

# Louise Benarroch

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/8524560/publications.pdf>

Version: 2024-02-01

10  
papers

240  
citations

1478280

6  
h-index

1372474

10  
g-index

11  
all docs

11  
docs citations

11  
times ranked

533  
citing authors

#	ARTICLE	IF	CITATIONS
1	Association of modifiers and other genetic factors explain Marfan syndrome clinical variability. <i>European Journal of Human Genetics</i> , 2018, 26, 1759-1772.	1.4	73
2	Genetic diversity and pathogenic variants as possible predictors of severity in a French sample of nonsyndromic heritable thoracic aortic aneurysms and dissections (nshTAAD). <i>Genetics in Medicine</i> , 2019, 21, 2015-2024.	1.1	39
3	Atypical antipsychotics and effects on feeding: from mice to men. <i>Psychopharmacology</i> , 2016, 233, 2629-2653.	1.5	38
4	Lamin A/C Assembly Defects in LMNA-Congenital Muscular Dystrophy Is Responsible for the Increased Severity of the Disease Compared with Emery-Dreifuss Muscular Dystrophy. <i>Cells</i> , 2020, 9, 844.	1.8	29
5	Pathogenic variants in THSD4, encoding the ADAMTS-like 6 protein, predispose to inherited thoracic aortic aneurysm. <i>Genetics in Medicine</i> , 2021, 23, 111-122.	1.1	25
6	Quantifying the Genetic Basis of Marfan Syndrome Clinical Variability. <i>Genes</i> , 2020, 11, 574.	1.0	11
7	Effects of acute olanzapine exposure on central insulin-mediated regulation of whole body fuel selection and feeding. <i>Psychoneuroendocrinology</i> , 2018, 98, 127-130.	1.3	6
8	Reference Expression Profile of Three FBN1 Transcript Isoforms and Their Association with Clinical Variability in Marfan Syndrome. <i>Genes</i> , 2019, 10, 128.	1.0	6
9	Marfan Syndrome Variability: Investigation of the Roles of Sarcolipin and Calcium as Potential Transregulator of FBN1 Expression. <i>Genes</i> , 2018, 9, 421.	1.0	4
10	Preclinical Advances of Therapies for Laminopathies. <i>Journal of Clinical Medicine</i> , 2021, 10, 4834.	1.0	4