

# Sarah F Pearce

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/8593973/publications.pdf>

Version: 2024-02-01

15  
papers

856  
citations

687220

13  
h-index

996849

15  
g-index

18  
all docs

18  
docs citations

18  
times ranked

1512  
citing authors

| #  | ARTICLE   | IF  | CITATIONS |
|----|---|-----|-----------|
| 1  | Human GTPBP5 is involved in the late stage of mitoribosome large subunit assembly. <i>Nucleic Acids Research</i> , 2021, 49, 354-370.   | 6.5 | 21        |
| 2  | Mitoribosome Profiling from Human Cell Culture: A High Resolution View of Mitochondrial Translation. <i>Methods in Molecular Biology</i> , 2021, 2192, 183-196.   | 0.4 | 5         |
| 3  | Differential processing and localization of human Nocturnin controls metabolism of mRNA and nicotinamide adenine dinucleotide cofactors. <i>Journal of Biological Chemistry</i> , 2020, 295, 15112-15133.     | 1.6 | 6         |
| 4  | Metabolic shift underlies recovery in reversible infantile respiratory chain deficiency. <i>EMBO Journal</i> , 2020, 39, e105364.   | 3.5 | 26        |
| 5  | C6orf203 is an RNA-binding protein involved in mitochondrial protein synthesis. <i>Nucleic Acids Research</i> , 2019, 47, 9386-9399.  | 6.5 | 26        |
| 6  | MitoRibo-Tag Mice Provide a Tool for In Vivo Studies of Mitoribosome Composition. <i>Cell Reports</i> , 2019, 29, 1728-1738.e9.   | 2.9 | 24        |
| 7  | Maturation of selected human mitochondrial tRNAs requires deadenylation. <i>ELife</i> , 2017, 6, .  | 2.8 | 72        |
| 8  | TRNT1 deficiency: clinical, biochemical and molecular genetic features. <i>Orphanet Journal of Rare Diseases</i> , 2016, 11, 90.  | 1.2 | 64        |
| 9  | Deficient methylation and formylation of mt-tRNAMet wobble cytosine in a patient carrying mutations in NSUN3. <i>Nature Communications</i> , 2016, 7, 12039.  | 5.8 | 178       |
| 10 | Mitochondrial transcript maturation and its disorders. <i>Journal of Inherited Metabolic Disease</i> , 2015, 38, 655-680.   | 1.7 | 69        |
| 11 | Mutations in GTPBP3 Cause a Mitochondrial Translation Defect Associated with Hypertrophic Cardiomyopathy, Lactic Acidosis, and Encephalopathy. <i>American Journal of Human Genetics</i> , 2014, 95, 708-720. | 2.6 | 123       |
| 12 | MPV17L2 is required for ribosome assembly in mitochondria. <i>Nucleic Acids Research</i> , 2014, 42, 8500-8515.   | 6.5 | 56        |
| 13 | MRM2 and MRM3 are involved in biogenesis of the large subunit of the mitochondrial ribosome. <i>Molecular Biology of the Cell</i> , 2014, 25, 2542-2555.  | 0.9 | 99        |
| 14 | Polyadenylation in Bacteria and Organelles. <i>Methods in Molecular Biology</i> , 2014, 1125, 211-227.  | 0.4 | 23        |
| 15 | Mitochondrial diseases: Translation matters. <i>Molecular and Cellular Neurosciences</i> , 2013, 55, 1-12.  | 1.0 | 62        |