Salvatore Romeo

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Absence of disruptive TP53 mutations in highâ€risk human papillomavirusâ€driven neck squamous cell carcinoma of unknown primary. Head and Neck, 2019, 41, 3833-3841.	2.0	2
2	Human papillomavirus as prognostic marker with rising prevalence in neck squamous cell carcinoma of unknown primary: A retrospective multicentre study. European Journal of Cancer, 2017, 74, 73-81.	2.8	59
3	Prognostic significance of LINE-1 hypomethylation in oropharyngeal squamous cell carcinoma. Clinical Epigenetics, 2017, 9, 58.	4.1	32
4	Head and Neck Extranodal Interdigitating Dendritic Cell Sarcoma: Case Report and Review of the Literature. Head and Neck Pathology, 2016, 10, 145-151.	2.6	18
5	Concomitant KIT/BRAF and PDGFRA/BRAF mutations are rare events in gastrointestinal stromal tumors. Oncotarget, 2016, 7, 30109-30118.	1.8	25
6	Investigating nasal cytology as a potential tool for diagnosing occupational rhinitis in woodworkers. International Forum of Allergy and Rhinology, 2015, 5, 814-819.	2.8	9
7	Primary Synovial Sarcoma (SS) of the digestive system: a molecular and clinicopathological study of fifteen cases. Clinical Sarcoma Research, 2015, 5, 7.	2.3	39
8	The prevalence of human papillomavirus in squamous cell carcinoma of unknown primary site metastatic to neck lymph nodes: a systematic review. Clinical and Experimental Metastasis, 2015, 32, 835-845.	3.3	41
9	Is Neck Dissection Necessary After Induction Plus Concurrent Chemoradiotherapy in Complete Responder Head and Neck Cancer Patients with Pretherapy Advanced Nodal Disease?. Annals of Surgical Oncology, 2013, 20, 250-256.	1.5	6
10	MEF2 Is a Converging Hub for Histone Deacetylase 4 and Phosphatidylinositol 3-Kinase/Akt-Induced Transformation. Molecular and Cellular Biology, 2013, 33, 4473-4491.	2.3	48
11	Dedifferentiation in Gastrointestinal Stromal Tumor to an Anaplastic KIT-negative Phenotype. American Journal of Surgical Pathology, 2013, 37, 385-392.	3.7	90
12	Malignant fibrous histiocytoma and fibrosarcoma of bone: a re-assessment in the light of currently employed morphological, immunohistochemical and molecular approaches. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2012, 461, 561-570.	2.8	78
13	The clinical impact of molecular techniques on diagnostic pathology of soft tissue and bone tumours. Diagnostic Histopathology, 2012, 18, 81-85.	0.4	2
14	Interobserver reliability in the histopathological diagnosis of cartilaginous tumors in patients with multiple osteochondromas. Modern Pathology, 2012, 25, 1275-1283.	5.5	37
15	Clear Cell Sarcoma of the Ileum. International Journal of Surgical Pathology, 2012, 20, 401-406.	0.8	24
16	Peripheral chondrosarcoma progression is associated with increased type X collagen and vascularisation. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2012, 460, 95-102.	2.8	16
17	Clinical application of molecular pathology in sarcomas. Current Opinion in Oncology, 2011, 23, 379-384.	2.4	15
18	Nora's lesion of the thumb and a differential diagnosis. European Journal of Plastic Surgery, 2011, 34,	0.6	0

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19	Reclassification and subtyping of so-called malignant fibrous histiocytoma of bone: comparison with cytogenetic features. Clinical Sarcoma Research, 2011, 1, 10.	2.3	13
20	Soft tissue tumors associated with EWSR1 translocation. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2010, 456, 219-234.	2.8	149
21	Hierarchical clustering of flow cytometry data for the study of conventional central chondrosarcoma. Journal of Cellular Physiology, 2010, 225, 601-611.	4.1	19
22	Primary cilia organization reflects polarity in the growth plate and implies loss of polarity and mosaicism in osteochondroma. Laboratory Investigation, 2010, 90, 1091-1101.	3.7	73
23	Heterogeneous and Complex Rearrangements of Chromosome Arm 6q in Chondromyxoid Fibroma. American Journal of Pathology, 2010, 177, 1365-1376.	3.8	32
24	Cell Cycle/Apoptosis Molecule Expression Correlates with Imatinib Response in Patients with Advanced Gastrointestinal Stromal Tumors. Clinical Cancer Research, 2009, 15, 4191-4198.	7.0	61
25	A balanced t(5;17) (p15;q22-23) in chondroblastoma: frequency of the re-arrangement and analysis of the candidate genes. BMC Cancer, 2009, 9, 393.	2.6	18
26	Osteosarcoma originates from mesenchymal stem cells in consequence of aneuploidization and genomic loss of <i>Cdkn2</i> . Journal of Pathology, 2009, 219, 294-305.	4.5	234
27	Profiling of high-grade central osteosarcoma and its putative progenitor cells identifies tumourigenic pathways. British Journal of Cancer, 2009, 101, 1909-1918.	6.4	67
28	Benign Cartilaginous Tumors of Bone. Advances in Anatomic Pathology, 2009, 16, 307-315.	4.3	59
29	Expression of Cellular FLICE Inhibitory Protein, Caspase-8, and Protease Inhibitor-9 in Ewing Sarcoma and Implications for Susceptibility to Cytotoxic Pathways. Clinical Cancer Research, 2007, 13, 206-214.	7.0	28
30	The role of noncartilage-specific molecules in differentiation of cartilaginous tumors. Cancer, 2007, 110, 385-394.	4.1	25
31	Functional imaging of multidrug resistance in an orthotopic model of osteosarcoma using 99mTc-sestamibi. European Journal of Nuclear Medicine and Molecular Imaging, 2007, 34, 1793-1803.	6.4	22
32	Multidrug resistance mediated by ABC transporters in osteosarcoma cell lines: mRNA analysis and functional radiotracer studies. Nuclear Medicine and Biology, 2006, 33, 831-840.	0.6	38
33	TGF-β1 drives partial myofibroblastic differentiation in chondromyxoid fibroma of bone. Journal of Pathology, 2006, 208, 26-34.	4.5	24
34	Chondromyxoid fibroma resemblesin vitro chondrogenesis, but differs in expression of signalling molecules. Journal of Pathology, 2005, 206, 135-142.	4.5	27
35	Expression of cartilage growth plate signalling molecules in chondroblastoma. Journal of Pathology, 2004, 202, 113-120.	4.5	45
36	Multiple Familial Facial Glomus: Case Report and Review of the Literature. Annals of Otology, Rhinology and Laryngology, 2003, 112, 287-292.	1.1	9