

Melanie B Gillingham

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.

47
papers

976
citations

18
h-index

31
g-index

49
ext. papers

1,142
ext. citations

3.2
avg, IF

3.75
L-index

#	Paper	IF	Citations
47	Current issues regarding treatment of mitochondrial fatty acid oxidation disorders. <i>Journal of Inherited Metabolic Disease</i> , 2010 , 33, 555-61	5.4	85
46	Beneficial and cautionary outcomes of resveratrol supplementation in pregnant nonhuman primates. <i>FASEB Journal</i> , 2014 , 28, 2466-77	0.9	84
45	Maternal high fat diet is associated with decreased plasma n-3 fatty acids and fetal hepatic apoptosis in nonhuman primates. <i>PLoS ONE</i> , 2011 , 6, e17261	3.7	80
44	Optimal dietary therapy of long-chain 3-hydroxyacyl-CoA dehydrogenase deficiency. <i>Molecular Genetics and Metabolism</i> , 2003 , 79, 114-23	3.7	69
43	Metabolic control during exercise with and without medium-chain triglycerides (MCT) in children with long-chain 3-hydroxy acyl-CoA dehydrogenase (LCHAD) or trifunctional protein (TFP) deficiency. <i>Molecular Genetics and Metabolism</i> , 2006 , 89, 58-63	3.7	62
42	Dietary management of long-chain 3-hydroxyacyl-CoA dehydrogenase deficiency (LCHADD). A case report and survey. <i>Journal of Inherited Metabolic Disease</i> , 1999 , 22, 123-31	5.4	61
41	Triheptanoin versus trioctanoin for long-chain fatty acid oxidation disorders: a double blinded, randomized controlled trial. <i>Journal of Inherited Metabolic Disease</i> , 2017 , 40, 831-843	5.4	57
40	Effect of optimal dietary therapy upon visual function in children with long-chain 3-hydroxyacyl CoA dehydrogenase and trifunctional protein deficiency. <i>Molecular Genetics and Metabolism</i> , 2005 , 86, 124-33	3.7	56
39	Observations regarding retinopathy in mitochondrial trifunctional protein deficiencies. <i>Molecular Genetics and Metabolism</i> , 2012 , 106, 18-24	3.7	45
38	Substrate oxidation and cardiac performance during exercise in disorders of long chain fatty acid oxidation. <i>Molecular Genetics and Metabolism</i> , 2012 , 105, 110-5	3.7	40
37	Effects of higher dietary protein intake on energy balance and metabolic control in children with long-chain 3-hydroxy acyl-CoA dehydrogenase (LCHAD) or trifunctional protein (TFP) deficiency. <i>Molecular Genetics and Metabolism</i> , 2007 , 90, 64-9	3.7	31
36	Evidence for an association between infant mortality and a carnitine palmitoyltransferase 1A genetic variant. <i>Pediatrics</i> , 2010 , 126, 945-51	7.4	28
35	Prevalence and distribution of the c.1436C>T sequence variant of carnitine palmitoyltransferase 1A among Alaska Native infants. <i>Journal of Pediatrics</i> , 2011 , 158, 124-9	3.6	27
34	Effect of Aerobic and Resistance Exercise on Glycemic Control in Adults With Type 1 Diabetes. <i>Canadian Journal of Diabetes</i> , 2019 , 43, 406-414.e1	2.1	27
33	Increased vitamin E intake is associated with higher alpha-tocopherol concentration in the maternal circulation but higher alpha-carboxyethyl hydroxychroman concentration in the fetal circulation. <i>American Journal of Clinical Nutrition</i> , 2011 , 93, 368-73	7	26
32	Influence of high fat diet and resveratrol supplementation on placental fatty acid uptake in the Japanese macaque. <i>Placenta</i> , 2015 , 36, 903-10	3.4	24
31	Impaired fasting tolerance among Alaska native children with a common carnitine palmitoyltransferase 1A sequence variant. <i>Molecular Genetics and Metabolism</i> , 2011 , 104, 261-4	3.7	24

