Kathrin Ludwig

List of Publications by Year
in descending order

27	539	933447 10	642732 23
papers	citations	h-index	g-index

28	28	28	1066
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	â€œWhile there is p57, there is hope.â€•The past and the present of diagnosis in first trimester abortions: Diagnostic dilemmas and algorithmic approaches. A review. Placenta, 2021, 116, 31-37.	1.5	5
2	Photosensitive epilepsy and long QT: expanding Timothy syndrome phenotype. Clinical Neurophysiology, 2019, 130, 2134-2136.	1.5	6
3	The Anatomy and Histology of theÂLiver and Biliary Tract. , 2019, , 41-55.		2
4	Diagnostic utility of cyclin D1 in the diagnosis of small round blue cell tumors in children and adolescents: beware of cyclin D1 expression in clear cell sarcoma of the kidney and CIC-DUX4 fusionâ€“positive sarcomas. Comment on Magro et al (2016)â€”reply. Human Pathology, 2017, 67, 226-228.	2.0	2
5	BCOR-CCNB3 Undifferentiated Sarcomaâ€”Does Immunohistochemistry Help in the Identification?. Pediatric and Developmental Pathology, 2017, 20, 321-329.	1.0	38
6	Antiâ€Glypican 3, a Novel Ancillary Maker in the Histological Assessment of Hirschsprung's Disease. Journal of Pediatric Gastroenterology and Nutrition, 2016, 62, 692-697.	1.8	2
7	Recurrent BCOR Internal Tandem Duplication and YWHAE-NUTM2B Fusions in Soft Tissue Undifferentiated Round Cell Sarcoma of Infancy. American Journal of Surgical Pathology, 2016, 40, 1009-1020.	3.7	155
8	Molecular Cytogenetics Detect an Unbalanced t(2;13)(q36;q14) and PAX3 -FOXO1 Fusion in Rhabdomyosarcoma With Mixed Embryonal/Alveolar Features. Pediatric Blood and Cancer, 2015, 62, 2238-2241.	1.5	5
9	Oral-facial-digital syndrome type VI: is C5orf42 really the major gene?. Human Genetics, 2015, 134, 123-126.	3.8	30
10	Ichthyosis and Kallmann syndrome: Not always a contiguous gene syndrome. Journal of Dermatological Science, 2015, 78, 158-160.	1.9	4
11	Omental mesenteric myxoid hamartoma, a subtype of inflammatory myofibroblastic tumor? Considerations based on the histopathological evaluation of four cases. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2015, 467, 741-747.	2.8	7
12	Rare uterine cancer: Carcinosarcomas. Review from histology to treatment. Critical Reviews in Oncology/Hematology, 2015, 94, 98-104.	4.4	44
13	â€œDouble Troubleâ€•or an Amplification of the Triploidy Phenotype?. Fetal and Pediatric Pathology, 2013, 32, 60-65.	0.7	2
14	PDCD4/miR-21 dysregulation in inflammatory bowel disease-associated carcinogenesis. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2013, 462, 57-63.	2.8	55
15	Comment on â€œDeterminants of Lymph Node Count in Endometrial Cancer Surgical Stagingâ€• International Journal of Gynecological Cancer, 2013, 23, 588.	2.5	0
16	Endometrial Polyps in Women Affected by Levothyroxine-Treated Hypothyroidismâ€”Histological Features, Immunohistochemical Findings, and Possible Explanation of Etiopathogenic Mechanism: A Pilot Study. BioMed Research International, 2013, 2013, 1-5.	1.9	14
17	Apert Syndrome with Fused Thalami. Fetal and Pediatric Pathology, 2012, 31, 410-414.	0.7	7
18	Congenital Pulmonary Airway Malformation (CPAM) [Congenital Cystic Adenomatoid Malformation] Associated with Tracheoesophageal Fistula and Agensesis of the Corpus Callosum. Fetal and Pediatric Pathology, 2012, 31, 169-175.	0.7	5

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19	Human epithelial growth factor receptor 2 (HER2) status in primary and metastatic esophagogastric junction adenocarcinomas. Human Pathology, 2012, 43, 1206-1212.	2.0	34
20	Pentalogy of Cantrell with Complete Ectopia Cordis in a Fetus with Asplenia. Pediatric and Developmental Pathology, 2012, 15, 495-498.	1.0	8
21	Cervical Follicular Dendritic Cell Sarcoma: A Case Report and Review of the Literature. International Journal of Immunopathology and Pharmacology, 2011, 24, 539-544.	2.1	11
22	PDCD4 nuclear loss inversely correlates with miR-21 levels in colon carcinogenesis. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2011, 458, 413-419.	2.8	72
23	Dandy-Walker Malformation Masking the Molar Tooth Sign: An Illustrative Case With Magnetic Resonance Imaging Follow-up. Journal of Child Neurology, 2010, 25, 1419-1422.	1.4	14
24	A CASE OF FEMUR-FIBULAR-ULNA COMPLEX WITH PECULIAR METAPHYSEAL CHANGES. Fetal and Pediatric Pathology, 2010, 29, 255-260.	0.7	0
25	Tracheal Agenesis with Bifurcating Common Airway Arising from Midesophagus. Pediatric and Developmental Pathology, 2010, 13, 252-254.	1.0	4
26	A case of diploid/triploid mosaicism with dental Blaschko lines. Clinical Dysmorphology, 2009, 18, 232-233.	0.3	3
27	The Impact of Single Nucleotide Polymorphisms of the Thrombin Activatable Fibrinolysis Inhibitor (TAFI) Gene on TAFI Antigen Levels in Healthy Children and Pediatric Oncology Patients. Seminars in Thrombosis and Hemostasis, 2003, 29, 575-584.	2.7	8