Joussef Hayek

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79 2,085 28 43 g-index

83 2,504 4.8 4.28 ext. papers ext. citations avg, IF L-index

#	Paper	IF	Citations
79	A comprehensive molecular study on Coffin-Siris and Nicolaides-Baraitser syndromes identifies a broad molecular and clinical spectrum converging on altered chromatin remodeling. <i>Human Molecular Genetics</i> , 2013 , 22, 5121-35	5.6	138
78	Systemic oxidative stress in classic Rett syndrome. Free Radical Biology and Medicine, 2009, 47, 440-8	7.8	108
77	Di-(2-ethylhexyl) phthalate and autism spectrum disorders. ASN Neuro, 2012 , 4, 223-9	5.3	100
76	Oxidative brain damage in Mecp2-mutant murine models of Rett syndrome. <i>Neurobiology of Disease</i> , 2014 , 68, 66-77	7.5	86
75	The role of oxidative stress in Rett syndrome: an overview. <i>Annals of the New York Academy of Sciences</i> , 2012 , 1259, 121-35	6.5	72
74	F2-dihomo-isoprostanes as potential early biomarkers of lipid oxidative damage in Rett syndrome. Journal of Lipid Research, 2011 , 52, 2287-2297	6.3	72
73	Altered gut microbiota in Rett syndrome. <i>Microbiome</i> , 2016 , 4, 41	16.6	69
72	Partial rescue of Rett syndrome by B polyunsaturated fatty acids (PUFAs) oil. <i>Genes and Nutrition</i> , 2012 , 7, 447-58	4.3	68
71	FEheuroprostanes mediate neurological severity in Rett syndrome. Clinica Chimica Acta, 2011 , 412, 1399	9 -4 . <u>0</u> 6	63
70	Oxidative stress in Rett syndrome: natural history, genotype, and variants. <i>Redox Report</i> , 2011 , 16, 145-	53 .9	59
69	Increased levels of 4HNE-protein plasma adducts in Rett syndrome. Clinical Biochemistry, 2011, 44, 368-	7 3 1 5	56
68	Cholesterol metabolism is altered in Rett syndrome: a study on plasma and primary cultured fibroblasts derived from patients. <i>PLoS ONE</i> , 2014 , 9, e104834	3.7	48
67	Subclinical inflammatory status in Rett syndrome. <i>Mediators of Inflammation</i> , 2014 , 2014, 480980	4.3	45
66	Revealing the complexity of a monogenic disease: rett syndrome exome sequencing. <i>PLoS ONE</i> , 2013 , 8, e56599	3.7	45
65	Genes related to mitochondrial functions, protein degradation, and chromatin folding are differentially expressed in lymphomonocytes of Rett syndrome patients. <i>Mediators of Inflammation</i> , 2013, 2013, 137629	4.3	44
64	Cytokine Dysregulation in MECP2- and CDKL5-Related Rett Syndrome: Relationships with Aberrant Redox Homeostasis, Inflammation, and B PUFAs. <i>Oxidative Medicine and Cellular Longevity</i> , 2015 , 2015, 421624	6.7	42
63	Scavenger receptor B1 post-translational modifications in Rett syndrome. FEBS Letters, 2013, 587, 2199	9-3.84	42

