Francois H Van Der Westhuizen

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

87
papers

2,181
25
h-index

92
ext. papers

2,490
ext. citations

4.7
avg, IF

4.76
L-index

#	Paper	IF	Citations
87	Inhibition of complex I of the electron transport chain causes O2mediated mitochondrial outgrowth. <i>American Journal of Physiology - Cell Physiology</i> , 2005 , 288, C1440-50	5.4	227
86	OXPHOS gene expression and control in mitochondrial disorders. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2009 , 1792, 1113-21	6.9	101
85	Chemoprotective properties of rooibos (Aspalathus linearis), honeybush (Cyclopia intermedia) herbal and green and black (Camellia sinensis) teas against cancer promotion induced by fumonisin B1 in rat liver. <i>Food and Chemical Toxicology</i> , 2009 , 47, 220-9	4.7	91
84	Aloe ferox leaf gel phytochemical content, antioxidant capacity, and possible health benefits. <i>Journal of Agricultural and Food Chemistry</i> , 2007 , 55, 6891-6	5.7	89
83	Recessive germline SDHA and SDHB mutations causing leukodystrophy and isolated mitochondrial complex II deficiency. <i>Journal of Medical Genetics</i> , 2012 , 49, 569-77	5.8	87
82	The effects of high walnut and cashew nut diets on the antioxidant status of subjects with metabolic syndrome. <i>European Journal of Nutrition</i> , 2007 , 46, 155-64	5.2	87
81	Metallothionein isoform 2A expression is inducible and protects against ROS-mediated cell death in rotenone-treated HeLa cells. <i>Biochemical Journal</i> , 2006 , 395, 405-15	3.8	81
80	Modulation of hepatic drug metabolizing enzymes and oxidative status by rooibos (Aspalathus linearis) and Honeybush (Cyclopia intermedia), green and black (Camellia sinensis) teas in rats. Journal of Agricultural and Food Chemistry, 2003 , 51, 8113-9	5.7	80
79	3-Methylglutaconic acidurialessons from 50 genes and 977 patients. <i>Journal of Inherited Metabolic Disease</i> , 2013 , 36, 913-21	5.4	66
78	Curcumin Rescues a PINK1 Knock Down SH-SY5Y Cellular Model of Parkinson's Disease from Mitochondrial Dysfunction and Cell Death. <i>Molecular Neurobiology</i> , 2017 , 54, 2752-2762	6.2	59
77	Metabolomics of mitochondrial disease. <i>Mitochondrion</i> , 2017 , 35, 97-110	4.9	52
76	Glycaemic control improves fibrin network characteristics in type 2 diabetes - a purified fibrinogen model. <i>Thrombosis and Haemostasis</i> , 2008 , 99, 691-700	7	52
75	Metabolomics of urinary organic acids in respiratory chain deficiencies in children. <i>Metabolomics</i> , 2012 , 8, 264-283	4.7	50
74	Glycation of fibrinogen in uncontrolled diabetic patients and the effects of glycaemic control on fibrinogen glycation. <i>Thrombosis Research</i> , 2007 , 120, 439-46	8.2	43
73	Polyphenol composition and antioxidant activity of Kei-apple (Dovyalis caffra) juice. <i>Journal of Agricultural and Food Chemistry</i> , 2006 , 54, 1271-6	5.7	41
72	A kinetic study into the hydrolysis of the ochratoxins and analogues by carboxypeptidase A. <i>Chemical Research in Toxicology</i> , 2001 , 14, 302-4	4	41
71	Human mitochondrial complex I deficiency: investigating transcriptional responses by microarray. <i>Neuropediatrics</i> , 2003 , 34, 14-22	1.6	40

(2015-2008)

