

# Abdallah Fayssol

## List of Publications by Year in descending order

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76  
papers

972  
citations

566801

15  
h-index

500791

28  
g-index

84  
all docs

84  
docs citations

84  
times ranked

1846  
citing authors

#	ARTICLE	IF	CITATIONS
1	Development and Validation of a New Risk Prediction Score for Life-Threatening Ventricular Tachyarrhythmias in Laminopathies. <i>Circulation</i> , 2019, 140, 293-302.	1.6	131
2	Cardiomyopathy in Duchenne muscular dystrophy: pathogenesis and therapeutics. <i>Heart Failure Reviews</i> , 2010, 15, 103-107.	1.7	89
3	Diaphragm: Pathophysiology and Ultrasound Imaging in Neuromuscular Disorders. <i>Journal of Neuromuscular Diseases</i> , 2018, 5, 1-10.	1.1	57
4	Prediction of long-term prognosis by heteroplasmy levels of the m.3243A>G mutation in patients with the <scp>mitochondrial encephalomyopathy, lactic acidosis and stroke-like episodes</scp> syndrome. <i>European Journal of Neurology</i> , 2017, 24, 255-261.	1.7	41
5	Cardiac Involvement Classification and Therapeutic Management in Patients with Duchenne Muscular Dystrophy. <i>Journal of Neuromuscular Diseases</i> , 2017, 4, 17-23.	1.1	39
6	Association Between Mutation Size and Cardiac Involvement in Myotonic Dystrophy Type 1. <i>Circulation: Cardiovascular Genetics</i> , 2017, 10, .	5.1	39
7	Natural History of Cardiac and Respiratory Involvement, Prognosis and Predictive Factors for Long-Term Survival in Adult Patients with Limb Girdle Muscular Dystrophies Type 2C and 2D. <i>PLoS ONE</i> , 2016, 11, e0153095.	1.1	36
8	Development and Validation of a New Scoring System to Predict Survival in Patients With Myotonic Dystrophy Type 1. <i>JAMA Neurology</i> , 2018, 75, 573.	4.5	32
9	Torsade de pointes induced by citalopram and amiodarone. <i>Annales De Cardiologie Et D'Angiologie</i> , 2011, 60, 165-168.	0.3	29
10	Heart Diseases in Mitochondrial Encephalomyopathy, Lactic Acidosis, and Stroke Syndrome. <i>Congestive Heart Failure</i> , 2009, 15, 284-287.	2.0	25
11	Diaphragm sniff ultrasound: Normal values, relationship with sniff nasal pressure and accuracy for predicting respiratory involvement in patients with neuromuscular disorders. <i>PLoS ONE</i> , 2019, 14, e0214288.	1.1	25
12	Association between prophylactic angiotensin-converting enzyme inhibitors and overall survival in Duchenne muscular dystrophy—analysis of registry data. <i>European Heart Journal</i> , 2021, 42, 1976-1984.	1.0	25
13	Atrial flutter in myotonic dystrophy type 1: Patient characteristics and clinical outcome. <i>Neuromuscular Disorders</i> , 2016, 26, 227-233.	0.3	19
14	Cardiac Characterization of mdx Mice Using High-Resolution Doppler Echocardiography. <i>Journal of Ultrasound in Medicine</i> , 2013, 32, 757-761.	0.8	19
15	Cardiomyopathy in Pompe's disease. <i>European Journal of Internal Medicine</i> , 2008, 19, 57-59.	1.0	18
16	Complete atrioventricular block in Duchenne muscular dystrophy. <i>Europace</i> , 2008, 10, 1351-1352.	0.7	15
17	Percutaneous Extracorporeal Membrane Oxygenation for Cardiogenic Shock Due to Acute Fulminant Myocarditis. <i>Annals of Thoracic Surgery</i> , 2010, 89, 614-616.	0.7	15
18	Natural history of cardiac function in Duchenne and Becker muscular dystrophies on home mechanical ventilation. <i>Medicine (United States)</i> , 2018, 97, e11381.	0.4	15

