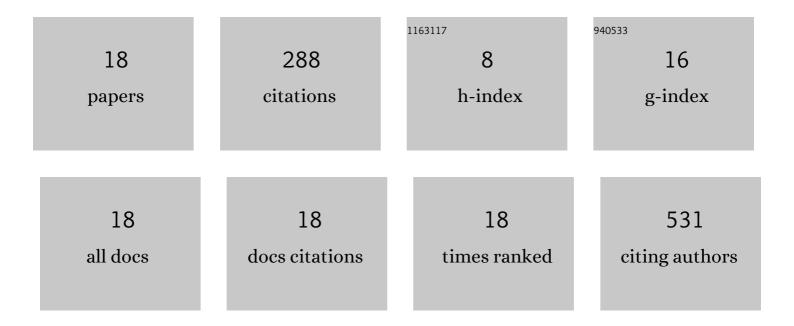
Essam Al Ageeli

List of Publications by Year in descending order

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| # | Article | IF | CITATIONS |
|----|---|-----|-----------|
| 1 | Duplication of the 15q11-q13 region: Clinical and genetic study of 30 new cases. European Journal of Medical Genetics, 2014, 57, 5-14. | 1.3 | 68 |
| 2 | Oncogenic long noncoding RNA MALAT1 and HCV-related hepatocellular carcinoma. Biomedicine and Pharmacotherapy, 2018, 102, 653-669. | 5.6 | 40 |
| 3 | A new 17p13.3 microduplication including the PAFAH1B1 and YWHAE genes resulting from an unbalanced X;17 translocation. European Journal of Medical Genetics, 2011, 54, 287-291. | 1.3 | 27 |
| 4 | Deciphering the role of circulating lncRNAs: RNCR2, NEAT2, CDKN2B-AS1, and PVT1 and the possible prediction of anti-VEGF treatment outcomes in diabetic retinopathy patients. Graefe's Archive for Clinical and Experimental Ophthalmology, 2019, 257, 1897-1913. | 1.9 | 26 |
| 5 | Long non-coding RNA MALAT1 and microRNA-499a expression profiles in diabetic ESRD patients undergoing dialysis: a preliminary cross-sectional analysis. Archives of Physiology and Biochemistry, 2020, 126, 172-182. | 2.1 | 24 |
| 6 | Structure and functional impact of seed region variant in MIR-499 gene family in bronchial asthma. Respiratory Research, 2017, 18, 169. | 3.6 | 17 |
| 7 | Unleash multifunctional role of long noncoding RNAs biomarker panel in breast cancer: a predictor classification model. Epigenomics, 2020, 12, 1215-1237. | 2.1 | 15 |
| 8 | MYD88, NFKB1, and IL6 transcripts overexpression are associated with poor outcomes and short survival in neonatal sepsis. Scientific Reports, 2021, 11, 13374. | 3.3 | 13 |
| 9 | Retinal involvement in two unrelated patients with Myhre syndrome. European Journal of Medical Genetics, 2012, 55, 541-547. | 1.3 | 11 |
| 10 | Alterations of mitochondria and related metabolic pathways in leukemia: A narrative review. Saudi Journal of Medicine and Medical Sciences, 2020, 8, 3. | 0.8 | 10 |
| 11 | Association of Angio-LncRNAs MIAT rs1061540/MALAT1 rs3200401 Molecular Variants with Gensini Score in Coronary Artery Disease Patients Undergoing Angiography. Biomolecules, 2022, 12, 137. | 4.0 | 9 |
| 12 | Long Non-Coding RNAs Gene Variants as Molecular Markers for Diabetic Retinopathy Risk and Response to Anti-VEGF Therapy. Pharmacogenomics and Personalized Medicine, 2021, Volume 14, 997-1014. | 0.7 | 8 |
| 13 | Noncoding RNAs orchestrate cell growth, death and drug resistance in renal cell carcinoma. Epigenomics, 2020, 12, 199-219. | 2.1 | 5 |
| 14 | Longevity-Related Gene Transcriptomic Signature in Glioblastoma Multiforme. Oxidative Medicine and Cellular Longevity, 2018, 2018, 1-12. | 4.0 | 4 |
| 15 | Group-specific component exon 11 haplotypes (D432E and T436K) and risk of albuminuria in type 2 diabetes mellitus patients. Archives of Physiology and Biochemistry, 2022, 128, 111-120. | 2.1 | 3 |
| 16 | MicroRNA-17-92a-1 Host Gene (MIR17HG) Expression Signature and rs4284505 Variant Association with Alopecia Areata: A Case–Control Study. Genes, 2022, 13, 505. | 2.4 | 3 |
| 17 | Genotype Triad for HOTAIR rs10783618, LINC-ROR rs1942347, and MALAT1 rs3200401 as Molecular Markers in Systemic Lupus Erythematous. Diagnostics, 2022, 12, 1197. | 2.6 | 3 |
| 18 | MicroRNA‑499a (rs3746444A/G) gene variant and susceptibility to typeÂ2 diabetes‑associated end‑stage renal disease. Experimental and Therapeutic Medicine, 2021, 23, 63. | 1.8 | 2 |