Sadia Saeed

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/8686512/publications.pdf

Version: 2024-02-01

13 papers	1,279 citations	759233 12 h-index	1125743 13 g-index
14 all docs	14 docs citations	14 times ranked	2520 citing authors

#	Article	IF	Citations
1	Rare Variant Analysis of Obesity-Associated Genes in Young Adults With Severe Obesity From a Consanguineous Population of Pakistan. Diabetes, 2022, 71, 694-705.	0.6	7
2	Genetic Causes of Severe Childhood Obesity: A Remarkably High Prevalence in an Inbred Population of Pakistan. Diabetes, 2020, 69, 1424-1438.	0.6	16
3	Genetics of Obesity in Consanguineous Populations: Toward Precision Medicine and the Discovery of Novel Obesity Genes. Obesity, 2018, 26, 474-484.	3.0	35
4	Loss-of-function mutations in ADCY3 cause monogenic severe obesity. Nature Genetics, 2018, 50, 175-179.	21.4	122
5	Genetic variants in LEP, LEPR, and <scp>MC4R</scp> explain 30% of severe obesity in children from a consanguineous population. Obesity, 2015, 23, 1687-1695.	3.0	82
6	Functional Characterization of Obesity-Associated Variants Involving the Î± and Î² Isoforms of Human SH2B1. Endocrinology, 2014, 155, 3219-3226.	2.8	39
7	Changes in levels of peripheral hormones controlling appetite are inconsistent with hyperphagia in leptin-deficient subjects. Endocrine, 2014, 45, 401-408.	2.3	19
8	Highly Sensitive Diagnosis of 43 Monogenic Forms of Diabetes or Obesity Through One-Step PCR-Based Enrichment in Combination With Next-Generation Sequencing. Diabetes Care, 2014, 37, 460-467.	8.6	69
9	Novel LEPR mutations in obese Pakistani children identified by PCRâ€based enrichment and next generation sequencing. Obesity, 2014, 22, 1112-1117.	3.0	51
10	Rare variants in single-minded 1 (SIM1) are associated with severe obesity. Journal of Clinical Investigation, 2013, 123, 3042-3050.	8.2	135
11	High prevalence of leptin and melanocortin-4 receptor gene mutations in children with severe obesity from Pakistani consanguineous families. Molecular Genetics and Metabolism, 2012, 106, 121-126.	1.1	70
12	Human SH2B1 mutations are associated with maladaptive behaviors and obesity. Journal of Clinical Investigation, 2012, 122, 4732-4736.	8.2	147
13	Large, rare chromosomal deletions associated with severe early-onset obesity. Nature, 2010, 463, 666-670.	27.8	487