

# Hela Fourati

## List of Publications by Year in descending order

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Version: 2024-02-01

21  
papers

176  
citations

1307594

7  
h-index

1199594

12  
g-index

35  
all docs

35  
docs citations

35  
times ranked

272  
citing authors

#	ARTICLE	IF	CITATIONS
1	Suprasellar Melanocytoma with Leptomeningeal Seeding: An Aggressive Clinical Course for a Histologically Benign Tumor. <i>Case Reports in Radiology</i> , 2021, 2021, 1-7.	0.3	1
2	Ontology-Based Approach for Liver Cancer Diagnosis and Treatment. <i>Journal of Digital Imaging</i> , 2019, 32, 116-130.	2.9	15
3	Factors Predicting Lung Contusions in Critically Ill Trauma Children. <i>Pediatric Emergency Care</i> , 2018, 34, 198-201.	0.9	7
4	CREM variant rs17583959 conferred susceptibility to T1D risk in the Tunisian families. <i>Immunology Letters</i> , 2017, 181, 1-5.	2.5	4
5	MRI features in 17 patients with l2 hydroxyglutaric aciduria. <i>European Journal of Radiology Open</i> , 2016, 3, 245-250.	1.6	23
6	Paucisymptomatic Dermoid Cyst with Fatal Outcome. <i>Pediatric Dermatology</i> , 2016, 33, e333-6.	0.9	2
7	Occipital dermoid cyst associated with dermal sinus complicated with meningitis: A case report. <i>Archives De Pediatrie</i> , 2016, 23, 197-200.	1.0	7
8	Unrepaired persistent truncus arteriosus in a 38-year-old woman with an uneventful pregnancy. <i>Cardiovascular Journal of Africa</i> , 2015, 26, e6-e8.	0.4	7
9	Presacral myelolipoma: Imaging features. <i>Presse Medicale</i> , 2015, 44, 1196-1198.	1.9	6
10	Masse h�patique unique et isol�e chez une femme �g�e. <i>Feuillets De Radiologie</i> , 2015, 55, 247-252.	0.0	0
11	Association of TCR/CD3, PTPN22, CD28 and ZAP70 gene polymorphisms with type 1 diabetes risk in Tunisian population: Family based association study. <i>Immunology Letters</i> , 2015, 163, 1-7.	2.5	19
12	Neurofibromatose de type 1 associ�e � un syndrome de Moya-Moya. <i>Archives De Pediatrie</i> , 2015, 22, 106-108.	1.0	0
13	Cholestasis and protein-losing enteropathy secondary to hyperthyroidism in a 6-year-old girl. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2014, 27, 1017-9.	0.9	2
14	Early-onset of multiple sclerosis in a 5-year-old girl. <i>Archives De Pediatrie</i> , 2014, 21, 291-295.	1.0	2
15	Actinomycotic brain abscess as the first clinical manifestation of hereditary hemorrhagic telangiectasia� case report and review of the literature. <i>Annals of Hematology</i> , 2013, 92, 1141-1143.	1.8	7
16	Resistant Invasive Aspergillosis in an Autosomal Recessive Chronic Granulomatous Disease. <i>Fetal and Pediatric Pathology</i> , 2013, 32, 241-245.	0.7	3
17	Arterial ischemic stroke in children: 22 cases from southern Tunisia. <i>Fetal and Pediatric Pathology</i> , 2013, 32, 271-275.	0.7	3
18	Brain magnetic resonance imaging findings in adult patients with congenital adrenal hyperplasia: Increased frequency of white matter impairment and temporal lobe structures dysgenesis. <i>Indian Journal of Endocrinology and Metabolism</i> , 2013, 17, 121.	0.4	23

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19	Complete endoscopic management of tubular esophageal duplication in a young woman. <i>Endoscopy</i> , 2012, 44, E261-E262.	1.8	7
20	Leydig Cell Tumor Associated with Testicular Adrenal Rest Tumors in a Patient with Congenital Adrenal Hyperplasia due to 11 $\beta$ -Hydroxylase Deficiency. <i>Case Reports in Urology</i> , 2012, 2012, 1-5.	0.3	12
21	HLA-DRB1/DQB1 susceptibility for autoimmune polyglandular syndrome type II and III in south of Tunisia. <i>Annales D'Endocrinologie</i> , 2011, 72, 232-238.	1.4	6