Ondrej Pös

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/8373794/publications.pdf

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18	602	11	13	
papers	citations	h-index	g-index	
19	19	19	711	
all docs	docs citations	times ranked	citing authors	

#	Article	IF	CITATIONS
1	Circulating cell-free nucleic acids: characteristics and applications. European Journal of Human Genetics, 2018, 26, 937-945.	2.8	163
2	Circulating Cell-Free Nucleic Acids: Main Characteristics and Clinical Application. International Journal of Molecular Sciences, 2020, 21, 6827.	4.1	110
3	DNA copy number variation: Main characteristics, evolutionary significance, and pathological aspects. Biomedical Journal, 2021, 44, 548-559.	3.1	89
4	Recent trends in prenatal genetic screening and testing. F1000Research, 2019, 8, 764.	1.6	46
5	Technical and Methodological Aspects of Cell-Free Nucleic Acids Analyzes. International Journal of Molecular Sciences, 2020, 21, 8634.	4.1	34
6	Endogenous H2S producing enzymes are involved in apoptosis induction in clear cell renal cell carcinoma. BMC Cancer, 2018, 18, 591.	2.6	33
7	Non-invasive prenatal testing (NIPT) by low coverage genomic sequencing: Detection limits of screened chromosomal microdeletions. PLoS ONE, 2020, 15, e0238245.	2.5	26
8	Copy Number Variation: Methods and Clinical Applications. Applied Sciences (Switzerland), 2021, 11, 819.	2.5	26
9	The Role of Exosomes in Cancer Progression. International Journal of Molecular Sciences, 2022, 23, 8.	4.1	23
10	Identification of Structural Variation from NGS-Based Non-Invasive Prenatal Testing. International Journal of Molecular Sciences, 2019, 20, 4403.	4.1	20
11	Quantification of peripheral whole blood, cell-free plasma and exosome encapsulated mitochondrial DNA copy numbers in patients with atrial fibrillation. Journal of Biotechnology, 2019, 299, 66-71.	3.8	19
12	Liquid Biopsy as a Source of Nucleic Acid Biomarkers in the Diagnosis and Management of Lynch Syndrome. International Journal of Molecular Sciences, 2022, 23, 4284.	4.1	10
13	Ultracentrifugation enrichment protocol followed by total RNA sequencing allows assembly of the complete mitochondrial genome. Journal of Biotechnology, 2019, 299, 8-12.	3.8	2
14	Prenatal genetic diagnosis: Fetal therapy as a possible solution to a positive test. Acta Biomedica, 2020, 91, e2020021.	0.3	0
15	Title is missing!. , 2020, 15, e0238245.		O
16	Title is missing!. , 2020, 15, e0238245.		0
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