62	Cytokines profile and peripheral blood mononuclear cells morphology in Rett and autistic patients. <i>Cytokine</i> , 2016 , 77, 180-8	4	39
61	Morphological changes and oxidative damage in Rett Syndrome erythrocytes. <i>Biochimica Et Biophysica Acta - General Subjects</i> , 2012 , 1820, 511-20	4	38
60	Non-protein-bound iron and 4-hydroxynonenal protein adducts in classic autism. <i>Brain and Development</i> , 2013 , 35, 146-54	2.2	36
59	Redox imbalance and morphological changes in skin fibroblasts in typical Rett syndrome. <i>Oxidative Medicine and Cellular Longevity</i> , 2014 , 2014, 195935	6.7	36
58	OxInflammation in Rett syndrome. International Journal of Biochemistry and Cell Biology, 2016, 81, 246-2	25.36	36
57	Impaired enzymatic defensive activity, mitochondrial dysfunction and proteasome activation are involved in RTT cell oxidative damage. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2015 , 1852, 2066-74	6.9	34
56	Erythrocyte shape abnormalities, membrane oxidative damage, and factin alterations: an unrecognized triad in classical autism. <i>Mediators of Inflammation</i> , 2013 , 2013, 432616	4.3	31
55	Isoprostanes and 4-hydroxy-2-nonenal: markers or mediators of disease? Focus on Rett syndrome as a model of autism spectrum disorder. <i>Oxidative Medicine and Cellular Longevity</i> , 2013 , 2013, 343824	6.7	31
54	Subclinical myocardial dysfunction in Rett syndrome. <i>European Heart Journal Cardiovascular Imaging</i> , 2012 , 13, 339-45	4.1	30
53	F(2)-Dihomo-isoprostanes and brain white matter damage in stage 1 Rett syndrome. <i>Biochimie</i> , 2013 , 95, 86-90	4.6	29
52	High frequency of COH1 intragenic deletions and duplications detected by MLPA in patients with Cohen syndrome. <i>European Journal of Human Genetics</i> , 2010 , 18, 1133-40	5.3	28
51	Coffin-Siris and Nicolaides-Baraitser syndromes are a common well recognizable cause of intellectual disability. <i>Brain and Development</i> , 2015 , 37, 527-36	2.2	26
50	Expression and oxidative modifications of plasma proteins in autism spectrum disorders: Interplay between inflammatory response and lipid peroxidation. <i>Proteomics - Clinical Applications</i> , 2016 , 10, 1103	3 ³ 1 ¹ 112	25
49	Unrecognized lung disease in classic Rett syndrome: a physiologic and high-resolution CT imaging study. <i>Chest</i> , 2010 , 138, 386-92	5.3	24
48	4-hydroxynonenal protein adducts: Key mediator in Rett syndrome oxinflammation. <i>Free Radical Biology and Medicine</i> , 2017 , 111, 270-280	7.8	23
47	Altered erythrocyte membrane fatty acid profile in typical Rett syndrome: effects of omega-3 polyunsaturated fatty acid supplementation. <i>Prostaglandins Leukotrienes and Essential Fatty Acids</i> , 2014 , 91, 183-93	2.8	23
46	Syndromic mental retardation with thrombocytopenia due to 21q22.11q22.12 deletion: Report of three patients. <i>American Journal of Medical Genetics, Part A</i> , 2010 , 152A, 1711-7	2.5	23
45	Retention of Mitochondria in Mature Human Red Blood Cells as the Result of Autophagy Impairment in Rett Syndrome. <i>Scientific Reports</i> , 2017 , 7, 12297	4.9	21

44	Relevance of 4-F-neuroprostane and 10-F-neuroprostane to neurological diseases. <i>Free Radical Biology and Medicine</i> , 2018 , 115, 278-287	7.8	21
43	Periventricular heterotopia with white matter abnormalities associated with 6p25 deletion. <i>American Journal of Medical Genetics, Part A</i> , 2012 , 158A, 1793-7	2.5	20
42	Inflammatory lung disease in Rett syndrome. <i>Mediators of Inflammation</i> , 2014 , 2014, 560120	4.3	18
41	Proteomic analysis of 4-hydroxynonenal and nitrotyrosine modified proteins in RTT fibroblasts. <i>International Journal of Biochemistry and Cell Biology</i> , 2016 , 81, 236-245	5.6	17
40	Lactonase Activity and Lipoprotein-Phospholipase A as Possible Novel Serum Biomarkers for the Differential Diagnosis of Autism Spectrum Disorders and Rett Syndrome: Results from a Pilot Study. <i>Oxidative Medicine and Cellular Longevity</i> , 2017 , 2017, 5694058	6.7	16
39	Effects of B PUFAs supplementation on myocardial function and oxidative stress markers in typical Rett syndrome. <i>Mediators of Inflammation</i> , 2014 , 2014, 983178	4.3	16
38	Alteration of serum lipid profile, SRB1 loss, and impaired Nrf2 activation in CDKL5 disorder. <i>Free Radical Biology and Medicine</i> , 2015 , 86, 156-65	7.8	15
37	Alterations of mitochondrial bioenergetics, dynamics, and morphology support the theory of oxidative damage involvement in autism spectrum disorder. <i>FASEB Journal</i> , 2020 , 34, 6521-6538	0.9	15
36	Rett syndrome: An autoimmune disease?. <i>Autoimmunity Reviews</i> , 2016 , 15, 411-6	13.6	15
35	Creatine transporter defect diagnosed by proton NMR spectroscopy in males with intellectual disability. <i>American Journal of Medical Genetics, Part A</i> , 2011 , 155A, 2446-52	2.5	15
34	MECP2 Duplication Syndrome: Evidence of Enhanced Oxidative Stress. A Comparison with Rett Syndrome. <i>PLoS ONE</i> , 2016 , 11, e0150101	3.7	15
33	Analysis of the Phenotypes in the Rett Networked Database. <i>International Journal of Genomics</i> , 2019 , 2019, 6956934	2.5	14
32	Compromised immune/inflammatory responses in Rett syndrome. <i>Free Radical Biology and Medicine</i> , 2020 , 152, 100-106	7.8	13
31	Altered inflammasome machinery as a key player in the perpetuation of Rett syndrome oxinflammation. <i>Redox Biology</i> , 2020 , 28, 101334	11.3	13
30	Persistent Unresolved Inflammation in the -308 Female Mutated Mouse Model of Rett Syndrome. <i>Mediators of Inflammation</i> , 2017 , 2017, 9467819	4.3	12
29	Beta-actin deficiency with oxidative posttranslational modifications in Rett syndrome erythrocytes: insights into an altered cytoskeletal organization. <i>PLoS ONE</i> , 2014 , 9, e93181	3.7	12
28	Red blood cells in Rett syndrome: oxidative stress, morphological changes and altered membrane organization. <i>Biological Chemistry</i> , 2015 , 396, 1233-40	4.5	11
27	Proteomic analysis of the Rett syndrome experimental model mecp2 mutant zebrafish. <i>Journal of Proteomics</i> , 2017 , 154, 128-133	3.9	10

