

Jan HojnÃ½

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4101732/publications.pdf>

Version: 2024-02-01

19
papers

350
citations

933447

10
h-index

839539

18
g-index

19
all docs

19
docs citations

19
times ranked

627
citing authors

#	ARTICLE	IF	CITATIONS
1	Identification of deleterious germline CHEK2 mutations and their association with breast and ovarian cancer. <i>International Journal of Cancer</i> , 2019, 145, 1782-1797.	5.1	62
2	Multiplex PCR and NGS-based identification of mRNA splicing variants: Analysis of BRCA1 splicing pattern as a model. <i>Gene</i> , 2017, 637, 41-49.	2.2	43
3	A comprehensive evaluation of pathogenic mutations in primary cutaneous melanomas, including the identification of novel loss-of-function variants. <i>Scientific Reports</i> , 2019, 9, 17050.	3.3	42
4	Expression of human BRCA1 ¹⁷ alternative splicing variant with a truncated BRCT domain in MCF-7 cells results in impaired assembly of DNA repair complexes and aberrant DNA damage response. <i>Cellular Signalling</i> , 2013, 25, 1186-1193.	3.6	35
5	Ovarian mesonephric-like adenocarcinoma arising in serous borderline tumor: a case report with complex morphological and molecular analysis. <i>Diagnostic Pathology</i> , 2020, 15, 91.	2.0	35
6	Association of Germline CHEK2 Gene Variants with Risk and Prognosis of Non-Hodgkin Lymphoma. <i>PLoS ONE</i> , 2015, 10, e0140819.	2.5	31
7	Leiomyoma with Bizarre Nuclei: a Study of 108 Cases Focusing on Clinicopathological Features, Morphology, and Fumarate Hydratase Alterations. <i>Pathology and Oncology Research</i> , 2020, 26, 1527-1537.	1.9	22
8	Germline mutation in the TP53 gene in uveal melanoma. <i>Scientific Reports</i> , 2018, 8, 7618.	3.3	13
9	Synchronous endometrioid endometrial and ovarian carcinomas are biologically related: A clinico-pathological and molecular (next generation sequencing) study of 22½ cases. <i>Oncology Letters</i> , 2019, 17, 2207-2214.	1.8	12
10	Results of targeted next-generation sequencing in children with cystic kidney diseases often change the clinical diagnosis. <i>PLoS ONE</i> , 2020, 15, e0235071.	2.5	12
11	Uterine cellular leiomyomas are characterized by common HMGA2 aberrations, followed by chromosome 1p deletion and MED12 mutation: morphological, molecular, and immunohistochemical study of 52 cases. <i>Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin</i> . 2022, 480, 281-291.	2.8	10
12	Expression, Epigenetic, and Genetic Changes of HNF1B in Colorectal Lesions: an Analysis of 145 Cases. <i>Pathology and Oncology Research</i> , 2020, 26, 2337-2350.	1.9	7
13	HNF1B, EZH2 and ECI2 in prostate carcinoma. Molecular, immunohistochemical and clinico-pathological study. <i>Scientific Reports</i> , 2020, 10, 14365.	3.3	6
14	Analysis of expression, epigenetic, and genetic changes of HNF1B in 130 kidney tumours. <i>Scientific Reports</i> , 2020, 10, 17151.	3.3	5
15	Identification of novel HNF1B mRNA splicing variants and their qualitative and semi-quantitative profile in selected healthy and tumour tissues. <i>Scientific Reports</i> , 2020, 10, 6958.	3.3	5
16	A comprehensive analysis of the expression, epigenetic and genetic changes of HNF1B and ECI2 in 122 cases of high-grade serous ovarian carcinoma. <i>Oncology Letters</i> , 2021, 21, 185.	1.8	4
17	Microscopic extraovarian sex cord proliferation: report of a case with bilateral Fallopian tube involvement and a comprehensive molecular analysis. <i>Polish Journal of Pathology</i> , 2020, 71, 175-180.	0.3	2
18	Comprehensive quantitative analysis of alternative splicing variants reveals the HNF1B mRNA splicing pattern in various tumour and non-tumour tissues. <i>Scientific Reports</i> , 2022, 12, 199.	3.3	2

#	ARTICLE	IF	CITATIONS
19	Desmoplastic Small Round Cell Tumor of the Uterus: A Report of Molecularly Confirmed Case with EWSR1-WT1 Fusion. <i>Diagnostics</i> , 2022, 12, 1184.	2.6	2