

# Shahram Misaghi

## List of Publications by Year in descending order

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23  
papers

495  
citations

840776

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677142

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#	ARTICLE	IF	CITATIONS
1	Association of C-Terminal Ubiquitin Hydrolase BRCA1-Associated Protein 1 with Cell Cycle Regulator Host Cell Factor 1. <i>Molecular and Cellular Biology</i> , 2009, 29, 2181-2192.	2.3	187
2	Chinese hamster ovary K1 host cell enables stable cell line development for antibody molecules which are difficult to express in DUXB11-derived dihydrofolate reductase deficient host cell. <i>Biotechnology Progress</i> , 2013, 29, 980-985.	2.6	40
3	FX knockout CHO hosts can express desired ratios of fucosylated or afucosylated antibodies with high titers and comparable product quality. <i>Biotechnology and Bioengineering</i> , 2017, 114, 632-644.	3.3	37
4	Probing the importance of clonality: Single cell subcloning of clonally derived CHO cell lines yields widely diverse clones differing in growth, productivity, and product quality. <i>Biotechnology Progress</i> , 2018, 34, 624-634.	2.6	36
5	Carboxypeptidase D is the only enzyme responsible for antibody C-terminal lysine cleavage in Chinese hamster ovary (CHO) cells. <i>Biotechnology and Bioengineering</i> , 2016, 113, 2100-2106.	3.3	30
6	It's time to regulate: Coping with product-induced nongenetic clonal instability in CHO cell lines via regulated protein expression. <i>Biotechnology Progress</i> , 2014, 30, 1432-1440.	2.6	24
7	Increased Targeting of Donor Switch Region and IgE in S $\beta$ 1-Deficient B Cells. <i>Journal of Immunology</i> , 2010, 185, 166-173.	0.8	18
8	Beating the odds: The poisson distribution of all input cells during limiting dilution grossly underestimates whether a cell line is clonally derived or not. <i>Biotechnology Progress</i> , 2018, 34, 559-569.	2.6	17
9	Tailoring translational strength using Kozak sequence variants improves bispecific antibody assembly and reduces product-related impurities in CHO cells. <i>Biotechnology and Bioengineering</i> , 2020, 117, 1946-1960.	3.3	16
10	Resilient immortals, characterizing and utilizing Bax/Bak deficient Chinese hamster ovary (CHO) cells for high titer antibody production. <i>Biotechnology Progress</i> , 2013, 29, 727-737.	2.6	15
11	Slashing the timelines: Opting to generate high-titer clonal lines faster via viability-based single cell sorting. <i>Biotechnology Progress</i> , 2016, 32, 198-207.	2.6	14
12	Taming hyperactive DNase I: Stable inducible expression of a hyperactive salt- and actin-resistant variant of human deoxyribonuclease I in CHO cells. <i>Biotechnology Progress</i> , 2017, 33, 523-533.	2.6	12
13	Pyruvate Kinase Muscle $\beta$ 1 Expression Appears to Drive Lactogenic Behavior in CHO Cell Lines, Triggering Lower Viability and Productivity: A Case Study. <i>Biotechnology Journal</i> , 2019, 14, 1800332.	3.5	7
14	Utilizing a regulated targeted integration cell line development approach to systematically investigate what makes an antibody difficult to express. <i>Biotechnology Progress</i> , 2019, 35, e2772.	2.6	7
15	Bax and Bak knockout apoptosis-resistant Chinese hamster ovary cell lines significantly improve culture viability and titer in intensified fed-batch culture process. <i>Biotechnology Progress</i> , 2022, 38, e3228.	2.6	7
16	Insulin degrading enzyme (IDE) expressed by Chinese hamster ovary (CHO) cells is responsible for degradation of insulin in culture media. <i>Journal of Biotechnology</i> , 2020, 320, 44-49.	3.8	5
17	Endothelial intercellular cell adhesion molecule 1 contributes to cell aggregate formation in CHO cells cultured in serum-free media. <i>Biotechnology Progress</i> , 2020, 36, e2951.	2.6	4
18	Preventing pyruvate kinase muscle expression in Chinese hamster ovary cells curbs lactogenic behavior by altering glycolysis, gating pyruvate generation, and increasing pyruvate flux into the TCA cycle. <i>Biotechnology Progress</i> , 2021, 37, e3193.	2.6	4

#	ARTICLE	IF	CITATIONS
19	UBR E3 ligases and the PDIA3 protease control degradation of unfolded antibody heavy chain by ERAD. Journal of Cell Biology, 2020, 219, .	5.2	4
20	High Intracellular Seed Train BiP Levels Correlate With Poor Production Culture Performance in CHO Cells. Biotechnology Journal, 2018, 13, e1700746.	3.5	3
21	Concurrent transfection of randomized transgene configurations into targeted integration CHO host is an advantageous and cost-effective method for expression of complex molecules. Biotechnology Journal, 2021, 16, e2000230.	3.5	3
22	A ribonucleoprotein-based decaplex CRISPR/Cas9 knockout strategy for CHO host engineering. Biotechnology Progress, 2022, 38, e3212.	2.6	3
23	Supplementation of Nucleosides During Selection can Reduce Sequence Variant Levels in CHO Cells Using GS/MSX Selection System. Biotechnology Journal, 2018, 13, 1700335.	3.5	2