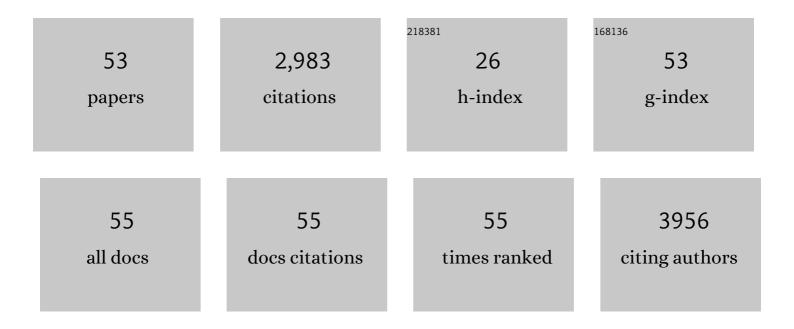
## Lajos KalmÃ;r

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	HAM-ART: An optimised culture-free Hi-C metagenomics pipeline for tracking antimicrobial resistance genes in complex microbial communities. PLoS Genetics, 2022, 18, e1009776.	1.5	14
2	Size-Exclusion Chromatography Separation Reveals That Vesicular and Non-Vesicular Small RNA Profiles Differ in Cell Free Urine. International Journal of Molecular Sciences, 2021, 22, 4881.	1.8	6
3	Abnormal patterns of sleep and EEG power distribution during non-rapid eye movement sleep in the sheep model of Huntington's disease. Neurobiology of Disease, 2021, 155, 105367.	2.1	21
4	Wake-Promoting and EEG Spectral Effects of Modafinil After Acute or Chronic Administration in the R6/2 Mouse Model of Huntington's Disease. Neurotherapeutics, 2020, 17, 1075-1086.	2.1	8
5	Characterisation of canine KCNIP4: A novel gene for cerebellar ataxia identified by whole-genome sequencing two affected Norwegian Buhund dogs. PLoS Genetics, 2020, 16, e1008527.	1.5	10
6	Intrinsically Disordered Linkers Impart Processivity on Enzymes by Spatial Confinement of Binding Domains. International Journal of Molecular Sciences, 2019, 20, 2119.	1.8	15
7	Truncation of GdpP mediates β-lactam resistance in clinical isolates of Staphylococcus aureus. Journal of Antimicrobial Chemotherapy, 2019, 74, 1182-1191.	1.3	40
8	Objective effectiveness of and indications for laserâ€essisted turbinectomy in brachycephalic obstructive airway syndrome. Veterinary Surgery, 2019, 48, 79-87.	0.5	18
9	Financial Stress Interacts With CLOCK Gene to Affect Migraine. Frontiers in Behavioral Neuroscience, 2019, 13, 284.	1.0	4
10	Endotracheal tube placement during computed tomography of brachycephalic dogs alters upper airway dimensional measurements. Veterinary Radiology and Ultrasound, 2018, 59, 289-304.	0.4	8
11	Brachycephalic obstructive airway syndrome. Veterinary Record, 2018, 182, 375-378.	0.2	33
12	Outcomes and prognostic factors of surgical treatments for brachycephalic obstructive airway syndrome in 3 breeds. Veterinary Surgery, 2017, 46, 271-280.	0.5	50
13	DisProt 7.0: a major update of the database of disordered proteins. Nucleic Acids Research, 2017, 45, D219-D227.	6.5	242
14	Conformational risk factors of brachycephalic obstructive airway syndrome (BOAS) in pugs, French bulldogs, and bulldogs. PLoS ONE, 2017, 12, e0181928.	1.1	74
15	Structural determinants governing S100A4â€induced isoformâ€selective disassembly of nonmuscle myosin <scp>II</scp> filaments. FEBS Journal, 2016, 283, 2164-2180.	2.2	13
16	Wholeâ€Body Barometric Plethysmography Characterizes Upper Airway Obstruction in 3 Brachycephalic Breeds of Dogs. Journal of Veterinary Internal Medicine, 2016, 30, 853-865.	0.6	74
17	The role of structural disorder in cell cycle regulation, related clinical proteomics, disease development and drug targeting. Expert Review of Proteomics, 2015, 12, 221-233.	1.3	14
18	Intrinsically disordered proteins: emerging interaction specialists. Current Opinion in Structural Biology, 2015, 35, 49-59.	2.6	177

