

# Silvia Favilli

## List of Publications by Year in descending order

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Version: 2024-02-01

69  
papers

834  
citations

687363

13  
h-index

552781

26  
g-index

80  
all docs

80  
docs citations

80  
times ranked

1109  
citing authors

#	ARTICLE	IF	CITATIONS
1	Clinical risks of beta-blockers in galenic preparation in children. <i>Minerva Pediatrics</i> , 2023, 75, .	0.4	1
2	Diagnosis and Management of Cardiovascular Involvement in Friedreich Ataxia. <i>Heart Failure Clinics</i> , 2022, 18, 31-37.	2.1	12
3	Eligibility criteria for pediatric patients who may benefit from anti SARS-CoV-2 monoclonal antibody therapy administration: an Italian inter-society consensus statement. <i>Italian Journal of Pediatrics</i> , 2022, 48, 7.	2.6	9
4	Multimodality imaging in complex aortic arch anomaly. <i>European Heart Journal - Case Reports</i> , 2022, 6, ytac048.	0.6	0
5	Diagnosis and Management of Rare Cardiomyopathies in Adult and Paediatric Patients. A Position Paper of the Italian Society of Cardiology (SIC) and Italian Society of Paediatric Cardiology (SICP). <i>International Journal of Cardiology</i> , 2022, 357, 55-71.	1.7	36
6	Prescribing, dosing and titrating exercise in patients with hypertrophic cardiomyopathy for prevention of comorbidities: Ready for prime time. <i>European Journal of Preventive Cardiology</i> , 2021, 28, 1093-1099.	1.8	15
7	Long-term follow-up of coronary artery lesions in children in Kawasaki syndrome. <i>European Journal of Pediatrics</i> , 2021, 180, 271-275.	2.7	6
8	Two-Dimensional Aortic Size Normalcy: A Novelty Detection Approach. <i>Diagnostics</i> , 2021, 11, 220.	2.6	4
9	Supraventricular tachycardias in the first year of life: what is the best pharmacological treatment? 24Åyears of experience in a single centre. <i>BMC Cardiovascular Disorders</i> , 2021, 21, 137.	1.7	8
10	Neonatal heart failure and noncompaction/dilated cardiomyopathy from mucopolysaccharidosis. First description in literature. <i>Molecular Genetics and Metabolism Reports</i> , 2021, 26, 100714.	1.1	2
11	Fast recovery of cardiac function in PIMS-TS patients early using intravenous anti-IL-1 treatment. <i>Critical Care</i> , 2021, 25, 131.	5.8	12
12	Pathophysiology and clinical presentation of paediatric heart failure related to congenital heart disease. <i>Acta Paediatrica, International Journal of Paediatrics</i> , 2021, 110, 2336-2343.	1.5	2
13	1â€...The role of the electrocardiographic phenotype in risk stratification for sudden cardiac death in childhood hypertrophic cardiomyopathy. , 2021, , .		2
14	Case Report: Perioperative Kounis Syndrome in an Adolescent With Congenital Glaucoma. <i>Frontiers in Cardiovascular Medicine</i> , 2021, 8, 676188.	2.4	2
15	Differential Diagnosis between Marfan Syndrome and Loey'sâ€Dietz Syndrome Type 4: A Novel Chromosomal Deletion Covering TGFB2. <i>Genes</i> , 2021, 12, 1462.	2.4	2
16	Clinical presentation and longâ€term outcomes of infantile hypertrophic cardiomyopathy: a European multicentre study. <i>ESC Heart Failure</i> , 2021, 8, 5057-5067.	3.1	22
17	Resilience and response of the congenital cardiac network in Italy during the COVID-19 pandemic. <i>Journal of Cardiovascular Medicine</i> , 2021, 22, 9-13.	1.5	7
18	Kounis Syndrome Associated With Takotsubo Syndrome in an Adolescent With Peutz-Jeghers Syndrome. <i>JACC: Case Reports</i> , 2021, 3, 1602-1606.	0.6	4

