

Lucia Sacchetti

List of Publications by Year in descending order

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Version: 2024-02-01

82
papers

2,911
citations

159525

30
h-index

182361

51
g-index

83
all docs

83
docs citations

83
times ranked

4957
citing authors

#	ARTICLE	IF	CITATIONS
1	Promelaxin Microenemas Are Non-inferior to Oral Polyethylene Glycol for the Treatment of Functional Constipation in Young Children: A Randomized Clinical Trial. <i>Frontiers in Pediatrics</i> , 2021, 9, 753938.	0.9	7
2	Gut microbiome investigation in celiac disease: from methods to its pathogenetic role. <i>Clinical Chemistry and Laboratory Medicine</i> , 2020, 58, 340-349.	1.4	33
3	Setup of Quantitative PCR for Oral <i>Neisseria</i> spp. Evaluation in Celiac Disease Diagnosis. <i>Diagnostics</i> , 2020, 10, 12.	1.3	6
4	Genetic analysis resolves differential diagnosis of a familial syndromic dilated cardiomyopathy: A new case of Alström syndrome. <i>Molecular Genetics & Genomic Medicine</i> , 2020, 8, e1260.	0.6	22
5	Characterization of the Duodenal Mucosal Microbiome in Obese Adult Subjects by 16S rRNA Sequencing. <i>Microorganisms</i> , 2020, 8, 485.	1.6	36
6	Very light physical activity amount in FTO genetically predisposed obese individuals. <i>Sport Sciences for Health</i> , 2019, 15, 689-697.	0.4	0
7	Celiac disease-associated <i>Neisseria flavescens</i> decreases mitochondrial respiration in CaCo2 epithelial cells: Impact of <i>Lactobacillus paracasei</i> CBA L74 on bacterial-induced cellular imbalance. <i>Cellular Microbiology</i> , 2019, 21, e13035.	1.1	21
8	Helper-dependent adenovirus-mediated gene transfer of a secreted LDL receptor/transferrin chimeric protein reduces aortic atherosclerosis in LDL receptor-deficient mice. <i>Gene Therapy</i> , 2019, 26, 121-130.	2.3	9
9	Effect of <i>CYP4F2</i> , <i>VKORC1</i> , and <i>CYP2C9</i> in Influencing Coumarin Dose: A Single-Patient Data Meta-Analysis in More Than 15,000 Individuals. <i>Clinical Pharmacology and Therapeutics</i> , 2019, 105, 1477-1491.	2.3	23
10	Altered Bioenergetic Profile in Umbilical Cord and Amniotic Mesenchymal Stem Cells from Newborns of Obese Women. <i>Stem Cells and Development</i> , 2018, 27, 199-206.	1.1	17
11	Oropharyngeal microbiome evaluation highlights <i>Neisseria</i> abundance in active celiac patients. <i>Scientific Reports</i> , 2018, 8, 11047.	1.6	33
12	Impaired Enterohormone Response Following a Liquid Test Meal in Gastrectomized Patients. <i>Annals of Nutrition and Metabolism</i> , 2017, 71, 211-216.	1.0	3
13	miR-138/miR-222 Overexpression Characterizes the miRNome of Amniotic Mesenchymal Stem Cells in Obesity. <i>Stem Cells and Development</i> , 2017, 26, 4-14.	1.1	17
14	The Cause of Death of a Child in the 18th Century Solved by Bone Microbiome Typing Using Laser Microdissection and Next Generation Sequencing. <i>International Journal of Molecular Sciences</i> , 2017, 18, 109.	1.8	10
15	Changes in the MicroRNA Profile Observed in the Subcutaneous Adipose Tissue of Obese Patients after Laparoscopic Adjustable Gastric Banding. <i>Journal of Obesity</i> , 2017, 2017, 1-6.	1.1	26
16	Metagenomics Reveals Dysbiosis and a Potentially Pathogenic <i>N. flavescens</i> Strain in Duodenum of Adult Celiac Patients. <i>American Journal of Gastroenterology</i> , 2016, 111, 879-890.	0.2	128
17	Orexin-A represses satiety-inducing POMC neurons and contributes to obesity via stimulation of endocannabinoid signaling. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016, 113, 4759-4764.	3.3	68
18	High Frequency of Haplotype HLA-DQ7 in Celiac Disease Patients from South Italy: Retrospective Evaluation of 5,535 Subjects at Risk of Celiac Disease. <i>PLoS ONE</i> , 2015, 10, e0138324.	1.1	23