70	Phytochemical contents and antioxidant capacities of two Aloe greatheadii var. davyana extracts. <i>Molecules</i> , 2008 , 13, 2169-80	4.8	38	
69	The involvement of metallothioneins in mitochondrial function and disease. <i>Current Protein and Peptide Science</i> , 2010 , 11, 292-309	2.8	37	
68	Blood glutathione and subclinical atherosclerosis in African men: the SABPA Study. <i>American Journal of Hypertension</i> , 2009 , 22, 1154-9	2.3	35	
67	Associations between reactive oxygen species, blood pressure and arterial stiffness in black South Africans: the SABPA study. <i>Journal of Human Hypertension</i> , 2012 , 26, 91-7	2.6	32	
66	Inhibition of N-acetylglutamate synthase by various monocarboxylic and dicarboxylic short-chain coenzyme A esters and the production of alternative glutamate esters. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2014 , 1842, 2510-6	6.9	29	
65	Altered Mitochondrial Respiration and Other Features of Mitochondrial Function in Parkin-Mutant Fibroblasts from Parkinson's Disease Patients. <i>Parkinson's Disease</i> , 2016 , 2016, 1819209	2.6	27	
64	Kinetic analysis, size profiling, and bioenergetic association of DNA released by selected cell lines in vitro. <i>Cellular and Molecular Life Sciences</i> , 2017 , 74, 2689-2707	10.3	26	
63	Obesity and metabolomics: metallothioneins protect against high-fat diet-induced consequences in metallothionein knockout mice. <i>OMICS A Journal of Integrative Biology</i> , 2015 , 19, 92-103	3.8	25	
62	The effect of glycaemic control on fibrin network structure of type 2 diabetic subjects. <i>Thrombosis and Haemostasis</i> , 2006 , 96, 623-629	7	25	
61	Use of metabolomics to elucidate the metabolic perturbation associated with hypertension in a black South African male cohort: the SABPA study. <i>Journal of the American Society of Hypertension</i> , 2015 , 9, 104-14		23	
60	DNA damage and repair detected by the comet assay in lymphocytes of african petrol attendants: a pilot study. <i>Annals of Occupational Hygiene</i> , 2008 , 52, 653-62		23	
59	In vitro antioxidant, antimutagenic and genoprotective activity of Rosa roxburghii fruit extract. <i>Phytotherapy Research</i> , 2008 , 22, 376-83	6.7	23	
58	The utilization of alanine, glutamic acid, and serine as amino acid substrates for glycine N-acyltransferase. <i>Journal of Biochemical and Molecular Toxicology</i> , 2000 , 14, 102-9	3.4	23	
57	Combined tarsal and carpal tunnel syndrome in mucolipidosis type III. A case study and review. <i>Annals of the New York Academy of Sciences</i> , 2009 , 1151, 77-84	6.5	22	
56	Characterization of mtDNA variation in a cohort of South African paediatric patients with mitochondrial disease. <i>European Journal of Human Genetics</i> , 2012 , 20, 650-6	5.3	21	
55	Disclosure of a putative biosignature for respiratory chain disorders through a metabolomics approach. <i>Metabolomics</i> , 2013 , 9, 379-391	4.7	20	
54	A significant decline in IGF-I may predispose young Africans to subsequent cardiometabolic vulnerability. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2010 , 95, 2503-7	5.6	20	
53	Untargeted urine metabolomics reveals a biosignature for muscle respiratory chain deficiencies. <i>Metabolomics</i> , 2015 , 11, 111-121	4.7	19	