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19	The Influence of Genotype on the Phenotype, Clinical Course, and Risk of Adverse Events in Children with Hypertrophic Cardiomyopathy. <i>Heart Failure Clinics</i> , 2021, 18, 1-8.	2.1	1
20	Prevalence of Inherited Cardiac Diseases Among Young Patients Requiring Permanent Pacing. <i>Circulation: Arrhythmia and Electrophysiology</i> , 2021, 14, CIRCEP121010562.	4.8	6
21	Impact of hard lockdown on interventional cardiology procedures in congenital heart disease: a survey on behalf of the Italian Society of Congenital Heart Disease. <i>Journal of Cardiovascular Medicine</i> , 2021, 22, 701-705.	1.5	3
22	Combined Surgical and Endoscopic Approach for Ringâ€“Sling Complex. <i>Thoracic and Cardiovascular Surgeon</i> , 2020, 68, 051-058.	1.0	2
23	Frequent Ventricular Premature Beats in Children and Adolescents: Natural History and Relationship with Sport Activity in a Long-Term Follow-Up. <i>Pediatric Cardiology</i> , 2020, 41, 123-128.	1.3	11
24	Incessant Automatic Atrial Tachycardia in a Neonate Successfully Treated with Nadolol and Closely Spaced Doses of Flecainide: A Case Report. <i>Pediatric Reports</i> , 2020, 12, 108-113.	1.3	3
25	Impact of cardiovascular involvement on the clinical course of paediatric mitochondrial disorders. <i>Orphanet Journal of Rare Diseases</i> , 2020, 15, 196.	2.7	8
26	A rare case of pediatric cardiomyopathy: AlstrÃ¶m syndrome identified by gene panel analysis. <i>Clinical Case Reports (discontinued)</i> , 2020, 8, 3369-3373.	0.5	2
27	Advances in Stem Cell Modeling of Dystrophin-Associated Disease: Implications for the Wider World of Dilated Cardiomyopathy. <i>Frontiers in Physiology</i> , 2020, 11, 368.	2.8	9
28	Kounis syndrome: a clinical entity penetrating from pediatrics to geriatrics. <i>Journal of Geriatric Cardiology</i> , 2020, 17, 294-299.	0.2	10
29	Kounis Syndrome: a pediatric perspective. <i>Minerva Pediatrica</i> , 2020, 72, 383-392.	2.7	12
30	Lifestyles and Cardiovascular Prevention in Childhood and Adolescence. <i>Pediatric Cardiology</i> , 2019, 40, 1113-1125.	1.3	54
31	Rare X-linked storage heart diseases are tougher on men but not kind to women. <i>International Journal of Cardiology</i> , 2019, 286, 113-114.	1.7	0
32	Comment on: Assessment of cardiac disease in MELAS requires comprehensive, prospective work-up. <i>International Journal of Cardiology</i> , 2019, 280, 162.	1.7	0
33	SAT0505â€“...LONG-TERM FOLLOW-UP IN KAWASAKI SYNDROME: EVIDENCE FROM RETROSPECTIVE MONOCENTRIC DATA IN REAL LIFE. , 2019, , .		0
34	Clinical profile and outcome of cardiac involvement in MELAS syndrome. <i>International Journal of Cardiology</i> , 2019, 276, 14-19.	1.7	21
35	Long-term Outcomes of Pediatric-Onset Hypertrophic Cardiomyopathy and Age-Specific Risk Factors for Lethal Arrhythmic Events. <i>JAMA Cardiology</i> , 2018, 3, 520.	6.1	78
36	Profiles of heart failure in adolescents and young adults with congenital heart disease. <i>Progress in Pediatric Cardiology</i> , 2018, 51, 37-45.	0.4	5

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37	Genetic testing in pediatric cardiomyopathies: Implications for diagnosis and management. <i>Progress in Pediatric Cardiology</i> , 2018, 51, 24-30.	0.4	3
38	Cardiomyopathies in children – inherited heart muscle disease. <i>Progress in Pediatric Cardiology</i> , 2018, 51, 8-15.	0.4	1
39	Age-related issues: From fetus to adolescent. <i>Progress in Pediatric Cardiology</i> , 2018, 51, 3-7.	0.4	2
40	Consensus Document of the Italian Association of Hospital Cardiologists (ANMCO), Italian Society of Pediatric Cardiology (SICP), and Italian Society of Gynaecologists and Obstetrics (SIGO): pregnancy and congenital heart diseases. <i>European Heart Journal Supplements</i> , 2017, 19, D256-D292.	0.1	13
41	Aortic Arch Interruption and Persistent Fifth Aortic Arch in Phace Syndrome: Prenatal Diagnosis and Postnatal Course. <i>Echocardiography</i> , 2015, 32, 1441-1443.	0.9	5
42	Determinants and Regression Equations for the Calculation of $\frac{V_{LVT}}{V_{LVT+RVT}}$ Scores of Left Ventricular Tissue Doppler Longitudinal Indexes in a Healthy Italian Pediatric Population. <i>Cardiology Research and Practice</i> , 2015, 2015, 1-8.	1.1	2
43	Advanced therapies in patients with congenital heart disease-related pulmonary arterial hypertension: results from a long-term, single center, real-world follow-up. <i>Internal and Emergency Medicine</i> , 2015, 10, 445-450.	2.0	3
44	Right Aortic Arch Detected Prenatally: A Rare Case With Bilateral Arterial Duct and Nonconfluent Pulmonary Arteries. <i>Canadian Journal of Cardiology</i> , 2015, 31, 1205.e1-1205.e2.	1.7	2
45	Clinical Outcome, Valve Dysfunction, and Progressive Aortic Dilatation in a Pediatric Population With Isolated Bicuspid Aortic Valve. <i>Pediatric Cardiology</i> , 2014, 35, 803-809.	1.3	15
46	ALCAPA and massive pulmonary atelectasis: How a stent in the airway can be life-saving. <i>International Journal of Pediatric Otorhinolaryngology</i> , 2014, 78, 2015-2017.	1.0	4
47	Prevalence and Long-Term Predictors of Left Ventricular Hypertrophy, Late Hypertension, and Hypertensive Response to Exercise After Successful Aortic Coarctation Repair. <i>Pediatric Cardiology</i> , 2013, 34, 620-629.	1.3	43
48	Giant aorto-pulmonary collaterals in pulmonary atresia and ventricular septal defect. <i>Journal of Cardiovascular Medicine</i> , 2013, 14, 613-615.	1.5	4
49	Prevalence and clinical characteristics of adult patients with congenital heart disease in Tuscany. <i>Journal of Cardiovascular Medicine</i> , 2012, 13, 805-809.	1.5	10
50	Atrial standstill disease progression documented after 13 years follow-up. <i>Internal and Emergency Medicine</i> , 2012, 7, 7-8.	2.0	3
51	Controversies in the therapy of isolated congenital complete heart block. <i>Journal of Cardiovascular Medicine</i> , 2010, 11, 426-430.	1.5	6
52	Sildenafil as –first line therapy– in pulmonary persistent hypertension of the newborn?. <i>Journal of Maternal-Fetal and Neonatal Medicine</i> , 2010, 23, 104-105.	1.5	4
53	Tunneled left anterior descending artery in a child with hypertrophic cardiomyopathy. <i>Nature Clinical Practice Cardiovascular Medicine</i> , 2009, 6, 134-139.	3.3	10
54	The use of B-type natriuretic peptide in paediatric patients: a review of literature. <i>Journal of Cardiovascular Medicine</i> , 2009, 10, 298-302.	1.5	15

