

# Yoshinao Wada

## List of Publications by Year in descending order

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87  
papers

3,399  
citations

172457

29  
h-index

144013

57  
g-index

88  
all docs

88  
docs citations

88  
times ranked

3579  
citing authors

#	ARTICLE	IF	CITATIONS
1	Comparison of the methods for profiling glycoprotein glycansâ€”HUPO Human Disease Glycomics/Proteome Initiative multi-institutional study. <i>Glycobiology</i> , 2007, 17, 411-422.	2.5	382
2	Hydrophilic Affinity Isolation and MALDI Multiple-Stage Tandem Mass Spectrometry of Glycopeptides for Glycoproteomics. <i>Analytical Chemistry</i> , 2004, 76, 6560-6565.	6.5	317
3	Multiple Phenotypes in Phosphoglucosyltransferase 1 Deficiency. <i>New England Journal of Medicine</i> , 2014, 370, 533-542.	27.0	236
4	Derivatization for Stabilizing Sialic Acids in MALDI-MS. <i>Analytical Chemistry</i> , 2005, 77, 4962-4968.	6.5	165
5	Comparison of Methods for Profiling O-Glycosylation. <i>Molecular and Cellular Proteomics</i> , 2010, 9, 719-727.	3.8	136
6	Structure of serum transferrin in carbohydrate-deficient glycoprotein syndrome. <i>Biochemical and Biophysical Research Communications</i> , 1992, 189, 832-836.	2.1	130
7	Differential analysis of site-specific glycans on plasma and cellular fibronectins: application of a hydrophilic affinity method for glycopeptide enrichment. <i>Glycobiology</i> , 2005, 15, 1332-1340.	2.5	125
8	Site-specific analysis of N-glycans on haptoglobin in sera of patients with pancreatic cancer: A novel approach for the development of tumor markers. <i>International Journal of Cancer</i> , 2008, 122, 2301-2309.	5.1	125
9	Interlaboratory Study on Differential Analysis of Protein Glycosylation by Mass Spectrometry: The ABRF Glycoprotein Research Multi-Institutional Study 2012. <i>Molecular and Cellular Proteomics</i> , 2013, 12, 2935-2951.	3.8	103
10	Requirements for Laser-Induced Desorption/Ionization on Submicrometer Structures. <i>Analytical Chemistry</i> , 2005, 77, 5364-5369.	6.5	95
11	Oligosaccharide Profiles of the Prostate Specific Antigen in Free and Complexed Forms from the Prostate Cancer Patient Serum and in Seminal Plasma: a Glycopeptide Approach. <i>Glycobiology</i> , 2008, 18, 2-8.	2.5	95
12	De Novo Mutations in SLC35A2 Encoding a UDP-Galactose Transporter Cause Early-Onset Epileptic Encephalopathy. <i>Human Mutation</i> , 2013, 34, 1708-1714.	2.5	85
13	Calponin 3 Regulates Actin Cytoskeleton Rearrangement in Trophoblastic Cell Fusion. <i>Molecular Biology of the Cell</i> , 2010, 21, 3973-3984.	2.1	70
14	Quantitation of Saccharide Compositions of O-glycans by Mass Spectrometry of Glycopeptides and Its Application to Rheumatoid Arthritis. <i>Journal of Proteome Research</i> , 2010, 9, 1367-1373.	3.7	66
15	Structure elucidation of hemoglobin variants and other proteins by digit-printing method. <i>Mass Spectrometry Reviews</i> , 1989, 8, 379-434.	5.4	62
16	Structural analysis of human hemoglobin variants with field desorption mass spectrometry. <i>Biochimica Et Biophysica Acta (BBA) - Protein Structure</i> , 1981, 667, 233-241.	1.7	60
17	Ordered Porous Alumina Geometries and Surface Metals for Surface-Assisted Laser Desorption/Ionization of Biomolecules: A Possible Mechanistic Implications of Metal Surface Melting. <i>Analytical Chemistry</i> , 2007, 79, 9122-9127.	6.5	59
18	cDNA cloning, genomic cloning, and tissue-specific regulation of mouse cerebroside sulfotransferase. <i>FEBS Journal</i> , 2000, 267, 1909-1917.	0.2	58

