

Julia Gj Dancourt

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5380810/publications.pdf>

Version: 2024-02-01

12
papers

1,206
citations

1040056

9
h-index

1125743

13
g-index

21
all docs

21
docs citations

21
times ranked

2425
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|--|------|-----------|
| 1 | Wide clinical spectrum in ALC8-CDG: clues from molecular findings suggest an explanation for a milder phenotype in the first-described patient. <i>Pediatric Research</i> , 2019, 85, 384-389. | 2.3 | 8 |
| 2 | Small cargoes pass through synthetically glued Golgi stacks. <i>FEBS Letters</i> , 2016, 590, 1675-1686. | 2.8 | 9 |
| 3 | Lipidation of the autophagy proteins LC3 and GABARAP is a membrane-curvature dependent process. <i>Autophagy</i> , 2014, 10, 1470-1471. | 9.1 | 36 |
| 4 | Lipidation of the LC3/GABARAP family of autophagy proteins relies on a membrane-curvature-sensing domain in Atg3. <i>Nature Cell Biology</i> , 2014, 16, 415-424. | 10.3 | 221 |
| 5 | The Lipidation Machinery Involved in Autophagosome Growth is Only Functional on Highly Curved Membranes. <i>Biophysical Journal</i> , 2013, 104, 97a. | 0.5 | 0 |
| 6 | The <i>Legionella</i> Effector RavZ Inhibits Host Autophagy Through Irreversible Atg8 Deconjugation. <i>Science</i> , 2012, 338, 1072-1076. | 12.6 | 401 |
| 7 | Protein Sorting Receptors in the Early Secretory Pathway. <i>Annual Review of Biochemistry</i> , 2010, 79, 777-802. | 11.1 | 271 |
| 8 | Erv26-Dependent Export of Alkaline Phosphatase from the ER Requires Luminal Domain Recognition. <i>Traffic</i> , 2009, 10, 1006-1018. | 2.7 | 17 |
| 9 | A New Intronic Mutation in the DPM1 Gene Is Associated With a Milder Form of CDG Ie in Two French Siblings. <i>Pediatric Research</i> , 2006, 59, 835-839. | 2.3 | 31 |
| 10 | Two Proteins Homologous to the N- and C-terminal Domains of the Bacterial Glycosyltransferase Murg Are Required for the Second Step of Dolichyl-linked Oligosaccharide Synthesis in <i>Saccharomyces cerevisiae</i> . <i>Journal of Biological Chemistry</i> , 2005, 280, 9236-9242. | 3.4 | 41 |
| 11 | A Deficiency in Dolichyl-P-glucose:Glc1Man9GlcNAc2-PP-dolichyl β 3-Glucosyltransferase Defines a New Subtype of Congenital Disorders of Glycosylation. <i>Journal of Biological Chemistry</i> , 2003, 278, 9962-9971. | 3.4 | 78 |
| 12 | Congenital Disorders of Glycosylation Type Ig Is Defined by a Deficiency in Dolichyl-P-mannose:Man7GlcNAc2-PP-dolichyl Mannosyltransferase. <i>Journal of Biological Chemistry</i> , 2002, 277, 25815-25822. | 3.4 | 87 |