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19	Clinical profiles and prognosis of acute heart failure in adult patients with dystrophinopathies on home mechanical ventilation. <i>ESC Heart Failure</i> , 2017, 4, 527-534.	1.4	14
20	Analyzing left ventricular function in mice with Doppler echocardiography. <i>Heart Failure Reviews</i> , 2013, 18, 511-516.	1.7	13
21	Successful cardiac resynchronisation therapy in Duchenne muscular dystrophy: A 5-year follow-up. <i>Presse Medicale</i> , 2014, 43, 330-331.	0.8	13
22	Neonatal COVID-19 Pneumonia: Report of the First Case in a Preterm Neonate in Mayotte, an Overseas Department of France. <i>Children</i> , 2020, 7, 87.	0.6	13
23	CD38â€NADase is a new major contributor to Duchenne muscular dystrophic phenotype. <i>EMBO Molecular Medicine</i> , 2022, 14, e12860.	3.3	13
24	The Right Ventricle in COVID-19 Patients. <i>American Journal of Cardiology</i> , 2020, 130, 166-167.	0.7	12
25	Cardiac diseases in sarcoglycanopathies. <i>International Journal of Cardiology</i> , 2010, 144, 67-68.	0.8	11
26	Cardiac implantable electronic devices in tracheotomized muscular dystrophy patients: Safety and risks. <i>International Journal of Cardiology</i> , 2016, 222, 975-977.	0.8	11
27	Long term longitudinal study of muscle function in patients with glycogen storage disease type IIIa. <i>Molecular Genetics and Metabolism</i> , 2017, 122, 108-116.	0.5	11
28	Echographic Assessment of Diaphragmatic Function in Duchenne Muscular Dystrophy from Childhood to Adulthood. <i>Journal of Neuromuscular Diseases</i> , 2019, 6, 55-64.	1.1	11
29	Cardiac Characterization of <i>mdx</i> Mice Using High-Resolution Doppler Echocardiography. <i>Journal of Ultrasound in Medicine</i> , 2013, 32, 757-761.	0.8	10
30	Pacemaker Implantation for Sinus Node Dysfunction in a Young Patient With Duchenne Muscular Dystrophy. <i>Congestive Heart Failure</i> , 2010, 16, 127-128.	2.0	9
31	Masseter muscle oxygen saturation is associated with central venous oxygen saturation in patients with severe sepsis. <i>Journal of Clinical Monitoring and Computing</i> , 2010, 24, 289-293.	0.7	9
32	Cardiac asynchrony in Duchenne muscular dystrophy. <i>Journal of Clinical Monitoring and Computing</i> , 2013, 27, 587-589.	0.7	9
33	1st International Workshop on Clinical trial readiness for sarcoglycanopathies 15â€“16 November 2016, Evry, France. <i>Neuromuscular Disorders</i> , 2017, 27, 683-692.	0.3	9
34	Renâ€© Laennec (1781â€“1826) and the Invention of the Stethoscope. <i>American Journal of Cardiology</i> , 2009, 104, 743-744.	0.7	8
35	Heart and anorexia nervosa. <i>Heart Failure Reviews</i> , 2021, 26, 65-70.	1.7	8
36	Nutritional status, swallowing disorders, and respiratory prognosis in adult Duchenne muscular dystrophy patients. <i>Pediatric Pulmonology</i> , 2021, 56, 2146-2154.	1.0	7

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37	Diastolic function in Steinert's disease. <i>Neurology International</i> , 2014, 6, 5140.	1.3	6
38	Be careful about abdominal discomfort in adult patients with muscular dystrophy. <i>Revue Neurologique</i> , 2014, 170, 548-550.	0.6	6
39	Left bundle branch block in Duchenne muscular dystrophy: Prevalence, genetic relationship and prognosis. <i>PLoS ONE</i> , 2018, 13, e0190518.	1.1	6
40	Right ventricular function in late-onset Pompe disease. <i>Journal of Clinical Monitoring and Computing</i> , 2014, 28, 419-421.	0.7	5
41	Left ventricular function in alpha-sarcoglycanopathy and gamma-sarcoglycanopathy. <i>Acta Neurologica Belgica</i> , 2014, 114, 257-259.	0.5	5
42	Risk for Complications after Pacemaker or Cardioverter Defibrillator Implantations in Patients with Myotonic Dystrophy Type 1. <i>Journal of Neuromuscular Diseases</i> , 2017, 4, 175-181.	1.1	5
43	Hypertrophic cardiomyopathy in Friedreich's ataxia. <i>International Journal of Cardiology</i> , 2008, 127, e122-e123.	0.8	4
44	Impact of Angiotensin-Converting Enzyme Inhibitors and Angiotensin II Receptor Blockers in Hypertensive Patients with COVID-19 (COVIDECA Study). <i>American Journal of Cardiology</i> , 2021, 147, 58-60.	0.7	4
45	Diaphragm Ultrasound in Cardiac Surgery: State of the Art. <i>Medicines (Basel, Switzerland)</i> , 2022, 9, 5.	0.7	4
46	Holter electrocardiogram should be systematic in Duchenne muscular dystrophy. <i>International Journal of Cardiology</i> , 2008, 128, 442-443.	0.8	3
47	Tissue Doppler characterization of cardiac phenotype in mouse. <i>European Journal of Radiology</i> , 2009, 72, 82-84.	1.2	3
48	Sudden death in Steinert's disease. <i>International Journal of Cardiology</i> , 2009, 136, e27-e28.	0.8	3
49	Left Atrial Volume Index: A Predictor of Adverse Outcome in Patients With Hypertrophic Cardiomyopathy. <i>Journal of the American Society of Echocardiography</i> , 2010, 23, 456.	1.2	3
50	Usefulness of myocardial strain imaging in Duchenne muscular dystrophy. <i>International Journal of Cardiology</i> , 2010, 140, 114-115.	0.8	3
51	Wolff-Parkinson-White syndrome in Duchenne muscular dystrophy. <i>International Journal of Cardiology</i> , 2013, 167, e53-e54.	0.8	3
52	Cardiac characterization of sgca-null mice using high resolution echocardiography. <i>Neurology International</i> , 2013, 5, 22.	1.3	3
53	Accuracy of B-natriuretic peptide for the diagnosis of decompensated heart failure in muscular dystrophies patients with chronic respiratory failure. <i>Neurology International</i> , 2018, 10, 7917.	1.3	3
54	Assessment of diaphragm motion using ultrasonography in a patient with facio-scapulo-humeral dystrophy. <i>Medicine (United States)</i> , 2019, 98, e13887.	0.4	3