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26	Immune dysfunction in Rett syndrome patients revealed by high levels of serum anti-N(Glc) IgM antibody fraction. <i>Journal of Immunology Research</i> , 2014 , 2014, 260973	4.5	10
25	Effects of EB polyunsaturated fatty acids on plasma proteome in Rett syndrome. <i>Mediators of Inflammation</i> , 2013 , 2013, 723269	4.3	10
24	Antibody recognition in multiple sclerosis and Rett syndrome using a collection of linear and cyclic N-glucosylated antigenic probes. <i>Biopolymers</i> , 2015 , 104, 560-76	2.2	9
23	A plasma proteomic approach in Rett syndrome: classical versus preserved speech variant. <i>Mediators of Inflammation</i> , 2013 , 2013, 438653	4.3	9
22	13-HODE, 9-HODE and ALOX15 as potential players in Rett syndrome OxInflammation. <i>Free Radical Biology and Medicine</i> , 2019 , 134, 598-603	7.8	8
21	Defective proteasome biogenesis into skin fibroblasts isolated from Rett syndrome subjects with MeCP2 non-sense mutations. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2020 , 1866, 165	793	8
20	Inflammatory protein response in CDKL5-Rett syndrome: evidence of a subclinical smouldering inflammation. <i>Inflammation Research</i> , 2017 , 66, 269-280	7.2	7
19	Isoprostanoids in Clinical and Experimental Neurological Disease Models. <i>Antioxidants</i> , 2018 , 7,	7.1	7
18	Proteomic profiling reveals mitochondrial alterations in Rett syndrome. <i>Free Radical Biology and Medicine</i> , 2020 , 155, 37-48	7.8	7
17	Oxygen exchange and energy metabolism in erythrocytes of Rett syndrome and their relationships with respiratory alterations. <i>Molecular and Cellular Biochemistry</i> , 2017 , 426, 205-213	4.2	6
16	Intestinal Candida parapsilosis isolates from Rett syndrome subjects bear potential virulent traits and capacity to persist within the host. <i>BMC Gastroenterology</i> , 2018 , 18, 57	3	6
15	Antiglycative Activity and RAGE Expression in Rett Syndrome. Cells, 2019, 8,	7.9	6
14	Increased isoprostanoid levels in brain from murine model of Krabbe disease - Relevance of isoprostanes, dihomo-isoprostanes and neuroprostanes to disease severity. <i>Free Radical Biology and Medicine</i> , 2019 , 139, 46-54	7.8	5
13	Abnormal N-glycosylation pattern for brain nucleotide pyrophosphatase-5 (NPP-5) in Mecp2-mutant murine models of Rett syndrome. <i>Neuroscience Research</i> , 2016 , 105, 28-34	2.9	5
12	Oxidative stress: a hallmark of Rett syndrome. <i>Future Neurology</i> , 2015 , 10, 179-182	1.5	4
11	Impaired mitochondrial quality control in Rett Syndrome. <i>Archives of Biochemistry and Biophysics</i> , 2021 , 700, 108790	4.1	4
10	Brain protein changes in Mecp2 mouse mutant models: Effects on disease progression of Mecp2 brain specific gene reactivation. <i>Journal of Proteomics</i> , 2020 , 210, 103537	3.9	4
9	Fatty Acids and Autism Spectrum Disorders: The Rett Syndrome Conundrum. <i>Food and Nutrition Sciences (Print)</i> , 2013 , 04, 71-75	0.4	3

8	Bone status in relation to ambulatory performance in girls with Rett syndrome: a 10-year longitudinal study. <i>Pediatric Research</i> , 2019 , 85, 639-643	3.2	3
7	The complexity of Rett syndrome models: Primary fibroblasts as a disease-in-a-dish reliable approach. <i>Drug Discovery Today: Disease Models</i> , 2020 , 31, 11-19	1.3	2
6	Altered Bone Status in Rett Syndrome. <i>Life</i> , 2021 , 11,	3	1
5	4HNE Protein Adducts in Autistic Spectrum Disorders: Rett Syndrome and Autism 2014 , 2667-2687		1
4	Breathing Abnormalities During Sleep and Wakefulness in Rett Syndrome: Clinical Relevance and Paradoxical Relationship With Circulating Pro-oxidant Markers <i>Frontiers in Neurology</i> , 2022 , 13, 83323	39 ^{4.1}	1
3	A proteomics approach to further highlight the altered inflammatory condition in Rett syndrome. <i>Archives of Biochemistry and Biophysics</i> , 2020 , 696, 108660	4.1	O

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