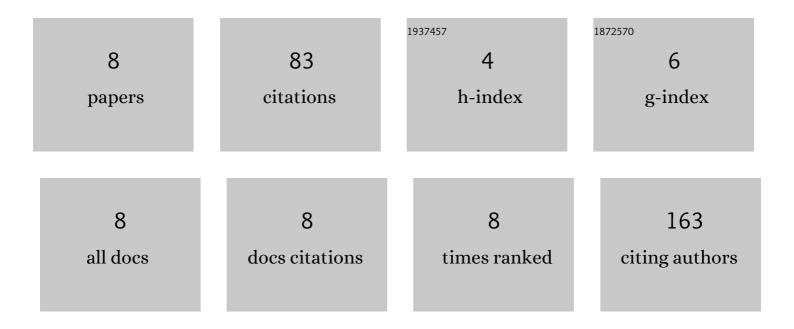
## Landry E Nfonsam

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/5131530/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Adapting the ACMG/AMP variant classification framework: A perspective from the ClinGen Hemoglobinopathy Variant Curation Expert Panel. Human Mutation, 2022, 43, 1089-1096.	1.1	20
2	Interplay between probe design and test performance: overlap between genomic regions of interest, capture regions and high quality reference calls influence performance of WES-based assays. Human Genetics, 2021, 140, 289-297.	1.8	0
3	<i>ALU</i> transposition induces familial hypertrophic cardiomyopathy. Molecular Genetics & Genomic Medicine, 2020, 8, e951.	0.6	1
4	Adopting High-Resolution Allele Frequencies Substantially Expedites Variant Interpretation in Genetic Diagnostic Laboratories. Journal of Molecular Diagnostics, 2019, 21, 602-611.	1.2	0
5	Genetic Diagnostic Testing for Inherited Cardiomyopathies. Journal of Molecular Diagnostics, 2019, 21, 437-448.	1.2	7
6	COMP Gene Coexpresses With EMT Genes and Is Associated With Poor Survival in Colon Cancer Patients. Journal of Surgical Research, 2019, 233, 297-303.	0.8	24
7	Leveraging the power of new molecular technologies in the clinical setting requires unprecedented awareness of limitations and drawbacks: experience of one diagnostic laboratory. Journal of Medical Genetics, 2019, 56, 408-412.	1.5	3
8	<i>SFRP4</i> expression correlates with epithelial mesenchymal transition-linked genes and poor overall survival in colon cancer patients. World Journal of Gastrointestinal Oncology, 2019, 11, 589-598.	0.8	28