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19	Chronic escitalopram treatment caused dissociative adaptation in serotonin (5-HT) 2C receptor antagonist-induced effects in REM sleep, wake and theta wave activity. Experimental Brain Research, 2014, 232, 935-946.	0.7	12
20	Contribution of proline to the pre-structuring tendency of transient helical secondary structure elements in intrinsically disordered proteins. Biochimica Et Biophysica Acta - General Subjects, 2014, 1840, 993-1003.	1.1	26
21	Narcolepsy patients have antibodies that stain distinct cell populations in rat brain and influence sleep patterns. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, E3735-44.	3.3	71
22	The other side of the coin: Functional and structural versatility of ADF/cofilins. European Journal of Cell Biology, 2014, 93, 238-251.	1.6	39
23	Discrete Molecular Dynamics Can Predict Helical Prestructured Motifs in Disordered Proteins. PLoS ONE, 2014, 9, e95795.	1.1	19
24	Acute escitalopram treatment inhibits REM sleep rebound and activation of MCH-expressing neurons in the lateral hypothalamus after long term selective REM sleep deprivation. Psychopharmacology, 2013, 228, 439-449.	1.5	12
25	Differential adaptation of REM sleep latency, intermediate stage and theta power effects of escitalopram after chronic treatment. Journal of Neural Transmission, 2013, 120, 169-176.	1.4	18
26	Factors affecting DNA quality in feathers used for non-invasive sampling. Journal of Ornithology, 2013, 154, 587-595.	0.5	19
27	The alphabet of intrinsic disorder. Intrinsically Disordered Proteins, 2013, 1, e24360.	1.9	208
28	Exon-phase symmetry and intrinsic structural disorder promote modular evolution in the human genome. Nucleic Acids Research, 2013, 41, 4409-4422.	6.5	15
29	Structural disorder in proteins brings order to crystal growth in biomineralization. Bone, 2012, 51, 528-534.	1.4	107
30	Long-Range Interactions in Nonsense-Mediated mRNA Decay Are Mediated by Intrinsically Disordered Protein Regions. Journal of Molecular Biology, 2012, 424, 125-131.	2.0	7
31	Characterization of ABL exon 7 deletion by molecular genetic and bioinformatic methods reveals no association with imatinib resistance in chronic myeloid leukemia. Medical Oncology, 2012, 29, 2136-2142.	1.2	8
32	Distinct Hydration Properties of Wild-Type and Familial Point Mutant A53T of α-Synuclein Associated with Parkinson's Disease. Biophysical Journal, 2011, 101, 2260-2266.	0.2	33
33	Verification of alternative splicing variants based on domain integrity, truncation length and intrinsic protein disorder. Nucleic Acids Research, 2011, 39, 1208-1219.	6.5	46
34	Reduction in Structural Disorder and Functional Complexity in the Thermal Adaptation of Prokaryotes. PLoS ONE, 2010, 5, e12069.	1.1	69
35	Dual coding in alternative reading frames correlates with intrinsic protein disorder. Proceedings of the United States of America, 2010, 107, 5429-5434.	3.3	92
36	Power Law Distribution Defines Structural Disorder as a Structural Element Directly Linked with Function. Journal of Molecular Biology, 2010, 403, 346-350.	2.0	29

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37	Genetic structure of the Imperial Eagle (Aquila heliaca) population in Slovakia. Slovak Raptor Journal, 2009, 3, 21-28.	0.4	11
38	TOPDOM: database of domains and motifs with conservative location in transmembrane proteins. Bioinformatics, 2008, 24, 1469-1470.	1.8	28
39	TOPDB: topology data bank of transmembrane proteins. Nucleic Acids Research, 2007, 36, D234-D239.	6.5	73
40	Analysis of the population heterogeneity in Hungary using fifteen forensically informative STR markers. Forensic Science International, 2006, 158, 244-249.	1.3	25
41	Asymmetric PCR increases efficiency of melting peak analysis on the LightCycler. Clinical Biochemistry, 2005, 38, 727-730.	0.8	22
42	An overlooked DNA source for non-invasive genetic analysis in birds. Journal of Avian Biology, 2005, 36, 84-88.	0.6	151
43	HAEdb: A novel interactive, locus-specific mutation database for the C1 inhibitor gene. Human Mutation, 2005, 25, 1-5.	1.1	88
44	Frequencies of two common mutations (c.35delG and c.167delT) of the connexin 26 gene in different populations of Hungary. International Journal of Molecular Medicine, 2004, 14, 1105.	1.8	1
45	Hereditary and acquired angioedema: Problems and progress: Proceedings of the third C1 esterase inhibitor deficiency workshop and beyond. Journal of Allergy and Clinical Immunology, 2004, 114, S51-S131.	1.5	582
46	Mutation History of the Roma/Gypsies. American Journal of Human Genetics, 2004, 75, 596-609.	2.6	148
47	Frequencies of two common mutations (c.35delG and c.167delT) of the connexin 26 gene in different populations of Hungary. International Journal of Molecular Medicine, 2004, 14, 1105-8.	1.8	14
48	Mutation screening of the C1 inhibitor gene among Hungarian patients with hereditary angioedema. Human Mutation, 2003, 22, 498-498.	1.1	63
49	Population genetic analysis in Hungarian populations using the Powerplexâ,,¢ 16 system. International Congress Series, 2003, 1239, 121-122.	0.2	0
50	Association of celiac disease and hereditary angioneurotic edema. American Journal of Gastroenterology, 2002, 97, 2682-2683.	0.2	16
51	The P28T Mutation in the GALK1 Gene Accounts for Galactokinase Deficiency in Roma (Gypsy) Patients across Europe. Pediatric Research, 2002, 51, 602-606.	1.1	40
52	Genotype Screening for Hereditary Hemochromatosis among Voluntary Blood Donors in Hungary. Blood Cells, Molecules, and Diseases, 2001, 27, 334-341.	0.6	17
53	High frequency of the haemochromatosis C282Y mutation in Hungary could argue against a Celtic origin of the mutation Journal of Medical Genetics, 1998, 35, 878-879.	1.5	13