Aleksandra Lisowska

List of Publications by Year
in descending order

60	612	687220 13	713332 21
papers	citations	h-index	g-index

60	60	60	832
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	The changing face of the exocrine pancreas in cystic fibrosis: pancreatic sufficiency, pancreatitis and genotype. European Journal of Gastroenterology and Hepatology, 2008, 20, 157-160.	0.8	59
2	Small intestine bacterial overgrowth is frequent in cystic fibrosis: combined hydrogen and methane measurements are required for its detection. Acta Biochimica Polonica, 2009, 56, 631-4.	0.3	37
3	Green tea extract decreases starch digestion and absorption from a test meal in humans: a randomized, placebo-controlled crossover study. Scientific Reports, 2015, 5, 12015.	1.6	33
4	Oral antibiotic therapy improves fat absorption in cystic fibrosis patients with small intestine bacterial overgrowth. Journal of Cystic Fibrosis, 2011, 10, 418-421.	0.3	32
5	Small intestine bacterial overgrowth does not correspond to intestinal inflammation in cystic fibrosis. Scandinavian Journal of Clinical and Laboratory Investigation, 2010, 70, 322-326.	0.6	29
6	Mulberry leaf extract decreases digestion and absorption of starch in healthy subjectsâ€”A randomized, placebo-controlled, crossover study. Advances in Medical Sciences, 2017, 62, 302-306.	0.9	27
7	Cystic fibrosis dyslipidaemia: A cross-sectional study. Journal of Cystic Fibrosis, 2019, 18, 566-571.	0.3	24
8	Cystic fibrosis is a risk factor for celiac disease. Acta Biochimica Polonica, 2010, 57, 115-8.	0.3	22
9	Adult-type hypolactasia and lactose malabsorption in Poland.. Acta Biochimica Polonica, 2010, 57, .	0.3	18
10	Gastroesophageal Reflux Disease in Children with Cystic Fibrosis. Advances in Experimental Medicine and Biology, 2015, 873, 1-7.	0.8	17
11	Single dose of green tea extract decreases lipid digestion and absorption from a test meal in humans.. Acta Biochimica Polonica, 2013, 60, .	0.3	17
12	Polyunsaturated Fatty Acids in Cystic Fibrosis Are Related to Nutrition and Clinical Expression of the Disease. Journal of Pediatric Gastroenterology and Nutrition, 2007, 45, 488-489.	0.9	14
13	Twelve weeks CLA supplementation decreases the hip circumference in overweight and obese women. A double-blind, randomized, placebo-controlled trial. Acta Scientiarum Polonorum, Technologia Alimentaria, 2016, 15, 107-113.	0.2	14
14	Chronic pouchitis is not related to small intestine bacterial overgrowth. Inflammatory Bowel Diseases, 2008, 14, 1102-1104.	0.9	13
15	Supplementation of ursodeoxycholic acid improves fat digestion and absorption in cystic fibrosis patients with mild liver involvement. European Journal of Gastroenterology and Hepatology, 2016, 28, 645-649.	0.8	13
16	Body Fat Changes and Liver Safety in Obese and Overweight Women Supplemented with Conjugated Linoleic Acid: A 12-Week Randomised, Double-Blind, Placebo-Controlled Trial. Nutrients, 2020, 12, 1811.	1.7	13
17	The effect of endurance and endurance-strength training on body composition and cardiometabolic markers in abdominally obese women: a randomised trial. Scientific Reports, 2021, 11, 12339.	1.6	13
18	Fat-Soluble Vitamin Supplementation Using Liposomes, Cyclodextrins, or Medium-Chain Triglycerides in Cystic Fibrosis: A Randomized Controlled Trial. Nutrients, 2021, 13, 4554.	1.7	12

