Rasmus Iversen

List of Publications by Year in descending order

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| # | Article | IF | CITATIONS |
|----|--|------|-----------|
| 1 | High abundance of plasma cells secreting transglutaminase 2–specific IgA autoantibodies with limited somatic hypermutation in celiac disease intestinal lesions. Nature Medicine, 2012, 18, 441-445. | 15.2 | 210 |
| 2 | Strong Clonal Relatedness between Serum and Gut IgA despite Different Plasma Cell Origins. Cell Reports, 2017, 20, 2357-2367. | 2.9 | 74 |
| 3 | Transglutaminase 2–Specific Autoantibodies in Celiac Disease Target Clustered, N-Terminal Epitopes Not Displayed on the Surface of Cells. Journal of Immunology, 2013, 190, 5981-5991. | 0.4 | 69 |
| 4 | High-Throughput Single-Cell Analysis of B Cell Receptor Usage among Autoantigen-Specific Plasma Cells in Celiac Disease. Journal of Immunology, 2017, 199, 782-791. | 0.4 | 62 |
| 5 | Plasma Cells Are the Most Abundant Gluten Peptide MHC-expressing Cells in Inflamed Intestinal Tissues FromÂPatients With Celiac Disease. Gastroenterology, 2019, 156, 1428-1439.e10. | 0.6 | 61 |
| 6 | Activity-regulating structural changes and autoantibody epitopes in transglutaminase 2 assessed by hydrogen/deuterium exchange. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 17146-17151. | 3.3 | 51 |
| 7 | Enhanced B-Cell Receptor Recognition of the Autoantigen Transglutaminase 2 by Efficient Catalytic Self-Multimerization. PLoS ONE, 2015, 10, e0134922. | 1.1 | 39 |
| 8 | Efficient T cell–B cell collaboration guides autoantibody epitope bias and onset of celiac disease. Proceedings of the National Academy of Sciences of the United States of America, 2019, 116, 15134-15139. | 3.3 | 39 |
| 9 | Evidence That Pathogenic Transglutaminase 2 in Celiac Disease Derives From Enterocytes. Gastroenterology, 2020, 159, 788-790. | 0.6 | 37 |
| 10 | lgs as Substrates for Transglutaminase 2: Implications for Autoantibody Production in Celiac Disease. Journal of Immunology, 2015, 195, 5159-5168. | 0.4 | 30 |
| 11 | Autoimmunity provoked by foreign antigens. Science, 2020, 368, 132-133. | 6.0 | 29 |
| 12 | Structural Basis for Antigen Recognition by Transglutaminase 2-specific Autoantibodies in Celiac Disease. Journal of Biological Chemistry, 2015, 290, 21365-21375. | 1.6 | 27 |
| 13 | Longevity, clonal relationship, and transcriptional program of celiac disease–specific plasma cells. Journal of Experimental Medicine, 2021, 218, . | 4.2 | 25 |
| 14 | Dissecting the interaction between transglutaminase 2 and fibronectin. Amino Acids, 2017, 49, 489-500. | 1.2 | 23 |
| 15 | Transglutaminase 2 interactions with extracellular matrix proteins as probed with celiac disease autoantibodies. FEBS Journal, 2015, 282, 2063-2075. | 2.2 | 20 |
| 16 | Epitope-dependent Functional Effects of Celiac Disease Autoantibodies on Transglutaminase 2. Journal of Biological Chemistry, 2016, 291, 25542-25552. | 1.6 | 20 |
| 17 | Transglutaminase 2 strongly binds to an extracellular matrix component other than fibronectin via its second Câ€ŧerminal betaâ€barrel domain. FEBS Journal, 2016, 283, 3994-4010. | 2.2 | 20 |
| 18 | Multivalent pIX phage display selects for distinct and improved antibody properties. Scientific Reports, 2016. 6. 39066. | 1.6 | 14 |