Pedro A Lage

List of Publications by Year
in descending order

16	727	840776 11	888059 17
papers	citations	h-index	g-index

17	17	17	1361
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	The 'just-right' signaling model: APC somatic mutations are selected based on a specific level of activation of the beta-catenin signaling cascade. Human Molecular Genetics, 2002, 11, 1549-1560.	2.9	317
2	Distinct patterns of KRAS mutations in colorectal carcinomas according to germline mismatch repair defects and hMLH1 methylation status. Human Molecular Genetics, 2004, 13, 2303-2311.	2.9	127
3	Pemphigus vulgaris with exclusive involvement of the esophagus: case report and review. Gastrointestinal Endoscopy, 2004, 60, 312-315.	1.0	55
4	Colorectal cancers show distinct mutation spectra in members of the canonical WNT signaling pathway according to their anatomical location and type of genetic instability. Genes Chromosomes and Cancer, 2010, 49, 746-759.	2.8	51
5	Gastric malt lymphoma: Analysis of a series of consecutive patients over 20 years. United European Gastroenterology Journal, 2016, 4, 395-402.	3.8	31
6	Familial colorectal cancer type X syndrome: two distinct molecular entities?. Familial Cancer, 2011, 10, 623-631.	1.9	27
7	High Rate of Percutaneous Endoscopic Gastrostomy Site Infections Due to Oropharyngeal Colonization. Digestive Diseases and Sciences, 2006, 51, 2384-2388.	2.3	20
8	Esophageal stenosis with sloughing esophagitis: A curious manifestation of graft-vs-host disease. World Journal of Gastroenterology, 2015, 21, 9217.	3.3	19
9	Colorectal Adenomas in Young Patients: Microsatellite Instability is not a Useful Marker to Detect New Cases of Lynch Syndrome. Diseases of the Colon and Rectum, 2008, 51, 909-915.	1.3	18
10	Aggressive Phenotype of MYH-Associated Polyposis with Jejunal Cancer and Intra-Abdominal Desmoid Tumor. Diseases of the Colon and Rectum, 2009, 52, 742-745.	1.3	17
11	Bethesda criteria for microsatellite instability testing: impact on the detection of new cases of Lynch syndrome. Familial Cancer, 2012, 11, 571-578.	1.9	15
12	Pediatric Colorectal Cancer: A Heterogenous Entity. Journal of Pediatric Hematology/Oncology, 2020, 42, 131-135.	0.6	7
13	Association of colonic and endometrial carcinomas in Portuguese families with hereditary nonpolyposis colorectal carcinoma significantly increases the probability of detecting a pathogenic mutation in mismatch repair genes, primarily theMSH2gene. Cancer, 2004, 101, 172-177.	4.1	6
14	APC Somatic Mosaicism in a Patient with Gardner Syndrome Carrying the E1573X Mutation: Report of a Case. Diseases of the Colon and Rectum, 2009, 52, 1516-1520.	1.3	6
15	Serrated polyposis associated with a family history of colorectal cancer and/or polyps: The preferential location of polyps in the colon and rectum defines two molecular entities. International Journal of Molecular Medicine, 2016, 38, 687-702.	4.0	3
16	The nonsense mutation MSH2 c.2152C>T shows a founder effect in Portuguese Lynch syndrome families. Genes Chromosomes and Cancer, 2019, 58, 657-664.	2.8	3