

Adele Clarkson

List of Publications by Year in descending order

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63
papers

4,167
citations

94381

37
h-index

118793

62
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63
all docs

63
docs citations

63
times ranked

5528
citing authors

#	ARTICLE	IF	CITATIONS
1	Succinate Dehydrogenase (SDH)-deficient Renal Carcinoma. American Journal of Surgical Pathology, 2014, 38, 1588-1602.	2.1	282
2	Immunohistochemistry for SDHB triages genetic testing of SDHB, SDHC, and SDHD in paraganglioma-pheochromocytoma syndromes. Human Pathology, 2010, 41, 805-814.	1.1	235
3	Loss of Nuclear Expression of Parafibromin Distinguishes Parathyroid Carcinomas and Hyperparathyroidism-Jaw Tumor (HPT-JT) Syndrome-related Adenomas From Sporadic Parathyroid Adenomas and Hyperplasias. American Journal of Surgical Pathology, 2006, 30, 1140-1149.	2.1	213
4	Immunohistochemistry for SDHB Divides Gastrointestinal Stromal Tumors (GISTs) into 2 Distinct Types. American Journal of Surgical Pathology, 2010, 34, 636-644.	2.1	210
5	Renal Tumors Associated With Germline SDHB Mutation Show Distinctive Morphology. American Journal of Surgical Pathology, 2011, 35, 1578-1585.	2.1	184
6	Assessing mutant p53 in primary high-grade serous ovarian cancer using immunohistochemistry and massively parallel sequencing. Scientific Reports, 2016, 6, 26191.	1.6	162
7	BRAFV600E Immunohistochemistry Facilitates Universal Screening of Colorectal Cancers for Lynch Syndrome. American Journal of Surgical Pathology, 2013, 37, 1592-1602.	2.1	125
8	Accuracy of Combined Protein Gene Product 9.5 and Parafibromin Markers for Immunohistochemical Diagnosis of Parathyroid Carcinoma. Journal of Clinical Endocrinology and Metabolism, 2009, 94, 434-441.	1.8	120
9	Renal Tumors and Hereditary Pheochromocytoma-Paraganglioma Syndrome Type 4. New England Journal of Medicine, 2011, 364, 885-886.	13.9	120
10	Loss of SDHA Expression Identifies SDHA Mutations in Succinate Dehydrogenase-deficient Gastrointestinal Stromal Tumors. American Journal of Surgical Pathology, 2013, 37, 226-233.	2.1	119
11	Immunohistochemistry for PMS2 and MSH6 alone can replace a four antibody panel for mismatch repair deficiency screening in colorectal adenocarcinoma. Pathology, 2010, 42, 409-413.	0.3	114
12	Wnt Pathway Inhibitors Are Strongly Down-Regulated in Pituitary Tumors. Endocrinology, 2008, 149, 1235-1242.	1.4	104
13	Loss of expression of BAP1 predicts longer survival in mesothelioma. Pathology, 2015, 47, 302-307.	0.3	102
14	Fumarate Hydratase-deficient Uterine Leiomyomas Occur in Both the Syndromic and Sporadic Settings. American Journal of Surgical Pathology, 2016, 40, 599-607.	2.1	102
15	Improving Diagnosis of Tumor-Induced Osteomalacia With Gallium-68 DOTATATE PET/CT. Journal of Clinical Endocrinology and Metabolism, 2013, 98, 687-694.	1.8	100
16	Loss of expression of BAP1 is a useful adjunct, which strongly supports the diagnosis of mesothelioma in effusion cytology. Modern Pathology, 2015, 28, 1360-1368.	2.9	95
17	Fibroblast Growth Factor 23: A New Clinical Marker for Oncogenic Osteomalacia. Journal of Clinical Endocrinology and Metabolism, 2003, 88, 4088-4094.	1.8	92
18	Variation of O6-methylguanine-DNA methyltransferase (MGMT) promoter methylation in serial samples in glioblastoma. Journal of Neuro-Oncology, 2008, 87, 71-78.	1.4	86

