

Anoop K Sendamarai

List of Publications by Year in descending order

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Version: 2024-02-01

18
papers

1,066
citations

623734

14
h-index

888059

17
g-index

20
all docs

20
docs citations

20
times ranked

1970
citing authors

| # | ARTICLE | IF | CITATIONS |
|----|--|------|-----------|
| 1 | Regulatory variants in TCF7L2 are associated with thoracic aortic aneurysm. American Journal of Human Genetics, 2021, 108, 1578-1589. | 6.2 | 17 |
| 2 | Mutations in the iron-sulfur cluster biogenesis protein HSCB cause congenital sideroblastic anemia. Journal of Clinical Investigation, 2020, 130, 5245-5256. | 8.2 | 13 |
| 3 | Lack of Gdf11 does not improve anemia or prevent the activity of RAP-536 in a mouse model of β^2 -thalassemia. Blood, 2019, 134, 568-572. | 1.4 | 56 |
| 4 | Downregulation of ribosome biogenesis during early forebrain development. ELife, 2018, 7, . | 6.0 | 72 |
| 5 | Bone marrow failure unresponsive to bone marrow transplant is caused by mutations in thrombopoietin. Blood, 2017, 130, 875-880. | 1.4 | 42 |
| 6 | UBE2O remodels the proteome during terminal erythroid differentiation. Science, 2017, 357, . | 12.6 | 121 |
| 7 | <i>Hscb</i> , a Mitochondrial Iron-Sulfur Cluster Assembly Co-Chaperone, Is a Novel Candidate Gene for Congenital Sideroblastic Anemia. Blood, 2017, 130, 79-79. | 1.4 | 4 |
| 8 | Pseudouridine synthase 1 deficient mice, a model for Mitochondrial Myopathy with Sideroblastic Anemia, exhibit muscle morphology and physiology alterations. Scientific Reports, 2016, 6, 26202. | 3.3 | 26 |
| 9 | A recurring mutation in the respiratory complex 1 protein NDUF11 is responsible for a novel form of X-linked sideroblastic anemia. Blood, 2016, 128, 1913-1917. | 1.4 | 33 |
| 10 | Congenital sideroblastic anemia due to mutations in the mitochondrial HSP70 homologue HSPA9. Blood, 2015, 126, 2734-2738. | 1.4 | 78 |
| 11 | Indolent T-lymphoblastic Proliferation With Disseminated Multinodal Involvement and Partial CD33 Expression. American Journal of Surgical Pathology, 2014, 38, 1298-1304. | 3.7 | 27 |
| 12 | X-linked sideroblastic anemia due to ALAS2 intron 1 enhancer element GATA binding site mutations. American Journal of Hematology, 2014, 89, 315-319. | 4.1 | 39 |
| 13 | Mutations in TRNT1 cause congenital sideroblastic anemia with immunodeficiency, fevers, and developmental delay (SIFD). Blood, 2014, 124, 2867-2871. | 1.4 | 162 |
| 14 | An RNAi therapeutic targeting Tmprss6 decreases iron overload in Hfe ^{-/-} mice and ameliorates anemia and iron overload in murine β^2 -thalassemia intermedia. Blood, 2013, 121, 1200-1208. | 1.4 | 180 |
| 15 | The Crystal Structure of Six-transmembrane Epithelial Antigen of the Prostate 4 (Steap4), a Ferri/Cuprioreductase, Suggests a Novel Interdomain Flavin-binding Site. Journal of Biological Chemistry, 2013, 288, 20668-20682. | 3.4 | 33 |
| 16 | Atomic structure of the 75 MDa extremophile <i>Sulfolobus</i> turreted icosahedral virus determined by CryoEM and X-ray crystallography. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, 5504-5509. | 7.1 | 77 |
| 17 | RNAi-Mediated Inhibition of Tmprss6 Ameliorates Anemia and Secondary Iron Overload in a Mouse Model of β^2 -Thalassemia Intermedia and Decreases Iron Overload in Hfe ^{-/-} Mice. Blood, 2012, 120, 1018-1018. | 1.4 | 0 |
| 18 | Structure of the membrane proximal oxidoreductase domain of human Steap3, the dominant ferrireductase of the erythroid transferrin cycle. Proceedings of the National Academy of Sciences of the United States of America, 2008, 105, 7410-7415. | 7.1 | 83 |