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List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Prevalence of Lynch syndrome in unselected patients with endometrial or ovarian cancer. Archives of Gynecology and Obstetrics, 2016, 294, 1299-1303.	1.7	4
2	An unusual case of Cowden syndrome associated with ganglioneuromatous polyposis. Hereditary Cancer in Clinical Practice, 2016, 14, 11.	1.5	6
3	Genomic rearrangements in MSH2, MLH1 or MSH6 are rare in HNPCC patients carrying point mutations. Cancer Letters, 2007, 248, 89-95.	7.2	18
4	N-acetyltransferase (NAT) 2 acetylator status and age of onset in patients with hereditary nonpolyposis colorectal cancer (HNPCC). Cancer Letters, 2006, 241, 150-157.	7.2	14
5	Resektionsausmaß und Therapiekonzept bei hereditäem, nicht Polyposis-assoziiertem kolorektalem Karzinom (HNPCC) – Indexpatient: chirurgische Strategie. Visceral Medicine, 2006, 22, 1-5.	1.3	0
6	N-Acetyltransferase (NAT) 2 acetylator status and age of tumour onset in patients with sporadic and familial, microsatellite stable (MSS) colorectal cancer. International Journal of Colorectal Disease, 2006, 22, 137-143.	2.2	9
7	Occult endometrial cancer and decision making for prophylactic hysterectomy in hereditary nonpolyposis colorectal cancer patients. Gynecologic Oncology, 2006, 102, 189-194.	1.4	30
8	Loss of MSH3 Protein Expression Is Frequent in MLH1-Deficient Colorectal Cancer and Is Associated with Disease Progression 1. Cancer Research, 2004, 64, 864-870.	0.9	48
9	Methylenetetrahydrofolate reductase polymorphisms and risk of sporadic and hereditary colorectal cancer with or without microsatellite instability. Cancer Letters, 2003, 191, 179-185.	7.2	40
10	Seven novel MLH1 and MSH2 germline mutations in hereditary nonpolyposis colorectal cancer. Human Mutation, 2002, 19, 82-82.	2.5	9
11	Involvement ofhMSH6in the development of hereditary and sporadic colorectal cancer revealed by immunostaining is based on germline mutations, but rarely on somatic inactivation. International Journal of Cancer, 2002, 97, 643-648.	5.1	72
12	Sequence analysis of the mismatch repair gene hMSH6 in the germline of patients with familial and sporadic colorectal cancer. , 2000, 85, 606-613.		53