

Sergio Guerrero-Castillo

List of Publications by Year in descending order

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Version: 2024-02-01

38
papers

1,882
citations

304743

22
h-index

315739

38
g-index

42
all docs

42
docs citations

42
times ranked

2947
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|--|------|-----------|
| 1 | ZBTB11 dysfunction: spectrum of brain abnormalities, biochemical signature and cellular consequences. <i>Brain</i> , 2022, 145, 2602-2616. | 7.6 | 5 |
| 2 | The plastid proteome of the nonphotosynthetic chlorophycean alga <i>Polytomella parva</i> . <i>Microbiological Research</i> , 2021, 243, 126649. | 5.3 | 5 |
| 3 | Neutropenia and intellectual disability are hallmarks of biallelic and de novo CLPB deficiency. <i>Genetics in Medicine</i> , 2021, 23, 1705-1714. | 2.4 | 22 |
| 4 | CEDAR, an online resource for the reporting and exploration of complexome profiling data. <i>Biochimica Et Biophysica Acta - Bioenergetics</i> , 2021, 1862, 148411. | 1.0 | 27 |
| 5 | Multiplexed complexome profiling using tandem mass tags. <i>Biochimica Et Biophysica Acta - Bioenergetics</i> , 2021, 1862, 148448. | 1.0 | 6 |
| 6 | Ablation of mitochondrial DNA results in widespread remodeling of the mitochondrial complexome. <i>EMBO Journal</i> , 2021, 40, e108648. | 7.8 | 18 |
| 7 | Complexome Profiling—Exploring Mitochondrial Protein Complexes in Health and Disease. <i>Frontiers in Cell and Developmental Biology</i> , 2021, 9, 796128. | 3.7 | 20 |
| 8 | Novel defect in phosphatidylinositol 4-kinase type 2 α (<i>PI4K2A</i>) at the membrane-enzyme interface is associated with metabolic cutis laxa. <i>Journal of Inherited Metabolic Disease</i> , 2020, 43, 1382-1391. | 3.6 | 7 |
| 9 | TMEM70 functions in the assembly of complexes I and V. <i>Biochimica Et Biophysica Acta - Bioenergetics</i> , 2020, 1861, 148202. | 1.0 | 31 |
| 10 | A salvage pathway maintains highly functional respiratory complex I. <i>Nature Communications</i> , 2020, 11, 1643. | 12.8 | 80 |
| 11 | Complexome analysis of the nitrite-dependent methanotroph <i>Methylomirabilis lanthanidiphila</i> . <i>Biochimica Et Biophysica Acta - Bioenergetics</i> , 2019, 1860, 734-744. | 1.0 | 18 |
| 12 | Complexome Profiling Alignment (COPAL) reveals remodeling of mitochondrial protein complexes in Barth syndrome. <i>Bioinformatics</i> , 2019, 35, 3083-3091. | 4.1 | 37 |
| 13 | Deactivation of mitochondrial complex I after hypoxia—“ischemia in the immature brain. <i>Journal of Cerebral Blood Flow and Metabolism</i> , 2019, 39, 1790-1802. | 4.3 | 24 |
| 14 | Bi-allelic Mutations in the Mitochondrial Ribosomal Protein MRPS2 Cause Sensorineural Hearing Loss, Hypoglycemia, and Multiple OXPHOS Complex Deficiencies. <i>American Journal of Human Genetics</i> , 2018, 102, 685-695. | 6.2 | 61 |
| 15 | Locking loop movement in the ubiquinone pocket of complex I disengages the proton pumps. <i>Nature Communications</i> , 2018, 9, 4500. | 12.8 | 80 |
| 16 | Barth syndrome cells display widespread remodeling of mitochondrial complexes without affecting metabolic flux distribution. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2018, 1864, 3650-3658. | 3.8 | 53 |
| 17 | Mutations in ATP6V1E1 or ATP6V1A Cause Autosomal-Recessive Cutis Laxa. <i>American Journal of Human Genetics</i> , 2017, 100, 216-227. | 6.2 | 82 |
| 18 | MR-1S Interacts with PET100 and PET117 in Module-Based Assembly of Human Cytochrome c Oxidase. <i>Cell Reports</i> , 2017, 18, 1727-1738. | 6.4 | 86 |

