

# Kati M Donner

## List of Publications by Year in descending order

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34  
papers

1,833  
citations

377584

21  
h-index

425179

34  
g-index

34  
all docs

34  
docs citations

34  
times ranked

1958  
citing authors

#	ARTICLE	IF	CITATIONS
1	Pharmacoeigenetics of hypertension: genome-wide methylation analysis of responsiveness to four classes of antihypertensive drugs using a double-blind crossover study design. <i>Epigenetics</i> , 2022, , 1-14.	1.3	7
2	Missense mutations in small muscle protein X-linked (SMPX) cause distal myopathy with protein inclusions. <i>Acta Neuropathologica</i> , 2021, 142, 375-393.	3.9	6
3	Dominant Distal Myopathy 3 (MPD3) Caused by a Deletion in the <i>HNRNPA1</i> Gene. <i>Neurology: Genetics</i> , 2021, 7, e632.	0.9	7
4	Genomics of asthma, allergy and chronic rhinosinusitis: novel concepts and relevance in airway mucosa. <i>Clinical and Translational Allergy</i> , 2020, 10, 45.	1.4	26
5	Heme oxygenase-1 repeat polymorphism in septic acute kidney injury. <i>PLoS ONE</i> , 2019, 14, e0217291.	1.1	16
6	Replicated evidence for aminoacylase 3 and nephrin gene variations to predict antihypertensive drug responses. <i>Pharmacogenomics</i> , 2017, 18, 445-458.	0.6	18
7	PRKDCBP (CAVIN3) and CRY2 associate with major depressive disorder. <i>Journal of Affective Disorders</i> , 2017, 207, 136-140.	2.0	20
8	CRY1 and CRY2 genetic variants in seasonality: A longitudinal and cross-sectional study. <i>Psychiatry Research</i> , 2016, 242, 101-110.	1.7	10
9	PTPRD gene associated with blood pressure response to atenolol and resistant hypertension. <i>Journal of Hypertension</i> , 2015, 33, 2278-2285.	0.3	38
10	TET2 and CSMD1 genes affect SBP response to hydrochlorothiazide in never-treated essential hypertensives. <i>Journal of Hypertension</i> , 2015, 33, 1301-1309.	0.3	29
11	Genetic Loci Associated with Allergic Sensitization in Lithuanians. <i>PLoS ONE</i> , 2015, 10, e0134188.	1.1	4
12	SIRT1 Polymorphisms Associate with Seasonal Weight Variation, Depressive Disorders, and Diastolic Blood Pressure in the General Population. <i>PLoS ONE</i> , 2015, 10, e0141001.	1.1	23
13	Pharmacogenomics of Hypertension: A Genome-Wide, Placebo-Controlled Cross-Over Study, Using Four Classes of Antihypertensive Drugs. <i>Journal of the American Heart Association</i> , 2015, 4, e001521.	1.6	74
14	CRY1, CRY2 and PRKDCBP genetic variants in metabolic syndrome. <i>Hypertension Research</i> , 2015, 38, 186-192.	1.5	35
15	Mutation Update: The Spectra of Nebulin Variants and Associated Myopathies. <i>Human Mutation</i> , 2014, 35, 1418-1426.	1.1	107
16	Generalized glucocorticoid resistance caused by a novel two-nucleotide deletion in the hormone-binding domain of the glucocorticoid receptor gene NR3C1. <i>European Journal of Endocrinology</i> , 2013, 168, K9-K18.	1.9	18
17	Genomic Association Analysis of Common Variants Influencing Antihypertensive Response to Hydrochlorothiazide. <i>Hypertension</i> , 2013, 62, 391-397.	1.3	96
18	CRY2 Genetic Variants Associate with Dysthymia. <i>PLoS ONE</i> , 2013, 8, e71450.	1.1	42

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19	STK39 variation predicts the ambulatory blood pressure response to losartan in hypertensive men. <i>Hypertension Research</i> , 2012, 35, 107-114.	1.5	21
20	Abnormal actin binding of aberrant $\beta$ -tropomyosins is a molecular cause of muscle weakness in <i>TPM2</i> -related nemaline and cap myopathy. <i>Biochemical Journal</i> , 2012, 442, 231-239.	1.7	48
21	Common genetic variation of $\beta$ 1- and $\beta$ 2-adrenergic receptor and response to four classes of antihypertensive treatment. <i>Pharmacogenetics and Genomics</i> , 2010, 20, 342-345.	0.7	33
22	Renin-Angiotensin System and $\beta$ -Adducin Gene Polymorphisms and Their Relation to Responses to Antihypertensive Drugs: Results From the GENRES Study. <i>American Journal of Hypertension</i> , 2009, 22, 169-175.	1.0	37
23	CYP2C9 genotype modifies activity of the renin-angiotensin-aldosterone system in hypertensive men. <i>Journal of Hypertension</i> , 2009, 27, 2001-2009.	0.3	16
24	Identification of a founder mutation in TPM3 in nemaline myopathy patients of Turkish origin. <i>European Journal of Human Genetics</i> , 2008, 16, 1055-1061.	1.4	36
25	Laboratory tests as predictors of the antihypertensive effects of amlodipine, bisoprolol, hydrochlorothiazide and losartan in men: results from the randomized, double-blind, crossover GENRES Study. <i>Journal of Hypertension</i> , 2008, 26, 1250-1256.	0.3	29
26	Developmental and muscle-type-specific expression of mouse nebulin exons 127 and 128. <i>Genomics</i> , 2006, 88, 489-495.	1.3	20
27	Identification of 45 novel mutations in the nebulin gene associated with autosomal recessive nemaline myopathy. <i>Human Mutation</i> , 2006, 27, 946-956.	1.1	112
28	Complete genomic structure of the human nebulin gene and identification of alternatively spliced transcripts. <i>European Journal of Human Genetics</i> , 2004, 12, 744-751.	1.4	101
29	Mutations in the $\beta$ -tropomyosin (TPM2) gene – a rare cause of nemaline myopathy. <i>Neuromuscular Disorders</i> , 2002, 12, 151-158.	0.3	210
30	Mutations in the nebulin gene can cause severe congenital nemaline myopathy. <i>Neuromuscular Disorders</i> , 2002, 12, 674-679.	0.3	85
31	Nebulin mutations in autosomal recessive nemaline myopathy: an update. <i>Neuromuscular Disorders</i> , 2002, 12, 680-686.	0.3	55
32	Nebulin expression in patients with nemaline myopathy. <i>Neuromuscular Disorders</i> , 2001, 11, 154-162.	0.3	39
33	Mutations in the skeletal muscle $\beta$ -actin gene in patients with actin myopathy and nemaline myopathy. <i>Nature Genetics</i> , 1999, 23, 208-212.	9.4	389
34	Refined Localisation of the Genes for Nebulin and Titin on Chromosome 2q Allows the Assignment of Nebulin as a Candidate Gene for Autosomal Recessive Nemaline Myopathy. <i>European Journal of Human Genetics</i> , 1997, 5, 229-234.	1.4	26