

# Eloise Giabicani

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/172506/publications.pdf>

Version: 2024-02-01

18  
papers

673  
citations

840119

11  
h-index

839053

18  
g-index

18  
all docs

18  
docs citations

18  
times ranked

965  
citing authors

#	ARTICLE	IF	CITATIONS
1	Diagnosis and management of Silver-Russell syndrome: first international consensus statement. <i>Nature Reviews Endocrinology</i> , 2017, 13, 105-124.	4.3	336
2	Overgrowth syndromes – clinical and molecular aspects and tumour risk. <i>Nature Reviews Endocrinology</i> , 2019, 15, 299-311.	4.3	59
3	Chromosome 14q32.2 Imprinted Region Disruption as an Alternative Molecular Diagnosis of Silver-Russell Syndrome. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2018, 103, 2436-2446.	1.8	48
4	Chromosomal rearrangements in the 11p15 imprinted region: 17 new 11p15.5 duplications with associated phenotypes and putative functional consequences. <i>Journal of Medical Genetics</i> , 2018, 55, 205-213.	1.5	36
5	Presentation of 493 Consecutive Girls with Idiopathic Central Precocious Puberty: A Single-Center Study. <i>PLoS ONE</i> , 2013, 8, e70931.	1.1	34
6	Transcriptional profiling at the <i>DLK1/MEG3</i> domain explains clinical overlap between imprinting disorders. <i>Science Advances</i> , 2019, 5, eaau9425.	4.7	29
7	Diagnosis and management of postnatal fetal growth restriction. <i>Best Practice and Research in Clinical Endocrinology and Metabolism</i> , 2018, 32, 523-534.	2.2	23
8	Increasing knowledge in <i>IGF1R</i> defects: lessons from 35 new patients. <i>Journal of Medical Genetics</i> , 2020, 57, 160-168.	1.5	20
9	<i>IGF2</i> : Development, Genetic and Epigenetic Abnormalities. <i>Cells</i> , 2022, 11, 1886.	1.8	18
10	New clinical and molecular insights into Silver-Russell syndrome. <i>Current Opinion in Pediatrics</i> , 2016, 28, 529-535.	1.0	14
11	Models for Predicting the Adult Height and Age at First Menstruation of Girls with Idiopathic Central Precocious Puberty. <i>PLoS ONE</i> , 2015, 10, e0120588.	1.1	14
12	Roles of Type 1 Insulin-Like Growth Factor (IGF) Receptor and IGF-II in Growth Regulation: Evidence From a Patient Carrying Both an 11p Paternal Duplication and 15q Deletion. <i>Frontiers in Endocrinology</i> , 2019, 10, 263.	1.5	10
13	Imprinted disorders and growth. <i>Annales D'Endocrinologie</i> , 2017, 78, 112-113.	0.6	9
14	Premature Pubarche before One Year of Age: Distinguishing between Mini-Puberty Variants and Precocious Puberty. <i>Medical Science Monitor</i> , 2015, 21, 955-963.	0.5	8
15	Sleep disordered breathing in Silver-Russell syndrome patients: a new outcome. <i>Sleep Medicine</i> , 2019, 64, 23-29.	0.8	5
16	Dental pulp stem cells as a promising model to study imprinting diseases. <i>International Journal of Oral Science</i> , 2022, 14, 19.	3.6	5
17	Height and body mass index in molecularly confirmed Silver-Russell syndrome and the long-term effects of growth hormone treatment. <i>Clinical Endocrinology</i> , 2022, 97, 284-292.	1.2	3
18	Screening of patients born small for gestational age with the Silver-Russell syndrome phenotype for <i>DLK1</i> variants. <i>European Journal of Human Genetics</i> , 2021, 29, 1756-1761.	1.4	2