52	Rosa roxburghii supplementation in a controlled feeding study increases plasma antioxidant capacity and glutathione redox state. <i>European Journal of Nutrition</i> , 2005 , 44, 452-7	5.2	19
51	The effect of glycaemic control on fibrin network structure of type 2 diabetic subjects. <i>Thrombosis and Haemostasis</i> , 2006 , 96, 623-9	7	18
50	Conservation of the coding regions of the glycine N-acyltransferase gene further suggests that glycine conjugation is an essential detoxification pathway. <i>Gene</i> , 2015 , 571, 126-34	3.8	16
49	Systemic and organ specific metabolic variation in metallothionein knockout mice challenged with swimming exercise. <i>Metabolomics</i> , 2013 , 9, 418-432	4.7	15
48	Characterisation of the influence of genetic variations on the enzyme activity of a recombinant human glycine N-acyltransferase. <i>Gene</i> , 2013 , 515, 447-53	3.8	15
47	Clinically proven mtDNA mutations are not common in those with chronic fatigue syndrome. <i>BMC Medical Genetics</i> , 2017 , 18, 29	2.1	14
46	DNA Methylation Associated with Mitochondrial Dysfunction in a South African Autism Spectrum Disorder Cohort. <i>Autism Research</i> , 2020 , 13, 1079-1093	5.1	14
45	Development and validation of LC-ESI-MS/MS methods for quantification of 27 free and conjugated estrogen-related metabolites. <i>Analytical Biochemistry</i> , 2020 , 590, 113531	3.1	14
44	MtDNA population variation in Myalgic encephalomyelitis/Chronic fatigue syndrome in two populations: a study of mildly deleterious variants. <i>Scientific Reports</i> , 2019 , 9, 2914	4.9	14
43	Using MutPred derived mtDNA load scores to evaluate mtDNA variation in hypertension and diabetes in a two-population cohort: The SABPA study. <i>Journal of Genetics and Genomics</i> , 2017 , 44, 139-	149	13
42	The First Case of Riboflavin Transporter Deficiency in sub-Saharan Africa. <i>Seminars in Pediatric Neurology</i> , 2018 , 26, 10-14	2.9	13
41	Metabolic and glutathione redox markers associated with brain-derived neurotrophic factor in depressed african men and women: evidence for counterregulation?. <i>Neuropsychobiology</i> , 2013 , 67, 33-	4 6	13
40	An overview of a cohort of South African patients with mitochondrial disorders. <i>Journal of Inherited Metabolic Disease</i> , 2010 , 33 Suppl 3, S95-104	5.4	13
39	The unresolved role of mitochondrial DNA in Parkinson's disease: An overview of published studies, their limitations, and future prospects. <i>Neurochemistry International</i> , 2019 , 129, 104495	4.4	12
38	8-Oxo-7,8-dihydro-2Sdeoxyguanosine, reactive oxygen species and ambulatory blood pressure in African and Caucasian men: the SABPA study. <i>Free Radical Research</i> , 2014 , 48, 1291-9	4	12
37	A call for global action for rare diseases in Africa. <i>Nature Genetics</i> , 2020 , 52, 21-26	36.3	11
36	A novel mutation in ETFDH manifesting as severe neonatal-onset multiple acyl-CoA dehydrogenase deficiency. <i>Journal of the Neurological Sciences</i> , 2018 , 384, 121-125	3.2	11
35	Attenuation of Endoplasmic Reticulum Stress, Impaired Calcium Homeostasis, and Altered Bioenergetic Functions in MPP-Exposed SH-SY5Y Cells Pretreated with Rutin. <i>Neurotoxicity</i>	4.3	10

(2014-2016)

