with enhanced sialylation using CHO-gmt4 glycosylation mutant cells. <i>Bioengineered</i> , 2014, 5, 269-273.	3.2	11
24	A quantitative study of the Golgi retention of glycosyltransferases. <i>Journal of Cell Science</i> , 2021, 134, .	2.0	10
25	Deciphering O-glycomics for the development and production of biopharmaceuticals. <i>Pharmaceutical Bioprocessing</i> , 2013, 1, 89-104.	0.8	8
26	Improving Antibody Production in Stably Transfected CHO Cells by CRISPR/Cas9-Mediated Inactivation of Genes Identified in a Large-Scale Screen with Chinese Hamster-Specific siRNAs. <i>Biotechnology Journal</i> , 2021, 16, e2000267.	3.5	7
27	Sodium butyrate enhances the acidic isoform content of recombinant human erythropoietin produced by Chinese hamster ovary cells. <i>Biotechnology Letters</i> , 2014, 36, 907-911.	2.2	5
28	Improving sialylation of recombinant biologics for enhanced therapeutic efficacy. <i>Pharmaceutical Bioprocessing</i> , 2014, 2, 363-366.	0.8	3
29	Bad plays a more significant role than Bid and Bim in mediating cell death signals in batch cultures of HEK 293 cells. <i>Biotechnology Letters</i> , 2008, 30, 819-827.	2.2	2
30	Production of Highly Sialylated Recombinant Glycoproteins Using <i>Ricinus communis</i> Agglutinin-I-Resistant CHO Glycosylation Mutants. <i>Methods in Molecular Biology</i> , 2015, 1321, 323-333.	0.9	1