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19	Adaptive changes of pancreatic protease secretion to a short-term vegan diet: influence of reduced intake and modification of protein. British Journal of Nutrition, 2012, 107, 272-276.	1.2	11
20	Pancreatic Elastase-1 Quick Test for rapid assessment of pancreatic status in cystic fibrosis patients. Journal of Cystic Fibrosis, 2016, 15, 664-668.	0.3	11
21	Increased Soluble VCAM-1 and Normal P-Selectin in Cystic Fibrosis: a Cross-Sectional Study. Lung, 2017, 195, 445-453.	1.4	11
22	Lactose malabsorption is a risk factor for decreased bone mineral density in pancreatic insufficient cystic fibrosis patients. European Journal of Human Genetics, 2012, 20, 1092-1095.	1.4	10
23	Serum Lipase After Secretin Stimulation Detects Mild Pancreatic Involvement in Cystic Fibrosis. Journal of Pediatric Gastroenterology and Nutrition, 2004, 38, 430-435.	0.9	9
24	Mild CFTR mutations and genetic predisposition to lactase persistence in cystic fibrosis. European Journal of Human Genetics, 2011, 19, 748-752.	1.4	9
25	Vitamin K status in cystic fibrosis patients with liver cirrhosis. Digestive and Liver Disease, 2017, 49, 672-675.	0.4	9
26	Determinants of Serum Glycerophospholipid Fatty Acids in Cystic Fibrosis. International Journal of Molecular Sciences, 2017, 18, 185.	1.8	9
27	Endurance Training Depletes Antioxidant System but Does Not Affect Endothelial Functions in Women with Abdominal Obesity: A Randomized Trial with a Comparison to Endurance-Strength Training. Journal of Clinical Medicine, 2021, 10, 1639.	1.0	9
28	Adult-type hypolactasia and lactose malabsorption in Poland. Acta Biochimica Polonica, 2010, 57, 585-8.	0.3	9
29	Acid steatocrit determination is not helpful in cystic fibrosis patients without or with mild steatorrhea. Pediatric Pulmonology, 2010, 45, 249-254.	1.0	7
30	Comparison of fecal pyruvate kinase isoform M2 and calprotectin in acute diarrhea in hospitalized children. Scientific Reports, 2014, 4, 4769.	1.6	7
31	Serum Phospholipid Fatty Acid Composition in Cystic Fibrosis Patients with and without Liver Cirrhosis. Annals of Nutrition and Metabolism, 2017, 71, 91-98.	1.0	7
32	No Difference in Lactoferrin Levels between Metabolically Healthy and Unhealthy Obese Women. Nutrients, 2019, 11, 1976.	1.7	7
33	Preclinical atherosclerosis in cystic fibrosis: Two distinct presentations are related to pancreatic status. Journal of Cystic Fibrosis, 2022, 21, 26-33.	0.3	7
34	Single dose of green tea extract decreases lipid digestion and absorption from a test meal in humans. Acta Biochimica Polonica, 2013, 60, 481-3.	0.3	7
35	Esophageal pH Monitoring in Children. Journal of Pediatric Gastroenterology and Nutrition, 2015, 61, 212-214.	0.9	6
36	The elimination of meat from the diet selectively decreases pancreatic elastase secretion. British Journal of Nutrition, 2007, 98, 154-158.	1.2	5