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55	Letter by Faysoil Regarding Article, "Excess Mortality Associated With Functional Tricuspid Regurgitation Complicating Heart Failure With Reduced Ejection Fraction" Circulation, 2020, 141, e1-e2.	1.6	3
56	Determinants of diaphragm inspiratory motion, diaphragm thickening, and its performance for predicting respiratory restrictive pattern in <scp>Duchenne</scp> muscular dystrophy. Muscle and Nerve, 2022, 65, 89-95.	1.0	3
57	Should we perform systematic electrophysiological study in Steinert's disease?. Journal of Cardiothoracic Surgery, 2008, 3, 56.	0.4	2
58	Prognostic Implications of the Doppler Restrictive Filling Pattern in Hypertrophic Cardiomyopathy. American Journal of Cardiology, 2010, 105, 1358-1359.	0.7	2
59	Effects of Home Mechanical Ventilation on Left Ventricular Function in Sarcoglycanopathies (Limb) Tj ETQq1 1 0.784314 rgBJ /Overlock	0.7	2
60	M Mode Ultrasound and Tissue Doppler Imaging to Assess Diaphragm Feature in Late Onset Pompe Disease. Neurology International, 2020, 12, 55-58.	1.3	2
61	Acute ischemic stroke in gamma-sarcoglycanopathy. Presse Medicale, 2013, 42, 484-486.	0.8	1
62	Left ventricular aneurysm in a patient with Duchenne muscular dystrophy. Presse Medicale, 2014, 43, 731-732.	0.8	1
63	Revisited the place for B-type natriuretic peptide in patients with muscular dystrophy in the area of mechanical ventilation. International Journal of Cardiology, 2018, 257, 315.	0.8	1
64	Diaphragm dysfunction after cardiac surgery: a global approach. Journal of Clinical Monitoring and Computing, 2020, 34, 615-615.	0.7	1
65	Analysis of inspiratory and expiratory muscles using ultrasound in rats: A reproducible and non-invasive tool to study respiratory function. Respiratory Physiology and Neurobiology, 2021, 285, 103596.	0.7	1
66	Cardiopulmonary Pathophysiological Aspects in the Context of COVID-19 and Obesity. SN Comprehensive Clinical Medicine, 2021, 3, 1848-1857.	0.3	1
67	Leadless intracardiac transcatheter pacing system: 20 months follow up in adult Duchenne muscular dystrophy. Neuromuscular Disorders, 2021, 31, 896-898.	0.3	1
68	Diaphragm ultrasound to stratify <scp>COVID</scp>"19 patients in the emergency department?. Journal of Clinical Ultrasound, 2022, 50, 106-107.	0.4	1
69	High parasternal intercostal muscle thickening prior to intubation in COVID-19 infection. Radiology Case Reports, 2022, 17, 843-846.	0.2	1
70	Aldosterone Antagonists in Patients With Heart Failure. JAMA - Journal of the American Medical Association, 2010, 303, 833.	3.8	0
71	Letter by Faysoil Regarding Article, "Determinants of Surgical Outcome in Patients With Isolated Tricuspid Regurgitation" Circulation, 2010, 122, e13; author reply e15.	1.6	0
72	Letter by Faysoil Regarding Article, "Early Treatment With Lisinopril and Spironolactone Preserves Cardiac and Skeletal Muscle in Duchenne Muscular Dystrophy Mice" Circulation, 2012, 125, e429.	1.6	0

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73	Right ventricular function in Steinert's disease. International Journal of Cardiology, 2013, 167, 291.	0.8	0
74	Which place for ivabradine in Duchenne muscular dystrophy with heart failure?. International Journal of Cardiology, 2016, 223, 768-769.	0.8	0
75	Focus on left ventricular systolic and diastolic function in the assisted 6â€­minute hand bike cycle test in muscular dystrophy. Muscle and Nerve, 2019, 59, E47.	1.0	0
76	Managing advanced heart failure in Duchenne muscular dystrophy. Progress in Pediatric Cardiology, 2020, 56, 101148.	0.2	0