30	Association of a genetic variant of carnitine palmitoyltransferase 1A with infections in Alaska Native children. <i>Journal of Pediatrics</i> , 2013 , 163, 1716-21	3.6	21
29	Altered body composition and energy expenditure but normal glucose tolerance among humans with a long-chain fatty acid oxidation disorder. <i>American Journal of Physiology - Endocrinology and Metabolism</i> , 2013 , 305, E1299-308	6	16
28	NORMAL FATTY ACID CONCENTRATIONS IN YOUNG CHILDREN WITH PHENYLKETONURIA (PKU). <i>Topics in Clinical Nutrition</i> , 2009 , 24, 333-340	0.4	16
27	Unique plasma metabolomic signatures of individuals with inherited disorders of long-chain fatty acid oxidation. <i>Journal of Inherited Metabolic Disease</i> , 2016 , 39, 399-408	5.4	13
26	Characterization of Chorioretinopathy Associated with Mitochondrial Trifunctional Protein Disorders: Long-Term Follow-up of 21 Cases. <i>Ophthalmology</i> , 2016 , 123, 2183-95	7.3	13
25	Maternal Weight Gain Regulates Omega-3 Fatty Acids in Male, Not Female, Neonates. <i>Reproductive Sciences</i> , 2017 , 24, 560-567	3	9
24	EFFECT OF FEEDING, EXERCISE AND GENOTYPE ON PLASMA 3-HYDROXYACYLCARNITINES IN CHILDREN WITH LCHAD DEFICIENCY. <i>Topics in Clinical Nutrition</i> , 2009 , 24, 359-365	0.4	9
23	Blood cytokine patterns suggest a modest inflammation phenotype in subjects with long-chain fatty acid oxidation disorders. <i>Physiological Reports</i> , 2019 , 7, e14037	2.6	8
22	Normal vitamin D levels and bone mineral density among children with inborn errors of metabolism consuming medical food-based diets. <i>Nutrition Research</i> , 2016 , 36, 101-8	4	7
21	Metabolism and energy requirements in pantothenate kinase-associated neurodegeneration. <i>Molecular Genetics and Metabolism</i> , 2013 , 110, 336-41	3.7	7
20	Higher dietary protein intake preserves lean body mass, lowers liver lipid deposition, and maintains metabolic control in participants with long-chain fatty acid oxidation disorders. <i>Journal of Inherited Metabolic Disease</i> , 2019 , 42, 857-869	5.4	5
19	More Time in Glucose Range During Exercise Days than Sedentary Days in Adults Living with Type 1 Diabetes. <i>Diabetes Technology and Therapeutics</i> , 2021 , 23, 376-383	8.1	5
18	Neonatal fatty acid profiles are correlated with infant growth measures at 6 months. <i>Journal of Developmental Origins of Health and Disease</i> , 2017 , 8, 474-482	2.4	4
17	Nutrient intake, body composition, and blood phenylalanine control in children with phenylketonuria compared to healthy controls. <i>Molecular Genetics and Metabolism Reports</i> , 2020 , 23, 100599	1.8	4
16	Use of propofol for short duration procedures in children with long chain 3-hydroxyacyl-CoA dehydrogenase (LCHAD) or trifunctional protein (TFP) deficiencies. <i>Molecular Genetics and Metabolism</i> , 2014 , 112, 139-42	3.7	3
15	Change in postprandial substrate oxidation after a high-fructose meal is related to body mass index in healthy men. <i>Nutrition Research</i> , 2013 , 33, 435-41	4	3
14	A Ketogenic & Low-Protein Diet Slows Retinal Degeneration in rd10 Mice. <i>Translational Vision Science and Technology</i> , 2020 , 9, 18	3.3	2
13	Cardiac tissue citric acid cycle intermediates in exercised very long-chain acyl-CoA dehydrogenase-deficient mice fed triheptanoin or medium-chain triglyceride. <i>Journal of Inherited Metabolic Disease</i> , 2020 , 43, 1232-1242	5.4	1

12	Inborn errors of fatty acid oxidation	587-602		1
11	Effects of fasting, feeding and exercise on plasma acylcarnitines among subjects with CPT2D, VLCADD and LCHADD/TFPD. <i>Molecular Genetics and Metabolism</i> , 2020 , 131, 90-97		3.7	1
10	Maternal dietary fat intake during pregnancy and newborn body composition. <i>Journal of Perinatology</i> , 2021 , 41, 1007-1013		3.1	1
9	Response to Letter to the editor. <i>Journal of Inherited Metabolic Disease</i> , 2019 , 42, 396-397		5.4	
8	Comparison of Blood Glucose Monitoring Systems for Use in Insulin Clamp Studies During Either Intralipid or Glycerol Co-infusions. <i>Journal of Diabetes Science and Technology</i> , 2018 , 12, 232-233		4.1	
7	Reply: To PMID 23992672. <i>Journal of Pediatrics</i> , 2014 , 165, 420-2		3.6	
6	Reply: To PMID 23992672. <i>Journal of Pediatrics</i> , 2014 , 165, 419-20		3.6	
5	Effects of low and high carbohydrate meals on postprandial plasma lipid concentrations. <i>FASEB Journal</i> , 2009 , 23, 722.8		0.9	
4	Acute consumption of very low carbohydrate meals lowers postprandial concentrations of insulin, leptin, and total ghrelin. <i>FASEB Journal</i> , 2009 , 23, 720.3		0.9	
3	Vitamin E Deficiency in Children with Long-chain 3-hydroxyacylCoA dehydrogenase deficiency (LCHADD). <i>FASEB Journal</i> , 2009 , 23, 904.3		0.9	
2	0035 Resting Metabolism and the Metabolic Response to Exercise Follow Circadian Patterns with Day/Night Differences in Substrate Utilization Between Lean and Obese Adults. <i>Sleep</i> , 2020 , 43, A14-A14 ^{1.1}			
1	0222 A shift in the circadian timing of calories and an increase in sleep variability are associated with changes in cardiometabolic health in a real-world setting. <i>Sleep</i> , 2022 , 45, A101-A101			1.1