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Relation between fatty acid composition and clinical status or genotype in cystic fibrosis patients

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Annals of Nutrition and Metabolism, 2007, 51, 541-9.

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#	Paper	IF	Citations
31	Update on fat-soluble vitamins in cystic fibrosis. <i>Current Opinion in Pulmonary Medicine</i> , 2008 , 14, 574-81		46
30	Persistence of essential fatty acid deficiency in cystic fibrosis despite nutritional therapy. <i>Pediatric Research</i> , 2009 , 66, 585-9	3.2	45
29	Increased tissue arachidonic acid and reduced linoleic acid in a mouse model of cystic fibrosis are reversed by supplemental glycerophospholipids enriched in docosahexaenoic acid. <i>Journal of Nutrition</i> , 2009 , 139, 2358-64	4.1	25
28	The depressive effects of 5,8,11-eicosatrienoic Acid (20:3n-9) on osteoblasts. <i>Lipids</i> , 2009 , 44, 97-102	1.6	17
27	Structural equations to model relationships between pulmonary function, fatty acids and oxidation in cystic fibrosis. <i>Scandinavian Journal of Clinical and Laboratory Investigation</i> , 2009 , 69, 36-44	2	2
26	Lipid metabolism in cystic fibrosis. <i>Current Opinion in Clinical Nutrition and Metabolic Care</i> , 2009 , 12, 105-9	3.8	33
25	Fatty acid composition of serum phospholipids in cystic fibrosis (CF) patients with or without CF related liver disease. <i>Clinical Chemistry and Laboratory Medicine</i> , 2010 , 48, 1751-5	5.9	10
24	Whole blood fatty acid analysis with micromethod in cystic fibrosis and pulmonary disease. <i>Journal of Cystic Fibrosis</i> , 2010 , 9, 228-33	4.1	14
23	Increased elongase 6 and Δ -desaturase activity are associated with n-7 and n-9 fatty acid changes in cystic fibrosis. <i>Lipids</i> , 2011 , 46, 669-77	1.6	22
22	(n-3) long-chain PUFA differentially affect resistance to <i>Pseudomonas aeruginosa</i> infection of male and female <i>cftr</i> ^{-/-} mice. <i>Journal of Nutrition</i> , 2011 , 141, 1101-7	4.1	20
21	Specificity and rate of human and mouse liver and plasma phosphatidylcholine synthesis analyzed in vivo. <i>Journal of Lipid Research</i> , 2011 , 52, 399-407	6.3	97
20	DHA and EPA reverse cystic fibrosis-related FA abnormalities by suppressing FA desaturase expression and activity. <i>Journal of Lipid Research</i> , 2012 , 53, 257-65	6.3	52
19	Oxidative stress and antioxidant therapy in cystic fibrosis. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2012 , 1822, 690-713	6.9	140
18	Interactions of linoleic and alpha-linolenic acids in the development of fatty acid alterations in cystic fibrosis. <i>Lipids</i> , 2013 , 48, 333-42	1.6	12
17	Abnormal n-6 fatty acid metabolism in cystic fibrosis is caused by activation of AMP-activated protein kinase. <i>Journal of Lipid Research</i> , 2014 , 55, 1489-97	6.3	15
16	Omega-3 Fatty Acids and Cystic Fibrosis. 2015 , 383-387		
15	Essential Fatty Acid Deficiency in Cystic Fibrosis: Malabsorption or Metabolic Abnormality?. 2015 , 365-371		

14	Polyunsaturated fatty acid supplementation reverses cystic fibrosis-related fatty acid abnormalities in CFTR ^{-/-} mice by suppressing fatty acid desaturases. <i>Journal of Nutritional Biochemistry</i> , 2015 , 26, 36-43	6.3	7
13	Effect of Oral Lipid Matrix Supplement on Fat Absorption in Cystic Fibrosis: A Randomized Placebo-Controlled Trial. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2016 , 63, 676-680	2.8	7
12	The clinical benefits of long-term supplementation with omega-3 fatty acids in cystic fibrosis patients - A pilot study. <i>Prostaglandins Leukotrienes and Essential Fatty Acids</i> , 2016 , 108, 45-50	2.8	17
11	Complex Relation Between Diet and Phospholipid Fatty Acids in Children With Cystic Fibrosis. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2017 , 64, 598-604	2.8	5
10	The effects of ivacaftor on CF fatty acid metabolism: An analysis from the GOAL study. <i>Journal of Cystic Fibrosis</i> , 2017 , 16, 132-138	4.1	11
9	Low linoleic and high docosahexaenoic acids in a severe phenotype of transgenic cystic fibrosis mice. <i>Experimental Biology and Medicine</i> , 2018 , 243, 496-503	3.7	4
8	Fatty acid profile in erythrocytes associated with serum cytokines in pediatric cystic fibrosis patients. <i>Revista De Nutricao</i> , 2018 , 31, 455-466	1.8	1
7	Glued in lipids: Lipointoxication in cystic fibrosis. <i>EBioMedicine</i> , 2020 , 61, 103038	8.8	1
6	Impaired Ratio of Unsaturated to Saturated Non-Esterified Fatty Acids in Saliva from Patients with Cystic Fibrosis. <i>Diagnostics</i> , 2020 , 10,	3.8	1
5	Abnormal n-6 fatty acid metabolism in cystic fibrosis contributes to pulmonary symptoms. <i>Prostaglandins Leukotrienes and Essential Fatty Acids</i> , 2020 , 160, 102156	2.8	5
4	Circulating biomarkers of antioxidant status and oxidative stress in people with cystic fibrosis: A systematic review and meta-analysis. <i>Redox Biology</i> , 2020 , 32, 101436	11.3	22
3	Influence of lung transplantation on the essential fatty acid profile in cystic fibrosis. <i>Prostaglandins Leukotrienes and Essential Fatty Acids</i> , 2020 , 158, 102060	2.8	2
2	Nutrition in Cystic Fibrosis-Some Notes on the Fat Recommendations.. <i>Nutrients</i> , 2022 , 14,	6.7	0
1	The fatty acid imbalance of cystic fibrosis exists at birth independent of feeding in pig and ferret models. 2022 , 136, 1773-1791		1