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19	The tumor suppressor CDC73 interacts with the ring finger proteins RNF20 and RNF40 and is required for the maintenance of histone 2B monoubiquitination. <i>Human Molecular Genetics</i> , 2012, 21, 559-568.	1.4	85
20	A Detailed Clinicopathologic Study of ALK-translocated Papillary Thyroid Carcinoma. <i>American Journal of Surgical Pathology</i> , 2015, 39, 652-659.	2.1	85
21	Phosphaturic mesenchymal tumors show positive staining for somatostatin receptor 2A (SSTR2A). <i>Human Pathology</i> , 2013, 44, 2711-2718.	1.1	80
22	Parafibromin-deficient (HPT-JT Type, CDC73 Mutated) Parathyroid Tumors Demonstrate Distinctive Morphologic Features. <i>American Journal of Surgical Pathology</i> , 2019, 43, 35-46.	2.1	74
23	Loss of BAP1 expression is very rare in peritoneal and gynecologic serous adenocarcinomas and can be useful in the differential diagnosis with abdominal mesothelioma. <i>Human Pathology</i> , 2016, 51, 9-15.	1.1	72
24	Nuclear Accumulation of E-Cadherin Correlates with Loss of Cytoplasmic Membrane Staining and Invasion in Pituitary Adenomas. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2009, 94, 1436-1442.	1.8	71
25	Succinate Dehydrogenase Deficiency Is Rare in Pituitary Adenomas. <i>American Journal of Surgical Pathology</i> , 2014, 38, 560-566.	2.1	71
26	Loss of ARID1A expression in colorectal carcinoma is strongly associated with mismatch repair deficiency. <i>Human Pathology</i> , 2014, 45, 1697-1703.	1.1	61
27	Increased SSTR2A and SSTR3 expression in succinate dehydrogenase-deficient pheochromocytomas and paragangliomas. <i>Human Pathology</i> , 2015, 46, 390-396.	1.1	58
28	Inflammatory Myofibroblastic Tumors of the Female Genital Tract Are Under-recognized. <i>American Journal of Surgical Pathology</i> , 2017, 41, 1433-1442.	2.1	56
29	BRAFV600E immunohistochemistry in conjunction with mismatch repair status predicts survival in patients with colorectal cancer. <i>Modern Pathology</i> , 2014, 27, 644-650.	2.9	53
30	NTRK gene rearrangements are highly enriched in MLH1/PMS2 deficient, BRAF wild-type colorectal carcinomas—a study of 4569 cases. <i>Modern Pathology</i> , 2020, 33, 924-932.	2.9	51
31	When used together SS18-specific and SSX C-terminus immunohistochemistry are highly specific and sensitive for the diagnosis of synovial sarcoma and can replace FISH or molecular testing in most cases. <i>Histopathology</i> , 2020, 77, 588-600.	1.6	50
32	Loss of BAP1 Expression Occurs Frequently in Intrahepatic Cholangiocarcinoma. <i>Medicine (United States)</i> , 2018, 97, 1048-1054.	0.4	48
33	Succinate dehydrogenase-deficient GISTs are characterized by IGF1R overexpression. <i>Modern Pathology</i> , 2012, 25, 1307-1313.	2.9	46
34	A Proposed Grading Scheme for Medullary Thyroid Carcinoma Based on Proliferative Activity (Ki-67). <i>Thyroid</i> , 2015, 25, 1419-1428.	2.1	46
35	Medullary Colorectal Carcinoma Revisited: A Clinical and Pathological Study of 102 Cases. <i>Annals of Surgical Oncology</i> , 2015, 22, 2988-2996.	0.7	45
36	ATRX loss is an independent predictor of poor survival in pancreatic neuroendocrine tumors. <i>Human Pathology</i> , 2018, 82, 249-257.	1.1	42

