Barbara Mandriani

List of Publications by Year
in descending order

16	830	687363 13	839539 18
papers	citations	h-index	g-index

18	18	18	4278
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	The psychological impact of COVIDâ€19 pandemic on patients with neuroendocrine tumors: Between resilience and vulnerability. Journal of Neuroendocrinology, 2021, 33, e13041.	2.6	3
2	Adoptive T-cell immunotherapy in digestive tract malignancies: Current challenges and future perspectives. Cancer Treatment Reviews, 2021, 100, 102288.	7.7	9
3	The Role of Cytotoxic Chemotherapy in Well-Differentiated Gastroenteropancreatic and Lung Neuroendocrine Tumors. Current Treatment Options in Oncology, 2019, 20, 72.	3.0	7
4	DAXX mutations as potential genomic markers of malignant evolution in small nonfunctioning pancreatic neuroendocrine tumors. Scientific Reports, 2019, 9, 18614.	3.3	26
5	gene2drug: a computational tool for pathway-based rational drug repositioning. Bioinformatics, 2018, 34, 1498-1505.	4.1	62
6	TRIM50 regulates Beclin 1 proautophagic activity. Biochimica Et Biophysica Acta - Molecular Cell Research, 2018, 1865, 908-919.	4.1	39
7	Dissecting KMT2D missense mutations in Kabuki syndrome patients. Human Molecular Genetics, 2018, 27, 3651-3668.	2.9	49
8	GNB5 Mutations Cause an Autosomal-Recessive Multisystem Syndrome with Sinus Bradycardia and Cognitive Disability. American Journal of Human Genetics, 2016, 99, 704-710.	6.2	58
9	Identification of p53-target genes in Danio rerio. Scientific Reports, 2016, 6, 32474.	3.3	10
10	BRF1 mutations alter RNA polymerase IIIâ€“dependent transcription and cause neurodevelopmental anomalies. Genome Research, 2015, 25, 155-166.	5.5	85
11	TRIM8 downregulation in glioma affects cell proliferation and it is associated with patients survival. BMC Cancer, 2015, 15, 470.	2.6	61
12	TBC1D7Mutations are Associated with Intellectual Disability, Macrocrania, Patellar Dislocation, and Celiac Disease. Human Mutation, 2014, 35, 447-451.	2.5	52
13	Smaller and larger deletions of the Williams Beuren syndrome region implicate genes involved in mild facial phenotype, epilepsy and autistic traits. European Journal of Human Genetics, 2014, 22, 64-70.	2.8	63
14	Molecular Analysis, Pathogenic Mechanisms, and Readthrough Therapy on a Large Cohort of<scp>K</scp>abuki Syndrome Patients. Human Mutation, 2014, 35, 841-850.	2.5	87
15	HDAC6 mediates the acetylation of TRIM50. Cellular Signalling, 2014, 26, 363-369.	3.6	17
16	Cutting Edge: The NLRP3 Inflammasome Links Complement-Mediated Inflammation and IL-1Î² Release. Journal of Immunology, 2013, 191, 1006-1010.	0.8	173