underlying mitochondrial dysfunction and oxidative stress. <i>International Journal of Biochemistry and Cell Biology</i> , 2016 , 78, 116-129	5.6	10
A urinary biosignature for mitochondrial myopathy, encephalopathy, lactic acidosis and stroke like episodes (MELAS). <i>Mitochondrion</i> , 2019 , 45, 38-45	4.9	10
A molecular analysis of the gene in Caucasian South Africans with Parkinson's disease. <i>Molecular Genetics</i> & Amp; Genomic Medicine, 2017, 5, 147-156	2.3	9
Panel-Based Nuclear and Mitochondrial Next-Generation Sequencing Outcomes of an Ethnically Diverse Pediatric Patient Cohort with Mitochondrial Disease. <i>Journal of Molecular Diagnostics</i> , 2019 , 21, 503-513	5.1	9
The dilemma of diagnosing coenzyme Q deficiency in muscle. <i>Molecular Genetics and Metabolism</i> , 2018 , 125, 38-43	3.7	9
Aberrant synthesis of ATP synthase resulting from a novel deletion in mitochondrial DNA in an African patient with progressive external ophthalmoplegia. <i>Journal of Inherited Metabolic Disease</i> , 2010 , 33 Suppl 3, S55-62	5.4	9
Increased excretion of c4-carnitine species after a therapeutic acetylsalicylic Acid dose: evidence for an inhibitory effect on short-chain Fatty Acid metabolism. <i>ISRN Pharmacology</i> , 2011 , 2011, 851870		9
The aetiology of cardiovascular disease: a role for mitochondrial DNA?. <i>Cardiovascular Journal of Africa</i> , 2018 , 29, 122-132	0.7	9
Metabolomics of Ndufs4 skeletal muscle: Adaptive mechanisms converge at the ubiquinone-cycle. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2019 , 1865, 98-106	6.9	9
Toward a mtDNA locus-specific mutation database using the LOVD platform. <i>Human Mutation</i> , 2012 , 33, 1352-8	4.7	7
Disorders of flavin adenine dinucleotide metabolism: MADD and related deficiencies. <i>International Journal of Biochemistry and Cell Biology</i> , 2021 , 132, 105899	5.6	7
Barriers and Considerations for Diagnosing Rare Diseases in Indigenous Populations. <i>Frontiers in Pediatrics</i> , 2020 , 8, 579924	3.4	6
Draft De Novo Genome Sequence of Agapornis roseicollis for Application in Avian Breeding. <i>Animal Biotechnology</i> , 2018 , 29, 241-246	1.4	5
The contribution of the C-824T tyrosine hydroxylase polymorphism to the prevalence of hypertension in a South African cohort: the SABPA study. <i>Clinical and Experimental Hypertension</i> , 2013 , 35, 614-9	2.2	5
Metallothionein 1 Overexpression Does Not Protect Against Mitochondrial Disease Pathology in Ndufs4 Knockout Mice. <i>Molecular Neurobiology</i> , 2021 , 58, 243-262	6.2	5
Curcumin pre-treatment may protect against mitochondrial damage in -mutant Parkinson's disease and healthy control fibroblasts. <i>Biochemistry and Biophysics Reports</i> , 2021 , 27, 101035	2.2	5
Exploring the link between serum peroxides and angiogenesis in a bi-ethnic population from South Africa: the SAfrEIC study. <i>Journal of the American Society of Hypertension</i> , 2013 , 7, 267-75		4
Leukocyte telomere length and hemostatic factors in a South African cohort: the SABPA Study. Journal of Thrombosis and Haemostasis, 2014 , 12, 1975-85	15.4	4
	Aurinary biosignature for mitochondrial myopathy, encephalopathy, lactic acidosis and stroke like episodes (MELAS). <i>Mitochondrion</i> , 2019, 45, 38-45 A molecular analysis of the gene in Caucasian South Africans with Parkinson's disease. <i>Molecular Genetics & Amp; Genomic Medicine</i> , 2017, 5, 147-156 Panel-Based Nuclear and Mitochondrial Next-Generation Sequencing Outcomes of an Ethnically Diverse Pediatric Patient Cohort with Mitochondrial Disease. <i>Journal of Molecular Diagnostics</i> , 2019, 21, 503-513 The dilemma of diagnosing coenzyme Q deficiency in muscle. <i>Molecular Genetics and Metabolism</i> , 2018, 125, 38-43 Aberrant synthesis of ATP synthase resulting from a novel deletion in mitochondrial DNA in an African patient with progressive external ophthalmoplegia. <i>Journal of Inherited Metabolic Disease</i> , 2010, 33 Suppl 3, 555-62 Increased excretion of c4-carnitine species after a therapeutic acetylsalicylic Acid dose: evidence for an inhibitory effect on short-chain Fatty Acid metabolism. <i>SRN Pharmacology</i> , 2011, 2011, 851870 The aetiology of cardiovascular disease: a role for mitochondrial DNA?. <i>Cardiovascular Journal of Africa</i> , 2018, 29, 122-132 Metabolomics of Ndufs4 skeletal muscle: Adaptive mechanisms converge at the ubiquinone-cycle. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2019, 1865, 98-106 Toward a mtDNA locus-specific mutation database using the LOVD platform. <i>Human Mutation</i> , 2012, 33, 1352-8 Disorders of flavin adenine dinucleotide metabolism: MADD and related deficiencies. <i>International Journal of Biochemistry and Cell Biology</i> , 2021, 132, 105899 Barriers and Considerations for Diagnosing Rare Diseases in Indigenous Populations. <i>Frantiers in Pediatrics</i> , 2020, 8, 579924 Draft De Novo Genome Sequence of Agapornis roseicollis for Application in Avian Breeding. <i>Animal Biotechnology</i> , 2018, 29, 241-246 The contribution of the C-824T tyrosine hydroxylase polymorphism to the prevalence of hypertension in a South African cohort: the SABPA Study. <i>Clinical and Experimen</i>	A urinary biosignature for mitochondrial myopathy, encephalopathy, lactic acidosis and stroke like episodes (MELAS). Mitochondrian, 2019, 45, 38-45 A molecular analysis of the gene in Caucasian South Africans with Parkinson's disease. Molecular Genetics & mp; Genomic Medicine, 2017, 5, 147-156 Panel-Based Nuclear and Mitochondrial Next-Generation Sequencing Outcomes of an Ethnically Diverse Pediatric Patient Cohort with Mitochondrial Disease. Journal of Molecular Diagnostics, 2019, 21, 503-513 The dilemma of diagnosing coenzyme Q deficiency in muscle. Molecular Genetics and Metabolism, 2018, 125, 38-43 Aberrant synthesis of ATP synthase resulting from a novel deletion in mitochondrial DNA in an African patient with progressive external ophthalmoplegia. Journal of Inherited Metabolic Disease, 2010, 33 Suppl 3, 555-62 Increased excretion of c4-carnitine species after a therapeutic acetylsalicylic Acid dose: evidence for an inhibitory effect on short-chain Fatty Acid metabolism. ISRN Pharmacology, 2011, 2011, 851870 The aetiology of cardiovascular disease: a role for mitochondrial DNA?. Cardiovascular Journal of Africa, 2018, 29, 122-132 Metabolomics of Ndufs4 skeletal muscle: Adaptive mechanisms converge at the ubiquinone-cycle. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2019, 1865, 98-106 Toward a mtDNA locus-specific mutation database using the LOVD platform. Human Mutation, 2012, 33, 1352-8 Disorders of flavin adenine dinucleotide metabolism: MADD and related deficiencies. International Journal of Biochemistry and Cell Biology, 2021, 132, 105899 Barriers and Considerations for Diagnosing Rare Diseases in Indigenous Populations. Frontiers in Pediatrics, 2020, 8, 579924 Draft De Novo Genome Sequence of Agapornis roseicollis for Application in Avian Breeding. Animal Biotechinology, 2014, 747-748. Metallothionein 1 Overexpression Does Not Protect Against Mitochondrial Disease Pathology in Nodufs4 Knockout Mice. Molecular Neurobiology, 2021, 58, 243-262 Exploring the link between serum