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19	Immune-metabolic profiling of anorexic patients reveals an anti-oxidant and anti-inflammatory phenotype. <i>Metabolism: Clinical and Experimental</i> , 2015, 64, 396-405.	1.5	37
20	Catechol-O-Methyltransferase (COMT) Gene Polymorphisms as Risk Factor in Temporomandibular Disorders Patients From Southern Italy. <i>Clinical Journal of Pain</i> , 2014, 30, 129-133.	0.8	31
21	The FTO gene polymorphism (rs9939609) is associated with metabolic syndrome in morbidly obese subjects from southern Italy. <i>Molecular and Cellular Probes</i> , 2014, 28, 195-199.	0.9	23
22	Laparoscopic Adjustable Gastric Banding Reduces Subcutaneous Adipose Tissue and Blood Inflammation in Nondiabetic Morbidly Obese Individuals. <i>Obesity Surgery</i> , 2014, 24, 2161-2168.	1.1	3
23	An Altered Gut Microbiome Profile in a Child Affected by Crohn's Disease Normalized After Nutritional Therapy. <i>American Journal of Gastroenterology</i> , 2013, 108, 851-852.	0.2	54
24	High Aminopeptidase N/CD13 Levels Characterize Human Amniotic Mesenchymal Stem Cells and Drive Their Increased Adipogenic Potential in Obese Women. <i>Stem Cells and Development</i> , 2013, 22, 2287-2297.	1.1	19
25	Haplogroup T Is an Obesity Risk Factor: Mitochondrial DNA Haplotyping in a Morbid Obese Population from Southern Italy. <i>BioMed Research International</i> , 2013, 2013, 1-5.	0.9	37
26	Warfarin Anticoagulant Therapy: A Southern Italy Pharmacogenetics-Based Dosing Model. <i>PLoS ONE</i> , 2013, 8, e71505.	1.1	31
27	Identification of Candidate Children for Maturity-Onset Diabetes of the Young Type 2 (MODY2) Gene Testing: A Seven-Item Clinical Flowchart (7-iF). <i>PLoS ONE</i> , 2013, 8, e79933.	1.1	33
28	miRNA and Protein Expression Profiles of Visceral Adipose Tissue Reveal miR-141/YWHAG and miR-520e/RAB11A as Two Potential miRNA/Protein Target Pairs Associated with Severe Obesity. <i>Journal of Proteome Research</i> , 2012, 11, 3358-3369.	1.8	53
29	Interleukin-1 β Causes Anxiety by Interacting with the Endocannabinoid System. <i>Journal of Neuroscience</i> , 2012, 32, 13896-13905.	1.7	96
30	Mitochondrial Diabetes in Children: Seek and You Will Find It. <i>PLoS ONE</i> , 2012, 7, e34956.	1.1	28
31	Glucokinase (GCK) Mutations and Their Characterization in MODY2 Children of Southern Italy. <i>PLoS ONE</i> , 2012, 7, e38906.	1.1	37
32	MicroRNA-449a Overexpression, Reduced NOTCH1 Signals and Scarce Goblet Cells Characterize the Small Intestine of Celiac Patients. <i>PLoS ONE</i> , 2011, 6, e29094.	1.1	85
33	Sequence Analysis of the <i>UCP1</i> Gene in a Severe Obese Population from Southern Italy. <i>Journal of Obesity</i> , 2011, 2011, 1-4.	1.1	15
34	Loss of striatal cannabinoid CB1 receptor function in attention deficit/hyperactivity disorder mice with point mutation of the dopamine transporter. <i>European Journal of Neuroscience</i> , 2011, 34, 1369-1377.	1.2	23
35	High Leptin/Adiponectin Ratio and Serum Triglycerides Are Associated With an "At-Risk" Phenotype in Young Severely Obese Patients. <i>Obesity</i> , 2011, 19, 1492-1496.	1.5	64
36	Improving the Estimation of Celiac Disease Sibling Risk by Non-HLA Genes. <i>PLoS ONE</i> , 2011, 6, e26920.	1.1	24