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37	Mismatch repair deficiency as a prognostic factor in mucinous colorectal cancer. <i>Modern Pathology</i> , 2016, 29, 266-274.	2.9	39
38	Immunohistochemistry for Myc Predicts Survival in Colorectal Cancer. <i>PLoS ONE</i> , 2014, 9, e87456.	1.1	38
39	Loss of expression of BAP1 is very rare in non-small cell lung carcinoma. <i>Pathology</i> , 2016, 48, 336-340.	0.3	35
40	Lessons learnt from implementation of a Lynch syndrome screening program for patients with gynaecological malignancy. <i>Pathology</i> , 2017, 49, 457-464.	0.3	34
41	Reflex ALK immunohistochemistry is feasible and highly specific for ALK gene rearrangements in lung cancer. <i>Pathology</i> , 2014, 46, 383-388.	0.3	32
42	NRASQ61R Mutation-specific Immunohistochemistry Also Identifies the HRASQ61R Mutation in Medullary Thyroid Cancer and May Have a Role in Triaging Genetic Testing for MEN2. <i>American Journal of Surgical Pathology</i> , 2017, 41, 75-81.	2.1	31
43	ALK and ROS1 Overexpression is Very Rare in Colorectal Adenocarcinoma. <i>Applied Immunohistochemistry and Molecular Morphology</i> , 2015, 23, 134-138.	0.6	27
44	Elevated serum FGF23 concentrations in plasma cell dyscrasias. <i>Bone</i> , 2006, 39, 369-376.	1.4	26
45	The RING finger domain E3 ubiquitin ligases BRCA1 and the RNF20/RNF40 complex in global loss of the chromatin mark histone H2B monoubiquitination (H2Bub1) in cell line models and primary high-grade serous ovarian cancer. <i>Human Molecular Genetics</i> , 2016, 25, ddd362.	1.4	26
46	Immunoregulatory Forkhead Box Protein p3-Positive Lymphocytes Are Associated with Overall Survival in Patients with Pancreatic Neuroendocrine Tumors. <i>Journal of the American College of Surgeons</i> , 2016, 222, 281-287.	0.2	24
47	Loss of INI1 expression in colorectal carcinoma is associated with high tumor grade, poor survival, BRAFV600E mutation, and mismatch repair deficiency. <i>Human Pathology</i> , 2016, 55, 83-90.	1.1	20
48	Expanding the clinicopathological spectrum of succinate dehydrogenase-deficient renal cell carcinoma with a focus on variant morphologies: a study of 62 new tumors in 59 patients. <i>Modern Pathology</i> , 2022, 35, 836-849.	2.9	20
49	Loss of Hes1 expression is associated with poor prognosis in colorectal adenocarcinoma. <i>Human Pathology</i> , 2016, 57, 91-97.	1.1	19
50	The epithelioid BAP1-negative and p16-positive phenotype predicts prolonged survival in pleural mesothelioma. <i>Histopathology</i> , 2018, 72, 509-515.	1.6	17
51	Mutation specific immunohistochemistry is highly specific for the presence of calreticulin mutations in myeloproliferative neoplasms. <i>Pathology</i> , 2016, 48, 319-324.	0.3	15
52	Utility of GATA-3 Expression in the Analysis of Pituitary Neuroendocrine Tumour (PitNET) Transcription Factors. <i>Endocrine Pathology</i> , 2020, 31, 150-155.	5.2	15
53	A Further Investigation of Combined Mismatch Repair and BRAFV600E Mutation Specific Immunohistochemistry as a Predictor of Overall Survival in Colorectal Carcinoma. <i>PLoS ONE</i> , 2014, 9, e106105.	1.1	13
54	PD-L1 Is Preferentially Expressed in PIT-1 Positive Pituitary Neuroendocrine Tumours. <i>Endocrine Pathology</i> , 2021, 32, 408-414.	5.2	12

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55	Loss of BAP1 Expression Is Very Rare in Pancreatic Ductal Adenocarcinoma. PLoS ONE, 2016, 11, e0150338.	1.1	11
56	NRASQ61R Mutation-specific Immunohistochemistry is Highly Specific for Either NRAS Q61R or KRAS Q61R Mutation in Colorectal Carcinoma. Applied Immunohistochemistry and Molecular Morphology, 2017, 25, 475-480.	0.6	11
57	EGFR mutation specific immunohistochemistry is a useful adjunct which helps to identify false negative mutation testing in lung cancer. Pathology, 2014, 46, 501-508.	0.3	9
58	Switch/sucrose non-fermentable (SWI/SNF) complex (SMARCA4), Tj ETQqO O O rgBT /Overlock strongly associated with microsatellite instability: an incidence study in 4508 colorectal carcinomas. Histopathology, 2022, 80, 906-921.	1.6	9
59	DNA damage-inducible transcript 3 immunohistochemistry is highly sensitive for the diagnosis of myxoid liposarcoma but care is required in interpreting the significance of focal expression. Histopathology, 2021, 79, 106-116.	1.6	8
60	BRAF V600E mutation specific immunohistochemistry with clone VE1 is not reliable in pituitary adenomas. Pathology, 2014, 46, 79-80.	0.3	7
61	Real world experience of BRAFV600E mutation specific immunohistochemistry in colorectal carcinoma. Pathology, 2018, 50, 342-344.	0.3	6
62	Immunohistochemical expression of somatostatin receptors SSTR2A and SSTR5 in 299 pituitary adenomas. Pathology, 2018, 50, 472-474.	0.3	4
63	Immunohistochemistry for PMS2 and MSH6 alone can replace a four antibody panel for mismatch repair deficiency screening in colorectal adenocarcinoma: authors' reply. Pathology, 2011, 43, 85-86.	0.3	0