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19	The Absence of Core Fucose Up-regulates GnT-III and Wnt Target Genes. <i>Journal of Biological Chemistry</i> , 2014, 289, 11704-11714.	3.4	50
20	Diagnosis of carbohydrate-deficient glycoprotein syndrome by matrix-assisted laser desorption time-of-flight mass spectrometry. <i>Biological Mass Spectrometry</i> , 1994, 23, 108-109.	0.5	49
21	Mass spectrometry for congenital disorders of glycosylation, CDG. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , 2006, 838, 3-8.	2.3	49
22	Calponin 3 regulates stress fiber formation in dermal fibroblasts during wound healing. <i>Archives of Dermatological Research</i> , 2013, 305, 571-584.	1.9	48
23	Reduction of organic dyes in matrix-assisted laser desorption/ionization and desorption/ionization on porous silicon. <i>Rapid Communications in Mass Spectrometry</i> , 2004, 18, 2811-2817.	1.5	44
24	Electrospray ionization mass spectra of hemoglobin and transferrin by a magnetic sector mass spectrometer. Comparison with theoretical isotopic distributions. <i>Rapid Communications in Mass Spectrometry</i> , 1992, 6, 9-13.	1.5	40
25	Antibody to annexin V has anti-phospholipid and lupus anticoagulant properties. <i>American Journal of Hematology</i> , 1995, 49, 347-348.	4.1	40
26	Mass spectrometry of apolipoprotein C-III, a simple analytical method for mucin-type O-glycosylation and its application to an autosomal recessive cutis laxa type-2 (ARCL2) patient. <i>Glycobiology</i> , 2012, 22, 1140-1144.	2.5	38
27	Limitations of galactose therapy in phosphoglucomutase 1 deficiency. <i>Molecular Genetics and Metabolism Reports</i> , 2017, 13, 33-40.	1.1	34
28	Quantitative analysis of polypropyleneglycol mixtures by desorption/ionization on porous silicon mass spectrometry. <i>International Journal of Mass Spectrometry</i> , 2005, 241, 43-48.	1.5	33
29	Advanced analytical methods for hemoglobin variants. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , 2002, 781, 291-301.	2.3	32
30	Rock-dependent calponin 3 phosphorylation regulates myoblast fusion. <i>Experimental Cell Research</i> , 2013, 319, 633-648.	2.6	30
31	Transferrin variants: Pitfalls in the diagnostics of Congenital disorders of glycosylation. <i>Clinical Biochemistry</i> , 2015, 48, 11-13.	1.9	30
32	Transferrin mutations at the glycosylation site complicate diagnosis of congenital disorders of glycosylation type I. <i>Journal of Inherited Metabolic Disease</i> , 2011, 34, 901-906.	3.6	29
33	Comparison of analytical methods for profiling N- and O-linked glycans from cultured cell lines. <i>Glycoconjugate Journal</i> , 2016, 33, 405-415.	2.7	25
34	N-Glycans of SREC-I (scavenger receptor expressed by endothelial cells): Essential role for ligand binding, trafficking and stability. <i>Glycobiology</i> , 2012, 22, 714-724.	2.5	24
35	O-Glycosylated IgA Rheumatoid Factor Induces IgA Deposits and Glomerulonephritis. <i>Journal of the American Society of Nephrology: JASN</i> , 2012, 23, 438-446.	6.1	23
36	Deficiency of N-acetylgalactosamine in O-linked oligosaccharides of IgA is a novel biologic marker for Crohn's disease. <i>Inflammatory Bowel Diseases</i> , 2012, 18, 1723-1734.	1.9	22