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37	A multilayer perceptron neural network-based approach for the identification of responsiveness to interferon therapy in multiple sclerosis patients. <i>Information Sciences</i> , 2010, 180, 4153-4163.	4.0	28
38	miR-19d Overexpression Is Associated With Human Obesity. <i>Obesity</i> , 2010, 18, 2170-2176.	1.5	149
39	Preservation of Striatal Cannabinoid CB1 Receptor Function Correlates with the Antianxiety Effects of Fatty Acid Amide Hydrolase Inhibition. <i>Molecular Pharmacology</i> , 2010, 78, 260-268.	1.0	73
40	Brain-Derived Neurotrophic Factor Controls Cannabinoid CB1 Receptor Function in the Striatum. <i>Journal of Neuroscience</i> , 2010, 30, 8127-8137.	1.7	59
41	Voluntary Exercise and Sucrose Consumption Enhance Cannabinoid CB1 Receptor Sensitivity in the Striatum. <i>Neuropsychopharmacology</i> , 2010, 35, 374-387.	2.8	74
42	Nonsynonymous mutation of catechol-O-methyl-transferase (COMT) gene in a patient with temporomandibular disorder. <i>Progress in Orthodontics</i> , 2010, 11, 174-179.	1.3	3
43	Functional analysis of melanocortin-4-receptor mutants identified in severely obese subjects living in Southern Italy. <i>Gene</i> , 2010, 457, 35-41.	1.0	13
44	Adaptations of Striatal Endocannabinoid System During Stress. <i>Molecular Neurobiology</i> , 2009, 39, 178-184.	1.9	7
45	Homocysteine levels and sustained virological response to pegylated interferon α 2b plus ribavirin therapy for chronic hepatitis C: a prospective study. <i>Liver International</i> , 2009, 29, 248-252.	1.9	30
46	Caffeine drinking potentiates cannabinoid transmission in the striatum: Interaction with stress effects. <i>Neuropharmacology</i> , 2009, 56, 590-597.	2.0	23
47	Connectivity Between Posterior Parietal Cortex and Ipsilateral Motor Cortex Is Altered in Schizophrenia. <i>Biological Psychiatry</i> , 2008, 64, 815-819.	0.7	51
48	Multiple Sclerosis and Hepatitis C Virus Infection Are Associated with Single Nucleotide Polymorphisms in Interferon Pathway Genes. <i>Journal of Interferon and Cytokine Research</i> , 2008, 28, 141-152.	0.5	42
49	Decreased Paraoxonase-2 Expression in Human Carotids During the Progression of Atherosclerosis. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2008, 28, 594-600.	1.1	34
50	Age-Related Reference Intervals of the Main Biochemical and Hematological Parameters in C57BL/6J, 129SV/EV and C3H/HeJ Mouse Strains. <i>PLoS ONE</i> , 2008, 3, e3772.	1.1	101
51	Glucokinase Gene Mutations: Structural and Genotype-Phenotype Analyses in MODY Children from South Italy. <i>PLoS ONE</i> , 2008, 3, e1870.	1.1	44
52	HLA related genetic risk for coeliac disease. <i>Gut</i> , 2007, 56, 1054-1059.	6.1	94
53	Hereditary Fructose Intolerance and Celiac Disease: A Novel Genetic Association. <i>Clinical Gastroenterology and Hepatology</i> , 2006, 4, 635-638.	2.4	13
54	The mtDNA 15497 G/A polymorphism in cytochrome b in severe obese subjects from Southern Italy. <i>Nutrition, Metabolism and Cardiovascular Diseases</i> , 2006, 16, 466-470.	1.1	12

