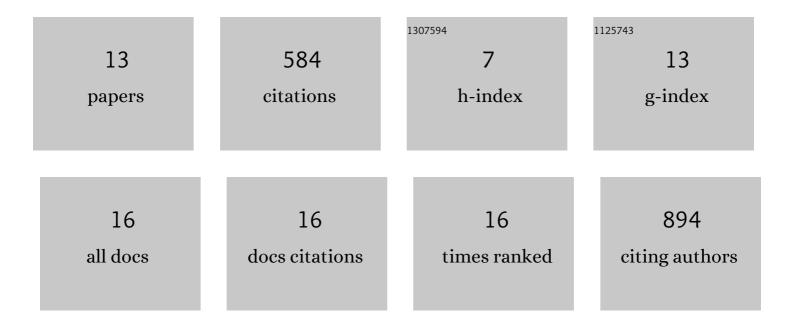
Kaoru Inoue

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	SHAPE reveals transcript-wide interactions, complex structural domains, and protein interactions across the <i>Xist</i> IncRNA in living cells. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, 10322-10327.	7.1	201
2	Functional classification of long non-coding RNAs by k-mer content. Nature Genetics, 2018, 50, 1474-1482.	21.4	198
3	Structural analysis by X-ray crystallography and calorimetry of a haemagglutinin component (HA1) of the progenitor toxin from Clostridium botulinum. Microbiology (United Kingdom), 2003, 149, 3361-3370.	1.8	69
4	LIN28A Modulates Splicing and Gene Expression Programs in Breast Cancer Cells. Molecular and Cellular Biology, 2015, 35, 3225-3243.	2.3	29
5	Early growth response 1 loops the <i>CYP2B6</i> promoter for synergistic activation by the distal and proximal nuclear receptors CAR and HNF4α. FEBS Letters, 2009, 583, 2126-2130.	2.8	23
6	Cohesin protein SMC1 represses the nuclear receptor CAR-mediated synergistic activation of a human P450 gene by xenobiotics. Biochemical Journal, 2006, 398, 125-133.	3.7	17
7	Multimodal regulatory elements within a hormone-specific super enhancer control a heterogeneous transcriptional response. Molecular Cell, 2022, 82, 803-815.e5.	9.7	14
8	Elements at the 5′ end of Xist harbor SPEN-independent transcriptional antiterminator activity. Nucleic Acids Research, 2020, 48, 10500-10517.	14.5	10
9	A ubiquitin-like domain is required for stabilizing the N-terminal ATPase module of human SMCHD1. Communications Biology, 2019, 2, 255.	4.4	8
10	Insight Into the Ontogeny of GnRH Neurons From Patients Born Without a Nose. Journal of Clinical Endocrinology and Metabolism, 2020, 105, 1538-1551.	3.6	7
11	A Statistical Method for Joint Estimation of <i>Cis</i> -eQTLs and Parent-of-Origin Effects Under Family Trio Design. Biometrics, 2019, 75, 864-874.	1.4	3
12	Cross-sectional Neuromuscular Phenotyping Study of Patients With Arhinia With <i>SMCHD1</i> Variants. Neurology, 2022, 98, .	1.1	3
13	Structural and functional consequences of SMCHD1 mutations associated with arhinia and muscular dystrophy. FASEB Journal, 2019, 33, 493.5.	0.5	0