

# Silvia Tognazzo

## List of Publications by Year in descending order

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33  
papers

2,636  
citations

218677

26  
h-index

395702

33  
g-index

33  
all docs

33  
docs citations

33  
times ranked

5494  
citing authors

#	ARTICLE	IF	CITATIONS
1	Identification of 12 new susceptibility loci for different histotypes of epithelial ovarian cancer. <i>Nature Genetics</i> , 2017, 49, 680-691.	21.4	356
2	Genome-Wide Association Study in BRCA1 Mutation Carriers Identifies Novel Loci Associated with Breast and Ovarian Cancer Risk. <i>PLoS Genetics</i> , 2013, 9, e1003212.	3.5	244
3	Mutational spectrum in a worldwide study of 29,700 families with <i>BRCA1</i> or <i>BRCA2</i> mutations. <i>Human Mutation</i> , 2018, 39, 593-620.	2.5	224
4	Identification of six new susceptibility loci for invasive epithelial ovarian cancer. <i>Nature Genetics</i> , 2015, 47, 164-171.	21.4	221
5	Prediction of Breast and Prostate Cancer Risks in Male <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers Using Polygenic Risk Scores. <i>Journal of Clinical Oncology</i> , 2017, 35, 2240-2250.	1.6	152
6	Value of Platelet Reactivity in Predicting Response to Treatment and Clinical Outcome in Patients Undergoing Primary Coronary Intervention. <i>Journal of the American College of Cardiology</i> , 2006, 48, 2178-2185.	2.8	140
7	Large scale multifactorial likelihood quantitative analysis of <i>BRCA1</i> and <i>BRCA2</i> variants: An ENIGMA resource to support clinical variant classification. <i>Human Mutation</i> , 2019, 40, 1557-1578.	2.5	102
8	A common mutation in the gene for coagulation factor XIII $\alpha$ (VAL34Leu): A risk factor for primary intracerebral hemorrhage is protective against atherothrombotic diseases. <i>American Journal of Hematology</i> , 2001, 67, 183-188.	4.1	97
9	<i>FANCM</i> c.5791C>T nonsense mutation (rs144567652) induces exon skipping, affects DNA repair activity and is a familial breast cancer risk factor. <i>Human Molecular Genetics</i> , 2015, 24, 5345-5355.	2.9	91
10	Hemochromatosis C282Y gene mutation increases the risk of venous leg ulceration. <i>Journal of Vascular Surgery</i> , 2005, 42, 309-314.	1.1	89
11	Polygenic risk scores and breast and epithelial ovarian cancer risks for carriers of <i>BRCA1</i> and <i>BRCA2</i> pathogenic variants. <i>Genetics in Medicine</i> , 2020, 22, 1653-1666.	2.4	82
12	Functional mechanisms underlying pleiotropic risk alleles at the 19p13.1 breast-ovarian cancer susceptibility locus. <i>Nature Communications</i> , 2016, 7, 12675.	12.8	78
13	Common breast cancer susceptibility alleles are associated with tumour subtypes in <i>BRCA1</i> and <i>BRCA2</i> mutation carriers: results from the Consortium of Investigators of Modifiers of <i>BRCA1/2</i> . <i>Breast Cancer Research</i> , 2011, 13, R110.	5.0	71
14	Common alleles at 6q25.1 and 1p11.2 are associated with breast cancer risk for <i>BRCA1</i> and <i>BRCA2</i> mutation carriers. <i>Human Molecular Genetics</i> , 2011, 20, 3304-3321.	2.9	68
15	The overlapping of local iron overload and HFE mutation in venous leg ulcer pathogenesis. <i>Free Radical Biology and Medicine</i> , 2006, 40, 1869-1873.	2.9	61
16	Characterization of <i>BRCA1</i> and <i>BRCA2</i> splicing variants: a collaborative report by ENIGMA consortium members. <i>Breast Cancer Research and Treatment</i> , 2012, 132, 1009-1023.	2.5	56
17	Methylenetetrahydrofolate reductase C677T and A1298C gene variants in adult non-Hodgkin's lymphoma patients: association with toxicity and survival. <i>Haematologica</i> , 2007, 92, 478-485.	3.5	53
18	DNA Glycosylases Involved in Base Excision Repair May Be Associated with Cancer Risk in <i>BRCA1</i> and <i>BRCA2</i> Mutation Carriers. <i>PLoS Genetics</i> , 2014, 10, e1004256.	3.5	47

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19	Prognostic role of factor XIII gene variants in nonhealing venous leg ulcers. <i>Journal of Vascular Surgery</i> , 2006, 44, 815-819.	1.1	45
20	Factor XIII V34L polymorphism modulates the risk of chronic venous leg ulcer progression and extension. <i>Wound Repair and Regeneration</i> , 2004, 12, 512-517.	3.0	43
21	Influence of gene polymorphisms in ulcer healing process after superficial venous surgery. <i>Journal of Vascular Surgery</i> , 2006, 44, 554-562.	1.1	43
22	Serum Iron and Matrix Metalloproteinase-9 Variations in Limbs Affected by Chronic Venous Disease and Venous Leg Ulcers. <i>Dermatologic Surgery</i> , 2005, 31, 644-649.	0.8	42
23	Factor XIII Contrasts the Effects of Metalloproteinases in Human Dermal Fibroblast Cultured Cells. <i>Vascular and Endovascular Surgery</i> , 2004, 38, 431-438.	0.7	37
24	Ovarian cancer susceptibility alleles and risk of ovarian cancer in BRCA1 and BRCA2 mutation carriers. <i>Human Mutation</i> , 2012, 33, 690-702.	2.5	34
25	Factor XIII A-V34L and Factor XIII B-H95R Gene Variants: Effects on Survival in Myocardial Infarction Patients. <i>Molecular Medicine</i> , 2007, 13, 112-120.	4.4	32
26	Serum Iron and Matrix Metalloproteinase-9 Variations in Limbs Affected by Chronic Venous Disease and Venous Leg Ulcers. <i>Dermatologic Surgery</i> , 2006, 31, 644-649.	0.8	31
27	Mendelian randomisation study of height and body mass index as modifiers of ovarian cancer risk in 22,588 BRCA1 and BRCA2 mutation carriers. <i>British Journal of Cancer</i> , 2019, 121, 180-192.	6.4	19
28	Association of breast cancer risk in BRCA1 and BRCA2 mutation carriers with genetic variants showing differential allelic expression: identification of a modifier of breast cancer risk at locus 11q22.3. <i>Breast Cancer Research and Treatment</i> , 2017, 161, 117-134.	2.5	18
29	CDKN2A Unclassified Variants in Familial Malignant Melanoma: Combining Functional and Computational Approaches for Their Assessment. <i>Human Mutation</i> , 2014, 35, 828-840.	2.5	17
30	Prevalence of factor XIII Val34Leu polymorphism in patients affected by spontaneous subconjunctival hemorrhage. <i>American Journal of Ophthalmology</i> , 2004, 138, 481-484.	3.3	14
31	Contribution of susceptibility gene variants to melanoma risk in families from the Veneto region of Italy. <i>Pigment Cell and Melanoma Research</i> , 2011, 24, 728-730.	3.3	12
32	Fine-Scale Mapping at 9p22.2 Identifies Candidate Causal Variants That Modify Ovarian Cancer Risk in BRCA1 and BRCA2 Mutation Carriers. <i>PLoS ONE</i> , 2016, 11, e0158801.	2.5	10
33	Vision Loss after PDT. <i>Ophthalmology</i> , 2006, 113, 157.e1-157.e4.	5.2	7