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55	Metabolic syndrome: Analogies with subtypes of protein energy malnutrition. <i>Clinical Nutrition</i> , 2006, 25, 537-539.	2.3	4
56	Comparison of the TaqMan and LightCycler systems in pharmacogenetic testing: evaluation of CYP2C9*2/*3 polymorphisms. <i>Clinical Chemistry and Laboratory Medicine</i> , 2006, 44, 285-7.	1.4	8
57	Carotid Artery Remodeling in Middle-Aged Women With the Metabolic Syndrome (from the "Progetto) Tj ETQq1 1 0.784314 rgBT 0.7 32	1.1	32
58	Six Novel Mutations in the Proopiomelanocortin and Melanocortin Receptor 4 Genes in Severely Obese Adults Living in Southern Italy. <i>Clinical Chemistry</i> , 2005, 51, 1358-1364.	1.5	47
59	Y-Chromosome short tandem repeat (STR) haplotypes in a Campania population sample. <i>Clinical Chemistry and Laboratory Medicine</i> , 2005, 43, 163-6.	1.4	0
60	Paraoxonase and Superoxide Dismutase Gene Polymorphisms and Noise-Induced Hearing Loss. <i>Clinical Chemistry</i> , 2004, 50, 2012-2018.	1.5	100
61	Direct Detection of Exon Deletions/Duplications in Female Carriers of and Male Patients with Duchenne/Becker Muscular Dystrophy. <i>Clinical Chemistry</i> , 2004, 50, 1435-1438.	1.5	17
62	A paraoxonase gene polymorphism, PON 1 (55), as an independent risk factor for increased carotid intima-media thickness in middle-aged women. <i>Atherosclerosis</i> , 2003, 167, 141-148.	0.4	50
63	Radical-Trapping Activity, Blood Pressure, and Carotid Enlargement in Women. <i>Hypertension</i> , 2003, 41, 289-296.	1.3	9
64	Serum oxidative and antioxidant parameters in a group of Italian patients with age-related maculopathy. <i>Clinica Chimica Acta</i> , 2002, 320, 111-115.	0.5	43
65	Association of Obesity and Central Fat Distribution With Carotid Artery Wall Thickening in Middle-Aged Women. <i>Stroke</i> , 2002, 33, 2923-2928.	1.0	181
66	Dietary and circulating antioxidant vitamins in relation to carotid plaques in middle-aged women. <i>American Journal of Clinical Nutrition</i> , 2002, 76, 582-587.	2.2	40
67	Site-Specific Atherosclerotic Plaques in the Carotid Arteries of Middle-Aged Women From Southern Italy. <i>Stroke</i> , 2001, 32, 1953-1959.	1.0	49
68	Serum $\hat{3}$ -Glutamyltransferase Isoform Complexed to LDL in the Diagnosis of Small Hepatocellular Carcinoma. <i>Clinical Chemistry</i> , 1999, 45, 1100a-1102.	1.5	6
69	IgA antibodies to tissue transglutaminase: An effective diagnostic test for celiac disease. <i>Journal of Pediatrics</i> , 1999, 134, 166-171.	0.9	183
70	Discrimination between Celiac and Other Gastrointestinal Disorders in Childhood by Rapid Human Lymphocyte Antigen Typing. <i>Clinical Chemistry</i> , 1998, 44, 1755-1757.	1.5	9
71	Multivariate discriminant analysis of biochemical parameters for the differentiation of clinically confounding liver diseases. <i>Clinica Chimica Acta</i> , 1997, 257, 41-58.	0.5	8
72	Serum Mn-Superoxide Dismutase in Acute Myocardial Infarction. <i>Clinical Biochemistry</i> , 1997, 30, 569-571.	0.8	9

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73	Problems and perspectives of clinical biochemistry training, and the example of Italy. <i>Clinica Chimica Acta</i> , 1996, 245, 113-124.	0.5	1
74	A quantitative polymerase chain reaction (PCR) assay completely discriminates between Duchenne and Becker muscular dystrophy deletion carriers and normal females. <i>Molecular and Cellular Probes</i> , 1996, 10, 129-137.	0.9	16
75	Allele frequency distributions at several variable number of tandem repeat (VNTR) and short tandem repeat (STR) loci in a restricted Caucasian population from South Italy and their evaluation for paternity and forensic use. <i>Molecular and Cellular Probes</i> , 1996, 10, 299-308.	0.9	12
76	Serum pseudouridine in the diagnosis of acute leukaemias and as a novel prognostic indicator in acute lymphoblastic leukaemia. <i>Clinical Biochemistry</i> , 1993, 26, 513-520.	0.8	15
77	Serum type-2 macro-creatine kinase isoenzyme is not a useful marker of severe liver diseases or neoplasia. <i>Clinical Biochemistry</i> , 1990, 23, 523-527.	0.8	11
78	Comparison of seven serum assays on four automatic analysers. <i>Journal of Automated Methods and Management in Chemistry</i> , 1989, 11, 124-128.	0.4	0
79	Electrophoretic behavior and partial characterization of disease-associated serum forms of gammaglutamyltransferase. <i>Electrophoresis</i> , 1989, 10, 619-627.	1.3	8
80	Evaluation of pancreatic amylase immunoassay in acute pancreatitis. <i>Clinica Chimica Acta</i> , 1989, 183, 95-100.	0.5	8
81	The Serum Gamma-glutamyltransferase Isoenzyme System and its Diagnostic Role in Hepatobiliary Diseases. <i>Progress in Clinical Biochemistry and Medicine</i> , 1989, , 17-46.	0.5	1
82	Diagnostic efficiency in discriminating liver malignancies from cirrhosis by serum gamma-glutamyltransferase isoforms. <i>Clinica Chimica Acta</i> , 1988, 177, 167-172.	0.5	19