

# Matthew R Lincoln

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/8599592/publications.pdf>

Version: 2024-02-01

28  
papers

2,856  
citations

430442

18  
h-index

580395

25  
g-index

31  
all docs

31  
docs citations

31  
times ranked

3918  
citing authors

| #  | ARTICLE   | IF  | CITATIONS |
|----|---|-----|-----------|
| 1  | Type I interferon transcriptional network regulates expression of coinhibitory receptors in human T cells. <i>Nature Immunology</i> , 2022, 23, 632-642.  | 7.0 | 54        |
| 2  | Vitamin D as disease-modifying therapy for multiple sclerosis?. <i>Expert Review of Clinical Immunology</i> , 2021, 17, 691-693.  | 1.3 | 3         |
| 3  | Epigenetic fine-mapping: identification of causal mechanisms for autoimmunity. <i>Current Opinion in Immunology</i> , 2020, 67, 50-56.  | 2.4 | 1         |
| 4  | Enhanced astrocyte responses are driven by a genetic risk allele associated with multiple sclerosis. <i>Nature Communications</i> , 2018, 9, 5337.  | 5.8 | 54        |
| 5  | Activated $\beta$ -catenin in Foxp3+ regulatory T cells links inflammatory environments to autoimmunity. <i>Nature Immunology</i> , 2018, 19, 1391-1402.  | 7.0 | 90        |
| 6  | Clinical Reasoning: A 34-year-old man with headache, diplopia, and hemiparesis. <i>Neurology</i> , 2016, 86, e24-8.   | 1.5 | 0         |
| 7  | Teaching Neuro Images : Large vagal nerve schwannoma presenting with hemorrhage and respiratory failure. <i>Neurology</i> , 2014, 82, e89-90.   | 1.5 | 0         |
| 8  | Exome sequencing identifies a novel multiple sclerosis susceptibility variant in the <i>TYK2</i> gene. <i>Neurology</i> , 2012, 79, 406-411.  | 1.5 | 56        |
| 9  | Robert Whytt, Benjamin Franklin, and the first probable case of multiple sclerosis. <i>Annals of Neurology</i> , 2012, 72, 307-311.   | 2.8 | 3         |
| 10 | Of mice and men: experimental autoimmune encephalitis and multiple sclerosis. <i>European Journal of Clinical Investigation</i> , 2011, 41, 1254-1258.  | 1.7 | 37        |
| 11 | Chronic cerebrospinal venous insufficiency and multiple sclerosis. <i>Annals of Neurology</i> , 2010, 68, 270-270.  | 2.8 | 1         |
| 12 | A ChIP-seq defined genome-wide map of vitamin D receptor binding: Associations with disease and evolution. <i>Genome Research</i> , 2010, 20, 1352-1360.  | 2.4 | 737       |
| 13 | Parent-of-origin effects at the major histocompatibility complex in multiple sclerosis. <i>Human Molecular Genetics</i> , 2010, 19, 3679-3689.  | 1.4 | 41        |
| 14 | Epistasis among <i>HLA-DRB1</i> , <i>HLA-DQA1</i> , and <i>HLA-DQB1</i> loci determines multiple sclerosis susceptibility. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009, 106, 7542-7547. | 3.3 | 148       |
| 15 | Expression of the Multiple Sclerosis-Associated MHC Class II Allele <i>HLA-DRB1*1501</i> Is Regulated by Vitamin D. <i>PLoS Genetics</i> , 2009, 5, e1000369.   | 1.5 | 442       |
| 16 | Analysis of 45 candidate genes for disease modifying activity in multiple sclerosis. <i>Journal of Neurology</i> , 2008, 255, 1215-1219.  | 1.8 | 19        |
| 17 | Parental transmission of <i>HLA-DRB1*15</i> in multiple sclerosis. <i>Human Genetics</i> , 2008, 122, 661-663.  | 1.8 | 47        |
| 18 | Methylation of class II transactivator gene promoter IV is not associated with susceptibility to Multiple Sclerosis. <i>BMC Medical Genetics</i> , 2008, 9, 63.   | 2.1 | 18        |

| #  | ARTICLE   | IF  | CITATIONS |
|----|---|-----|-----------|
| 19 | Parental non-inherited HLA resistance alleles do not confer protection against multiple sclerosis. <i>Journal of Neuroimmunology</i> , 2008, 196, 170-172.  | 1.1 | 3         |
| 20 | HLA class I alleles tag <i>HLA-DRB1</i> * <i>1501</i> haplotypes for differential risk in multiple sclerosis susceptibility. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2008, 105, 13069-13074. | 3.3 | 86        |
| 21 | Epigenetics in multiple sclerosis susceptibility: difference in transgenerational risk localizes to the major histocompatibility complex. <i>Human Molecular Genetics</i> , 2008, 18, 261-266.  | 1.4 | 89        |
| 22 | Evidence for genetic regulation of vitamin D status in twins with multiple sclerosis. <i>American Journal of Clinical Nutrition</i> , 2008, 88, 441-447.  | 2.2 | 223       |
| 23 | The Inheritance of Resistance Alleles in Multiple Sclerosis. <i>PLoS Genetics</i> , 2007, 3, e150.  | 1.5 | 109       |
| 24 | Transmission of class I/II multi-locus MHC haplotypes and multiple sclerosis susceptibility: accounting for linkage disequilibrium. <i>Human Molecular Genetics</i> , 2007, 16, 1951-1958.  | 1.4 | 33        |
| 25 | PRKCA and Multiple Sclerosis: Association in Two Independent Populations. <i>PLoS Genetics</i> , 2006, 2, e42.  | 1.5 | 45        |
| 26 | A predominant role for the HLA class II region in the association of the MHC region with multiple sclerosis. <i>Nature Genetics</i> , 2005, 37, 1108-1112.  | 9.4 | 295       |
| 27 | Complex interactions among MHC haplotypes in multiple sclerosis: susceptibility and resistance. <i>Human Molecular Genetics</i> , 2005, 14, 2019-2026.  | 1.4 | 212       |
| 28 | Suppressor Alleles in Multiple Sclerosis: Inheritance and Interactions. <i>PLoS Genetics</i> , 2005, preprint, e150.  | 1.5 | 0         |