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|----|---|------|-----------|
| 19 | A homozygous missense mutation in ERAL1, encoding a mitochondrial rRNA chaperone, causes Perrault syndrome. <i>Human Molecular Genetics</i> , 2017, 26, 2541-2550. | 2.9 | 61 |
| 20 | Identification and evolutionary analysis of tissue-specific isoforms of mitochondrial complex I subunit NDUFV3. <i>Biochimica Et Biophysica Acta - Bioenergetics</i> , 2017, 1858, 208-217. | 1.0 | 33 |
| 21 | The Assembly Pathway of Mitochondrial Respiratory Chain Complex I. <i>Cell Metabolism</i> , 2017, 25, 128-139. | 16.2 | 325 |
| 22 | Mutations in Complex I Assembly Factor TMEM126B Result in Muscle Weakness and Isolated Complex I Deficiency. <i>American Journal of Human Genetics</i> , 2016, 99, 208-216. | 6.2 | 51 |
| 23 | Mitochondrial Dysfunction during the Early Stages of Excitotoxic Spinal Motor Neuron Degeneration in Vivo. <i>ACS Chemical Neuroscience</i> , 2016, 7, 886-896. | 3.5 | 18 |
| 24 | Unraveling the complexity of mitochondrial complex I assembly: A dynamic process. <i>Biochimica Et Biophysica Acta - Bioenergetics</i> , 2016, 1857, 980-990. | 1.0 | 110 |
| 25 | The m-AAA Protease Associated with Neurodegeneration Limits MCU Activity in Mitochondria. <i>Molecular Cell</i> , 2016, 64, 148-162. | 9.7 | 153 |
| 26 | Evolution and structural organization of the mitochondrial contact site (MICOS) complex and the mitochondrial intermembrane space bridging (MIB) complex. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 2016, 1863, 91-101. | 4.1 | 150 |
| 27 | <i>Staphylococcus epidermidis</i> : metabolic adaptation and biofilm formation in response to different oxygen concentrations. <i>Pathogens and Disease</i> , 2016, 74, ftv111. | 2.0 | 35 |
| 28 | Accessory NUMM (NDUFS6) subunit harbors a Zn-binding site and is essential for biogenesis of mitochondrial complex I. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015, 112, 5685-5690. | 7.1 | 64 |
| 29 | The <i>Saccharomyces cerevisiae</i> mitochondrial unselective channel behaves as a physiological uncoupling system regulated by Ca ²⁺ , Mg ²⁺ , phosphate and ATP. <i>Journal of Bioenergetics and Biomembranes</i> , 2015, 47, 477-491. | 2.3 | 10 |
| 30 | Oxidative phosphorylation in <i>Debaryomyces hansenii</i> : Physiological uncoupling at different growth phases. <i>Biochimie</i> , 2014, 102, 124-136. | 2.6 | 5 |
| 31 | The branched mitochondrial respiratory chain from <i>Debaryomyces hansenii</i> : Components and supramolecular organization. <i>Biochimica Et Biophysica Acta - Bioenergetics</i> , 2014, 1837, 73-84. | 1.0 | 19 |
| 32 | A critical tyrosine residue determines the uncoupling protein-like activity of the yeast mitochondrial oxaloacetate carrier. <i>Biochemical Journal</i> , 2012, 443, 317-325. | 3.7 | 4 |
| 33 | Effect of glycolysis inhibition on mitochondrial function in rat brain. <i>Journal of Biochemical and Molecular Toxicology</i> , 2012, 26, 206-211. | 3.0 | 12 |
| 34 | During the stationary growth phase, <i>Yarrowia lipolytica</i> prevents the overproduction of reactive oxygen species by activating an uncoupled mitochondrial respiratory pathway. <i>Biochimica Et Biophysica Acta - Bioenergetics</i> , 2012, 1817, 353-362. | 1.0 | 25 |
| 35 | Physiological uncoupling of mitochondrial oxidative phosphorylation. Studies in different yeast species. <i>Journal of Bioenergetics and Biomembranes</i> , 2011, 43, 323-331. | 2.3 | 38 |
| 36 | Mitochondrial Unselective Channels throughout the eukaryotic domain. <i>Mitochondrion</i> , 2011, 11, 382-390. | 3.4 | 35 |

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|----|---|-----|-----------|
| 37 | Mitochondria from the salt-tolerant yeast <i>Debaryomyces hansenii</i> (halophilic organelles?). <i>Journal of Bioenergetics and Biomembranes</i> , 2010, 42, 11-19. | 2.3 | 25 |
| 38 | In <i>Yarrowia lipolytica</i> mitochondria, the alternative NADH dehydrogenase interacts specifically with the cytochrome complexes of the classic respiratory pathway. <i>Biochimica Et Biophysica Acta - Bioenergetics</i> , 2009, 1787, 75-85. | 1.0 | 42 |