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37	Comparison of Subjective and Objective Methods to Measure the Physical Activity of Non-Depressed Middle-Aged Healthy Subjects with Normal Cognitive Function and Mild Cognitive Impairmentâ€"A Cross-Sectional Study. International Journal of Environmental Research and Public Health, 2021, 18, 8042.	1.2	5
38	Antibiotic therapy and fat digestion and absorption in cystic fibrosis.. Acta Biochimica Polonica, 2011, 58, .	0.3	5
39	Unresponsive or non-compliant steatorrhea in cystic fibrosis?. Journal of Cystic Fibrosis, 2006, 5, 253-255.	0.3	4
40	Exocrine pancreatic function in children with Alagille syndrome. Scientific Reports, 2016, 6, 35229.	1.6	4
41	Comparison of the Effect of Amaranth Oil vs. Rapeseed Oil on Selected Atherosclerosis Markers in Overweight and Obese Subjects: A Randomized Double-Blind Cross-Over Trial. International Journal of Environmental Research and Public Health, 2021, 18, 8540.	1.2	4
42	Lactose intolerance, lactose malabsorption and genetic predisposition to adult-type hypolactasia in patients after restorative proctocolectomy. Acta Biochimica Polonica, 2019, 66, 173-175.	0.3	4
43	Comparison of the effects of endurance and enduranceâ€'strength training programmes on the level of endothelial dysfunction in women with abdominal obesity: study protocol for aÁrandomised controlled trial. Journal of Medical Science, 2019, 88, 266-272.	0.2	4
44	Fat-Soluble Vitamins in Standard vs. Liposomal Form Enriched with Vitamin K2 in Cystic Fibrosis: A Randomized Multi-Center Trial. Journal of Clinical Medicine, 2022, 11, 462.	1.0	4
45	Conjugated linoleic acid does not affect digestion and absorption of fat and starchâ€"a randomized, double-blinded, placebo-controlled parallel study. Journal of Breath Research, 2018, 12, 016010.	1.5	3
46	Severe Genotype, Pancreatic Insufficiency and Low Dose of Pancreatic Enzymes Associate with Abnormal Serum Sterol Profile in Cystic Fibrosis. Biomolecules, 2021, 11, 313.	1.8	3
47	Small intestinal bacterial overgrowth in patients with progressive familial intrahepatic cholestasis. Acta Biochimica Polonica, 2014, 61, 103-7.	0.3	3
48	Pancreatic enzyme therapy and gastrointestinal symptoms in patients with cystic fibrosis. Journal of Pediatrics, 2005, 147, 870-871.	0.9	2
49	Re: Fecal Elastase: Pancreatic Status Verification and Influence on Nutritional Status in Children with Cystic Fibrosis.. Journal of Pediatric Gastroenterology and Nutrition, 2006, 42, 117.	0.9	2
50	Gastroesophageal reflux is not associated with short-term variability of parasympathetic activity in children. Advances in Medical Sciences, 2017, 62, 103-109.	0.9	2
51	Amaranth Oil Increases Total and LDL Cholesterol Levels without Influencing Early Markers of Atherosclerosis in an Overweight and Obese Population: A Randomized Double-Blind Cross-Over Study in Comparison with Rapeseed Oil Supplementation. Nutrients, 2019, 11, 3069.	1.7	2
52	The Effect of Endurance and Endurance-Strength Training on Bone Mineral Density and Content in Abdominally Obese Postmenopausal Women: A Randomized Trial. Healthcare (Switzerland), 2021, 9, 1074.	1.0	2
53	Smaller Width of the Pancreatic Duct During Secretin-Enhanced Magnetic Resonance Cholangiopancreatography in Pancreatic-Sufficient Cystic Fibrosis Patients. Pancreas, 2016, 45, 1175-1178.	0.5	1
54	Normal levels of serum pancreatic enzymes in patients with progressive familial intrahepatic cholestasis type 2. Acta Biochimica Polonica, 2010, 57, 573-5.	0.3	1

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55	Znaczenie witaminy K w mukowiscydozie. Pediatria Polska, 2009, 84, 265-269.	0.1	0
56	Early intestinal manifestation in infants with cystic fibrosis â€“ a pilot study. Przeglad Gastroenterologiczny, 2010, 3, 164-167.	0.3	0
57	Leukocyte Telomere Length Is Not Reduced in Children and Adults with Cystic Fibrosis but Associates with Clinical Characteristicsâ€”A Cross-Sectional Study. Journal of Clinical Medicine, 2021, 10, 590.	1.0	0
58	Helicobacter pylori Infection in Children with Phenylketonuria Does Not Depend on Metabolic Control and Is Not More Frequent Than in Healthy Subjectsâ€”A Cross-Sectional Study. Children, 2021, 8, 713.	0.6	0
59	Antibiotic therapy and fat digestion and absorption in cystic fibrosis. Acta Biochimica Polonica, 2011, 58, 345-7.	0.3	0
60	Fecal pyruvate kinase is not suitable for discrimination between inflammatory bowel disease exacerbation and acute gastroenteritis. Medycyna Wieku Rozwojowego, 2015, 19, 167-73.	0.2	0