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37	Suppression of Heregulin $\hat{I}^2$ Signaling by the Single N-Glycan Deletion Mutant of Soluble ErbB3 Protein. <i>Journal of Biological Chemistry</i> , 2013, 288, 32910-32921.	3.4	22
38	Fast atom bombardment and tandem mass spectrometry for the characterization of hemoglobin variants including a new variant. <i>International Journal of Mass Spectrometry and Ion Processes</i> , 1992, 122, 219-229.	1.8	21
39	Dissociation Profile of Protonated Fucosyl Glycopeptides and Quantitation of Fucosylation Levels of Glycoproteins by Mass Spectrometry. <i>Journal of Proteome Research</i> , 2009, 8, 688-693.	3.7	21
40	Distinct Features of Matrix-Assisted 6 $\hat{I}^4$ m Infrared Laser Desorption/Ionization Mass Spectrometry in Biomolecular Analysis. <i>Analytical Chemistry</i> , 2009, 81, 6750-6755.	6.5	19
41	Quantitative change of IgA hinge O-glycan composition is a novel marker of therapeutic responses of IgA nephropathy. <i>Biochemical and Biophysical Research Communications</i> , 2012, 428, 339-342.	2.1	19
42	Mass spectrometry of transferrin and apolipoprotein C-III for diagnosis and screening of congenital disorder of glycosylation. <i>Glycoconjugate Journal</i> , 2016, 33, 297-307.	2.7	18
43	NUS1 mutation in a family with epilepsy, cerebellar ataxia, and tremor. <i>Epilepsy Research</i> , 2020, 164, 106371.	1.6	18
44	Mass spectrometry of transferrin glycoforms to detect congenital disorders of glycosylation: Site-specific profiles and pitfalls. <i>Proteomics</i> , 2016, 16, 3105-3110.	2.2	17
45	Cancer-associated alternative usage of multiple promoters of human GalCer sulfotransferase gene. <i>FEBS Journal</i> , 2000, 267, 2672-2679.	0.2	14
46	Determination of unique amino acid substitutions in protein variants by peptide mass mapping with FT-ICR MS. <i>Journal of the American Society for Mass Spectrometry</i> , 2006, 17, 508-513.	2.8	14
47	Mass Spectrometry in the Detection and Diagnosis of Congenital Disorders of Glycosylation. <i>European Journal of Mass Spectrometry</i> , 2007, 13, 101-103.	1.0	13
48	Evaluation of IgA1 O-glycosylation in Henoch-Schönlein Purpura Nephritis Using Mass Spectrometry. <i>Transplantation Proceedings</i> , 2019, 51, 1481-1487.	0.6	13
49	It Is Not Always Alcohol Abuse—A Transferrin Variant Impairing the CDT Test. <i>Alcohol and Alcoholism</i> , 2016, 51, 148-153.	1.6	12
50	Enantioselective Collision-Activated Dissociation of Gas-Phase Tryptophan Induced by Chiral Recognition of Protonated L-Alanine Peptides. <i>Origins of Life and Evolution of Biospheres</i> , 2017, 47, 161-167.	1.9	11
51	L-Fucose treatment of FUT8-CDG. <i>Molecular Genetics and Metabolism Reports</i> , 2020, 25, 100680.	1.1	11
52	The novel transferrin E592A variant impairs the diagnostics of congenital disorders of glycosylation. <i>Clinica Chimica Acta</i> , 2014, 436, 135-139.	1.1	10
53	Infrared matrix-assisted laser desorption/ionization mass spectrometry for quantification of glycosaminoglycans and gangliosides. <i>International Journal of Mass Spectrometry</i> , 2011, 305, 164-169.	1.5	9
54	Glycan Profiling: Label-Free Analysis of Glycoproteins. <i>Methods in Molecular Biology</i> , 2013, 951, 245-253.	0.9	9

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55	Anti-Ro52 antibody level is an important marker of fetal congenital heart block risk in anti-Ro/SSA antibody positive pregnancy. <i>Modern Rheumatology</i> , 2018, 28, 690-696.	1.8	9
56	Regulation of Vimentin Expression and Protease-mediated Vimentin Degradation during Differentiation of Human Monocytic Leukemia Cells. <i>Japanese Journal of Cancer Research</i> , 1997, 88, 484-491.	1.7	8
57	Measurement of serum salicylate levels by solid-phase extraction and desorption/ionization on silicon mass spectrometry. <i>Journal of Mass Spectrometry</i> , 2005, 40, 1000-1004.	1.6	8
58	Evaluation of serum carbohydrate-deficient transferrin by HPLC and MALDI-TOF MS. <i>Clinica Chimica Acta</i> , 2015, 448, 8-12.	1.1	8
59	Disruption of the Responsible Gene in a Phosphoglucomutase 1 Deficiency Patient by Homozygous Chromosomal Inversion. <i>JIMD Reports</i> , 2018, 43, 85-90.	1.5	8
60	Matrix-Assisted Laser Desorption/Ionization Mass Spectrometry to Detect Diagnostic Glycopeptide Markers of Congenital Disorders of Glycosylation. <i>Mass Spectrometry</i> , 2020, 9, A0084-A0084.	0.6	7
61	Mass Spectrometric Analysis of Synthetic Polymers Using Desorption/Ionization on Porous Silicon (DIOS)-Optimal Etching Conditions for DIOS Chips-. <i>Journal of the Mass Spectrometry Society of Japan</i> , 2004, 52, 142-148.	0.1	7
62	Chiral and Molecular Recognition through Protonation between Aromatic Amino Acids and Tripeptides Probed by Collision-Activated Dissociation in the Gas Phase. <i>Molecules</i> , 2018, 23, 162.	3.8	6
63	Congenital disorders of glycosylation type IIb with MOGS mutations cause early infantile epileptic encephalopathy, dysmorphic features, and hepatic dysfunction. <i>Brain and Development</i> , 2021, 43, 402-410.	1.1	6
64	Congenital disorder of glycosylation type Ic: Report of a Japanese case. <i>Brain and Development</i> , 2013, 35, 586-589.	1.1	5
65	Siblings with MAN1B1-CDG Showing Novel Biochemical Profiles. <i>Cells</i> , 2021, 10, 3117.	4.1	5
66	Decreased sialylation of IgA1 glycans associated with pneumococcal hemolytic uremic syndrome. <i>Pediatrics International</i> , 2013, 55, e143-5.	0.5	4
67	O-Linked Glycosylation Determines the Nephritogenic Potential of IgA Rheumatoid Factor. <i>Journal of the American Society of Nephrology: JASN</i> , 2014, 25, 1282-1290.	6.1	4
68	Apolipoprotein O glycoform profiling of 500 serum samples by matrix-assisted laser desorption/ionization mass spectrometry for diagnosis of congenital disorders of glycosylation. <i>Journal of Mass Spectrometry</i> , 2021, 56, e4597.	1.6	4
69	Primary ovarian insufficiency in a female with phosphomannomutase-2 gene ( <i>PMM2</i> ) mutations for congenital disorder of glycosylation. <i>Endocrine Journal</i> , 2021, 68, 605-611.	1.6	4
70	Trends in Glycoscience and Glycotechnology, 2021, 1, 1-10.		
71	Label-Free Analysis of O-glycosylation Site-Occupancy Based on the Signal Intensity of Glycopeptide/Peptide Ions. <i>Mass Spectrometry</i> , 2012, 1, A0008-A0008.	0.6	3
72	Translational balancing questioned: Unaltered glycosylation during disulfiram treatment in mannosyloligosaccharide alpha-1,2-mannosidase congenital disorders of glycosylation (MAN1B1-CDG). <i>JIMD Reports</i> , 2021, 60, 42-55.	1.5	3