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55	Severe Hypoplasia of the Posterior Mitral Leaflet. <i>Annals of Thoracic Surgery</i> , 2008, 86, 1978-1979.	1.3	21
56	What Is the Effective Diagnostic Role of Pediatric Cardiac Assessment in the Offspring of Women With Congenital Heart Disease?. <i>Pediatrics</i> , 2008, 122, 472-472.	2.1	0
57	Acute cardiac failure following pacing in an adult patient with congenital complete heart block. <i>Journal of Cardiovascular Medicine</i> , 2008, 9, 301-303.	1.5	3
58	Prenatal diagnosis and postnatal outcome in patients with absent pulmonary valve syndrome not associated with tetralogy of Fallot: report of one case and review of the literature. <i>Journal of Cardiovascular Medicine</i> , 2008, 9, 1127-1129.	1.5	9
59	An unusual case of tricuspid lesion in congenital corrected transposition of the great arteries. <i>Journal of Cardiovascular Medicine</i> , 2007, 8, 281-283.	1.5	0
60	Usefulness of integrated imaging in the diagnosis of a rare coronary artery anomaly in a young athlete. <i>Journal of Cardiovascular Medicine</i> , 2007, 8, 527-530.	1.5	1
61	Hyponatraemic hypertensive syndrome in a 15-month-old child with renal artery stenosis. <i>Pediatric Nephrology</i> , 2006, 21, 1027-1030.	1.7	15
62	Pulmonary Hypertension of the Neonate Resistant to Inhaled Nitric Oxide. <i>Journal of Pediatrics</i> , 2005, 147, 867.	1.8	0
63	A Major Involvement of the Cardiovascular System in Patients Affected by Marfan Syndrome: Novel Mutations in Fibrillin 1 Gene. <i>Journal of Molecular and Cellular Cardiology</i> , 1997, 29, 1877-1884.	1.9	27
64	Long-term cardiac follow-up of children with perinatally acquired human immunodeficiency virus-type 1 infection. <i>Cardiology in the Young</i> , 1996, 6, 143-148.	0.8	0
65	Natural history of mitral valve prolapse. <i>American Journal of Cardiology</i> , 1995, 75, 1028-1032.	1.6	154
66	Arrhythmias in mitral valve prolapse: Relation to anterior mitral leaflet thickening, clinical variables, and color Doppler echocardiographic parameters. <i>American Heart Journal</i> , 1994, 128, 919-927.	2.7	65
67	Transient ventricular septal hypertrophy in the first year of life associated with neonatal brain injury. <i>Pediatric Cardiology</i> , 1992, 13, 63-64.	1.3	5
68	Diastolic time intervals before and after nadolol in patients with hypertrophic cardiomyopathy. <i>Clinical Cardiology</i> , 1986, 9, 573-574.	1.8	0
69	Echocardiographic features of right ventricular infarction. <i>Clinical Cardiology</i> , 1984, 7, 405-412.	1.8	10