

# Hiroyuki Ohashi

## List of Publications by Year in descending order

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Version: 2024-02-01

28  
papers

1,033  
citations

623734

14  
h-index

501196

28  
g-index

28  
all docs

28  
docs citations

28  
times ranked

962  
citing authors

#	ARTICLE	IF	CITATIONS
1	Safety of 222 nm UVC Irradiation to the Surgical Site in a Rabbit Model. <i>Photochemistry and Photobiology</i> , 2022, 98, 1365-1371.	2.5	8
2	Effect of ultraviolet C emitted from KrCl excimer lamp with or without bandpass filter to mouse epidermis. <i>PLoS ONE</i> , 2022, 17, e0267957.	2.5	7
3	Evaluation of Acute Reactions on Mouse Skin Irradiated with 222 and 235 nm UV-C. <i>Photochemistry and Photobiology</i> , 2021, 97, 770-777.	2.5	18
4	Re-Evaluation of Rat Corneal Damage by Short-Wavelength UV Revealed Extremely Less Hazardous Property of Far-UV-C. <i>Photochemistry and Photobiology</i> , 2021, 97, 505-516.	2.5	31
5	Exploratory clinical trial on the safety and bactericidal effect of 222-nm ultraviolet C irradiation in healthy humans. <i>PLoS ONE</i> , 2020, 15, e0235948.	2.5	85
6	Long-term Effects of 222-nm ultraviolet radiation C Sterilizing Lamps on Mice Susceptible to Ultraviolet Radiation. <i>Photochemistry and Photobiology</i> , 2020, 96, 853-862.	2.5	113
7	Ultraviolet C light with wavelength of 222 nm inactivates a wide spectrum of microbial pathogens. <i>Journal of Hospital Infection</i> , 2020, 105, 459-467.	2.9	114
8	Antigen-responsive fluorescent antibody probes generated by selective N-terminal modification of IgGs. <i>Chemical Communications</i> , 2018, 54, 12734-12737.	4.1	13
9	Cell-Free Technologies for Proteomics and Protein Engineering. <i>Protein and Peptide Letters</i> , 2016, 23, 819-827.	0.9	2
10	One-pot construction of Quenchbodies using antibody-binding proteins. <i>Analytical Methods</i> , 2016, 8, 7774-7779.	2.7	11
11	Insight into the Working Mechanism of Quenchbody: Transition of the Dye around Antibody Variable Region That Fluoresces upon Antigen Binding. <i>Bioconjugate Chemistry</i> , 2016, 27, 2248-2253.	3.6	34
12	Development of a novel immunoassay for herbal cannabis using a new fluorescent antibody probe, "Ultra Quenchbody". <i>Forensic Science International</i> , 2016, 266, 541-548.	2.2	12
13	Catalytic subunits of the phosphatase calcineurin interact with NF- $\kappa$ B-inducing kinase (NIK) and attenuate NIK-dependent gene expression. <i>Scientific Reports</i> , 2015, 5, 10758.	3.3	13
14	Next-Generation Technologies for Multiomics Approaches Including Interactome Sequencing. <i>BioMed Research International</i> , 2015, 2015, 1-9.	1.9	33
15	Optimal fusion of antibody binding domains resulted in higher affinity and wider specificity. <i>Journal of Bioscience and Bioengineering</i> , 2015, 120, 504-509.	2.2	14
16	Development of a rapid method for the quantitative determination of deoxynivalenol using Quenchbody. <i>Analytica Chimica Acta</i> , 2015, 888, 126-130.	5.4	28
17	Mitochondria-Nucleus Shuttling FK506-Binding Protein 51 Interacts with TRAF Proteins and Facilitates the RIG-I-Like Receptor-Mediated Expression of Type I IFN. <i>PLoS ONE</i> , 2014, 9, e95992.	2.5	31
18	Towards Personalized Medicine Mediated by in Vitro Virus-Based Interactome Approaches. <i>International Journal of Molecular Sciences</i> , 2014, 15, 6717-6724.	4.1	8

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19	Ultra Q-bodies: quench-based antibody probes that utilize dye-dye interactions with enhanced antigen-dependent fluorescence. <i>Scientific Reports</i> , 2014, 4, 4640.	3.3	70
20	Analysis of Transcription Factor Networks Using IVV Method. <i>Methods in Molecular Biology</i> , 2014, 1164, 15-22.	0.9	1
21	Detection of vimentin serine phosphorylation by multicolor Quenchbodies. <i>Biosensors and Bioelectronics</i> , 2013, 40, 17-23.	10.1	37
22	Efficiency of puromycin-based technologies mediated by release factors and a ribosome recycling factor. <i>Protein Engineering, Design and Selection</i> , 2013, 26, 533-537.	2.1	2
23	Peptide Screening Using PURE Ribosome Display. <i>Methods in Molecular Biology</i> , 2012, 805, 251-259.	0.9	13
24	Next-generation sequencing coupled with a cell-free display technology for high-throughput production of reliable interactome data. <i>Scientific Reports</i> , 2012, 2, 691.	3.3	25
25	“Quenchbodies” Quench-Based Antibody Probes That Show Antigen-Dependent Fluorescence. <i>Journal of the American Chemical Society</i> , 2011, 133, 17386-17394.	13.7	129
26	A Highly Controllable Reconstituted Cell-Free System -a Breakthrough in Protein Synthesis Research. <i>Current Pharmaceutical Biotechnology</i> , 2010, 11, 267-271.	1.6	52
27	Ribosome Display with the PURE Technology. <i>Methods in Molecular Biology</i> , 2010, 607, 219-225.	0.9	14
28	Efficient protein selection based on ribosome display system with purified components. <i>Biochemical and Biophysical Research Communications</i> , 2007, 352, 270-276.	2.1	115