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73	Electrospray Ionization Mass Spectrometry of Transferrin: Use of Quadrupole Mass Analyzers for Congenital Disorders of Glycosylation. <i>Mass Spectrometry</i> , 2022, 11, A0103-A0103.	0.6	3
74	Molecular Diagnosis of Congenital Disorders of Glycosylation. , 2008, , 319-322.		2
75	Congenital nephrotic syndrome with dysmorphic features and death in early infancy: Answers. <i>Pediatric Nephrology</i> , 2016, 31, 1283-1286.	1.7	2
76	Mass Spectrometry of Glycopeptides. , 2008, , 98-99.		2
77	Electrospray Ionization Mass Spectrometry of Apolipoprotein CIII to Evaluate &lt;i>O</i>-glycan Site Occupancy and Sialylation in Congenital Disorders of Glycosylation. <i>Mass Spectrometry</i> , 2022, 11, A0104-A0104.	0.6	2
78	Detection and Characterization of Protein Mutations by Mass Spectrometry. , 2002, , 681-692.		1
79	Congenital nephrotic syndrome with dysmorphic features and death in early infancy: Questions. <i>Pediatric Nephrology</i> , 2016, 31, 1281-1281.	1.7	1
80	Novel ALG12 variants and hydronephrosis in siblings with impaired N-glycosylation. <i>Brain and Development</i> , 2021, 43, 945-951.	1.1	1
81	Congenital Disorders of Glycosylation, Analytical Aspects. , 2015, , 119-128.		1
82	Tackling Difficulties in the Determination of O-Glycosylation Sites: Approaches to Mucin-type Glycoproteins. <i>Trends in Glycoscience and Glycotechnology</i> , 2008, 20, 173.	0.1	1
83	Glycomics and glycoproteomics.. <i>Seibutsu Butsuri Kagaku</i> , 2006, 50, 37-40.	0.1	0
84	Mass Spectrometry of Molecular Disease: Development of $\Delta$ m Diagnostics, Recollection. <i>Journal of the Mass Spectrometry Society of Japan</i> , 2013, 61, 35-41.	0.1	0
85	Congenital Disorders of Glycosylation: Analytical Aspects. , 2014, , 1-9.		0
86	Congenital Disorders of Glycosylation (CDG), Neuromuscular Related Diseases. , 2019, , 289-295.		0
87	Technical aspects of gel-based proteomics designed for elucidating an aryl hydrocarbon receptor complex. <i>Environmental Sciences: an International Journal of Environmental Physiology and Toxicology</i> , 2004, 11, 25-31.	0.1	0