

# Fãtima V Ventura

## List of Publications by Year in descending order

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17  
papers

1,081  
citations

687363

13  
h-index

888059

17  
g-index

17  
all docs

17  
docs citations

17  
times ranked

1905  
citing authors

#	ARTICLE	IF	CITATIONS
1	Nanopharmaceutics: Part II&#x2014;Production Scales and Clinically Compliant Production Methods. <i>Nanomaterials</i> , 2020, 10, 455.	4.1	55
2	Unveiling the Pathogenic Molecular Mechanisms of the Most Common Variant (p.K329E) in Medium-Chain Acyl-CoA Dehydrogenase Deficiency by <i>in Vitro</i> and <i>in Silico</i> Approaches. <i>Biochemistry</i> , 2016, 55, 7086-7098.	2.5	5
3	Insights into Medium&#x2014;chain Acyl&#x2014;CoA Dehydrogenase Structure by Molecular Dynamics Simulations. <i>Chemical Biology and Drug Design</i> , 2016, 88, 281-292.	3.2	9
4	The Biochemistry and Physiology of Mitochondrial Fatty Acid $\hat{2}$ -Oxidation and Its Genetic Disorders. <i>Annual Review of Physiology</i> , 2016, 78, 23-44.	13.1	490
5	Retrospective study of the medium&#x2014;chain acyl&#x2014;CoA dehydrogenase deficiency in Portugal. <i>Clinical Genetics</i> , 2014, 85, 555-561.	2.0	11
6	Peroxisomes contribute to the acylcarnitine production when the carnitine shuttle is deficient. <i>Biochimica Et Biophysica Acta - Molecular and Cell Biology of Lipids</i> , 2013, 1831, 1467-1474.	2.4	74
7	Carnitine palmitoyltransferase 2 and carnitine/acylcarnitine translocase are involved in the mitochondrial synthesis and export of acylcarnitines. <i>FASEB Journal</i> , 2013, 27, 2039-2044.	0.5	58
8	Substrate specificity of human carnitine acetyltransferase: Implications for fatty acid and branched-chain amino acid metabolism. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2013, 1832, 773-779.	3.8	69
9	Carnitine palmitoyltransferase 2: New insights on the substrate specificity and implications for acylcarnitine profiling. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2010, 1802, 728-732.	3.8	37
10	Inhibition of adenine nucleotide transport in rat liver mitochondria by long-chain acyl-coenzyme A $\hat{2}$ -oxidation intermediates. <i>Biochemical and Biophysical Research Communications</i> , 2007, 352, 873-878.	2.1	14
11	Differential inhibitory effect of long-chain acyl-CoA esters on succinate and glutamate transport into rat liver mitochondria and its possible implications for long-chain fatty acid oxidation defects. <i>Molecular Genetics and Metabolism</i> , 2005, 86, 344-352.	1.1	34
12	Quantitative acylcarnitine profiling in fibroblasts using [U-13C] palmitic acid: an improved tool for the diagnosis of fatty acid oxidation defects. <i>Clinica Chimica Acta</i> , 1999, 281, 1-17.	1.1	79
13	Lactic acidosis in long-chain fatty acid $\hat{2}$ -oxidation disorders. <i>Journal of Inherited Metabolic Disease</i> , 1998, 21, 645-654.	3.6	31
14	Carnitine palmitoyltransferase II specificity towards beta-oxidation intermediates. Evidence for a reverse carnitine cycle in mitochondria. <i>FEBS Journal</i> , 1998, 253, 614-618.	0.2	30
15	Broad specificity of carnitine palmitoyltransferase II towards long-chain acyl-CoA $\hat{2}$ -oxidation intermediates and its practical approach to the synthesis of various long-chain acylcarnitines. <i>Journal of Inherited Metabolic Disease</i> , 1997, 20, 423-426.	3.6	5
16	Inhibitory effect of 3-hydroxyacyl-CoAs and other long-chain fatty acid $\hat{2}$ -oxidation intermediates on mitochondrial oxidative phosphorylation. <i>Journal of Inherited Metabolic Disease</i> , 1996, 19, 161-164.	3.6	40
17	Inhibition of oxidative phosphorylation by palmitoyl-CoA in digitonin permeabilized fibroblasts: implications for long-chain fatty acid $\hat{2}$ -oxidation disorders. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 1995, 1272, 14-20.	3.8	40