## Thierry Hennet

List of Publications by Year in descending order

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201674 276875 2,834 42 27 41 h-index citations g-index papers 42 42 42 4421 all docs docs citations times ranked citing authors

#	Article	IF	CITATIONS
1	Glycosylation-Dependent Induction of Programmed Cell Death in Murine Adenocarcinoma Cells. Frontiers in Immunology, 2022, 13, 797759.	4.8	2
2	Intestinal inflammation alters mucosal carbohydrate foraging and monosaccharide incorporation into microbial glycans. Cellular Microbiology, 2021, 23, e13269.	2.1	10
3	Significance of fucose in intestinal health and disease. Molecular Microbiology, 2021, 115, 1086-1093.	2.5	28
4	Synthesis of photoactivable oligosaccharide derivatives from 1,2-cyclic carbamate building blocks and study of their interaction with carbohydrate-binding proteins. Carbohydrate Research, 2021, 508, 108399.	2.3	0
5	Increased Antibody Response to Fucosylated Oligosaccharides and Fucose-Carrying Bacteroides Species in Crohn's Disease. Frontiers in Microbiology, 2020, 11, 1553.	3.5	10
6	Emergence and significance of carbohydrate-specific antibodies. Genes and Immunity, 2020, 21, 224-239.	4.1	58
7	Limited Neonatal Carbohydrate-Specific Antibody Repertoire Consecutive to Partial Prenatal Transfer of Maternal Antibodies. Frontiers in Immunology, 2020, 11, 573629.	4.8	1
8	Antiviral potential of 3′-sialyllactose- and 6′-sialyllactose-conjugated dendritic polymers against human and avian influenza viruses. Scientific Reports, 2020, 10, 768.	3.3	45
9	Alleviation of Intestinal Inflammation by Oral Supplementation With 2-Fucosyllactose in Mice. Frontiers in Microbiology, 2019, 10, 1385.	3.5	49
10	Maternal Human Milk Oligosaccharide Profile Modulates the Impact of an Intervention with Iron and Galacto-Oligosaccharides in Kenyan Infants. Nutrients, 2019, 11, 2596.	4.1	35
11	Collagen glycosylation. Current Opinion in Structural Biology, 2019, 56, 131-138.	5.7	92
12	Biallelic <i>COLGALT1</i> variants are associated with cerebral small vessel disease. Annals of Neurology, 2018, 84, 843-853.	5.3	46
13	Mechanisms and consequences of intestinal dysbiosis. Cellular and Molecular Life Sciences, 2017, 74, 2959-2977.	5.4	401
14	Custom Glycosylation of Cells and Proteins Using Cyclic Carbamate-Derivatized Oligosaccharides. Cell Chemical Biology, 2017, 24, 1336-1346.e3.	5.2	5
15	Oral supplementation of healthy adults with 2′- <i>O</i> -fucosyllactose and lacto- <i>N</i> -neotetraose is well tolerated and shifts the intestinal microbiota. British Journal of Nutrition, 2016, 116, 1356-1368.	2.3	148
16	NANS-mediated synthesis of sialic acid is required for brain and skeletal development. Nature Genetics, 2016, 48, 777-784.	21.4	125
17	Breastfed at Tiffany's. Trends in Biochemical Sciences, 2016, 41, 508-518.	7.5	69
18	Collagen Accumulation in Osteosarcoma Cells lacking GLT25D1 Collagen Galactosyltransferase. Journal of Biological Chemistry, 2016, 291, 18514-18524.	3.4	49

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19	Giant mimivirus R707 encodes a glycogenin paralogue polymerizing glucose through $\hat{l}_{\pm}$ - and $\hat{l}^2$ -glycosidic linkages. Biochemical Journal, 2016, 473, 3451-3462.	3.7	7
20	Glycosylation site occupancy in health, congenital disorder of glycosylation and fatty liver disease. Scientific Reports, 2016, 6, 33927.	3.3	17
21	Congenital disorders of glycosylation: a concise chart of glycocalyx dysfunction. Trends in Biochemical Sciences, 2015, 40, 377-384.	7.5	109
22	Sialic acid catabolism drives intestinal inflammation and microbial dysbiosis in mice. Nature Communications, 2015, 6, 8141.	12.8	168
23	Selective proliferation of intestinal <i>Barnesiella</i> under fucosyllactose supplementation in mice. British Journal of Nutrition, 2014, 111, 1602-1610.	2.3	81
24	O-Linked glycosylation in Acanthamoeba polyphaga mimivirus. Glycobiology, 2014, 24, 703-714.	2.5	10
25	Decoding breast milk oligosaccharides. Swiss Medical Weekly, 2014, 144, w13927.	1.6	14
26	Milk oligosaccharide sialyl ( $\hat{l}\pm 2,3$ ) lactose activates intestinal CD11c $\langle \sup \rangle + \langle \sup \rangle$ cells through TLR4. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, 17444-17449.	7.1	89
27	Diseases of glycosylation beyond classical congenital disorders of glycosylation. Biochimica Et Biophysica Acta - General Subjects, 2012, 1820, 1306-1317.	2.4	113
28	Identification of Domains and Amino Acids Essential to the Collagen Galactosyltransferase Activity of GLT25D1. PLoS ONE, 2011, 6, e29390.	2.5	14
29	Mimivirus Collagen Is Modified by Bifunctional Lysyl Hydroxylase and Glycosyltransferase Enzyme. Journal of Biological Chemistry, 2011, 286, 43701-43709.	3.4	42
30	Milk sialyllactose influences colitis in mice through selective intestinal bacterial colonization. Journal of Experimental Medicine, 2010, 207, 2843-2854.	8.5	110
31	Core Glycosylation of Collagen Is Initiated by Two $\hat{I}^2(1-\langle i > O < /i >)$ Galactosyltransferases. Molecular and Cellular Biology, 2009, 29, 943-952.	2.3	126
32	Molecular Basis for Galactosylation of Core Fucose Residues in Invertebrates. Journal of Biological Chemistry, 2009, 284, 36223-36233.	3.4	48
33	Deficiency in COG5 causes a moderate form of congenital disorders of glycosylation. Human Molecular Genetics, 2009, 18, 4350-4356.	2.9	104
34	Congenital disorders of glycosylation: an update on defects affecting the biosynthesis of dolichol-linked oligosaccharides. Human Mutation, 2009, 30, 1628-1641.	2.5	166
35	From glycosylation disorders back to glycosylation: What have we learned?. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2009, 1792, 921-924.	3.8	12
36	How does a medical doctor become a glycobiologist. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2009, 1792, 824.	3.8	1

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37	CDG nomenclature: Time for a change!. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2009, 1792, 825-826.	3.8	123
38	Human RFT1 Deficiency Leads to a Disorder of N-Linked Glycosylation. American Journal of Human Genetics, 2008, 82, 600-606.	6.2	53
39	MPDU1 mutations underlie a novel human congenital disorder of glycosylation, designated type If. Journal of Clinical Investigation, 2001, 108, 1687-1695.	8.2	115
40	Multi-allelic origin of congenital disorder of glycosylation (CDG)-lc. Human Genetics, 2000, 106, 538-545.	3.8	25
41	Multi-allelic origin of congenital disorder of glycosylation (CDG)-lc. Human Genetics, 2000, 106, 538-545.	3 <b>.</b> 8	62
42	Molecular cloning of a human UDP-galactose:GlcNAcbeta1,3GalNAc beta1,3 galactosyltransferase gene encoding an O-linked core3-elongation enzyme. FEBS Journal, 1999, 263, 571-576.	0.2	52