16	One mutation, three phenotypes: novel metabolic insights on MELAS, MIDD and myopathy caused by the m.3243A > G mutation. <i>Metabolomics</i> , 2021 , 17, 10	4.7	4
15	Investigating the effects of the presence of foreign DNA on DNA methylation and DNA repair events in cultured eukaryotic cells. <i>Gene</i> , 2013 , 512, 117-22	3.8	3
14	Mitochondrial DNA replication and OXPHOS gene transcription show varied responsiveness to Rieske protein knockdown in 143B cells. <i>Biochimie</i> , 2011 , 93, 758-65	4.6	3
13	Carnitine palmitoyltransferase I activity monitoring in fibroblasts and leukocytes using electrospray ionization mass spectrometry. <i>Analytical Biochemistry</i> , 1998 , 256, 178-84	3.1	3
12	Contribution of regions 3Sand 5Sto the hIL-5 gene on the expression of rhIL-5 in CHO-cells. <i>Biochemical and Biophysical Research Communications</i> , 1996 , 227, 576-80	3.4	3
11	Aberrant BCAA and glutamate metabolism linked to regional neurodegeneration in a mouse model of Leigh syndrome. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2021 , 1867, 166082	6.9	3
10	Nuclear Genes Associated with Mitochondrial DNA Processes as Contributors to Parkinson Disease Risk. <i>Movement Disorders</i> , 2021 , 36, 815-831	7	3
9	Implementing a new variant load model to investigate the role of mtDNA in oxidative stress and inflammation in a bi-ethnic cohort: the SABPA study. <i>Mitochondrial DNA Part A: DNA Mapping, Sequencing, and Analysis</i> , 2019 , 30, 440-447	1.3	2
8	A bioinformatics pipeline for rare genetic diseases in South African patients. <i>South African Journal of Science</i> , 2019 , 115,	1.3	2
7	Proteomics and metabolomics of HIV-associated neurocognitive disorders: A systematic review. Journal of Neurochemistry, 2021 , 157, 429-449	6	2
6	S-Nitrosylation and Attenuation of Excessive Calcium Flux by Pentacycloundecane Derivatives. <i>Medicinal Chemistry</i> , 2012 , 8, 367-371	1.8	1
5	Simple, rapid technique for cloning eukaryotic colonies. <i>Technical Tips Online</i> , 2002 , 7, 1-2		1
4	Health Status Is Affected, and Phase I/II Biotransformation Activity Altered in Young Women Using Oral Contraceptives Containing Drospirenone/Ethinyl Estradiol. <i>International Journal of Environmental Research and Public Health</i> , 2021 , 18,	4.6	1
3	Data on the optimisation of a solid phase extraction method for fractionating estrogen metabolites from small urine volumes. <i>Data in Brief</i> , 2020 , 29, 105222	1.2	O
2	Cross-comparison of systemic and tissue-specific metabolomes in a mouse model of Leigh syndrome. <i>Metabolomics</i> , 2021 , 17, 101	4.7	О
1	NON-PROTEIN AMINO ACIDS AND ORGANIC ACIDS INHIBIT SERINE HYDROXYMETHYLTRANSFERASE ACTIVITY IN THE CHICKEN EMBRYO NEURAL TUBE DEFECT MODEL. <i>Biochemical Society Transactions</i> , 2000 , 28, A438-A438	5.1	