## Judy H Cho

## List of Publications by Year in descending order

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		41344	27406
109	40,465	49	106
papers	citations	h-index	g-index
121	121	121	48822
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	The mutational constraint spectrum quantified from variation in 141,456 humans. Nature, 2020, 581, 434-443.	27.8	6,140
2	A frameshift mutation in NOD2 associated with susceptibility to Crohn's disease. Nature, 2001, 411, 603-606.	27.8	4,589
3	Host–microbe interactions have shaped the genetic architecture of inflammatory bowel disease. Nature, 2012, 491, 119-124.	27.8	4,038
4	A Genome-Wide Association Study Identifies $\langle i \rangle IL23R \langle i \rangle$ as an Inflammatory Bowel Disease Gene. Science, 2006, 314, 1461-1463.	12.6	2,739
5	Genome-wide association defines more than 30 distinct susceptibility loci for Crohn's disease. Nature Genetics, 2008, 40, 955-962.	21.4	2,422
6	Inflammatory Bowel Disease. New England Journal of Medicine, 2009, 361, 2066-2078.	27.0	2,369
7	Genome-wide meta-analysis increases to 71 the number of confirmed Crohn's disease susceptibility loci. Nature Genetics, 2010, 42, 1118-1125.	21.4	2,284
8	Association analyses identify 38 susceptibility loci for inflammatory bowel disease and highlight shared genetic risk across populations. Nature Genetics, 2015, 47, 979-986.	21.4	1,965
9	Genome-wide association study identifies new susceptibility loci for Crohn disease and implicates autophagy in disease pathogenesis. Nature Genetics, 2007, 39, 596-604.	21.4	1,633
10	Immunoglobulin A Coating Identifies Colitogenic Bacteria in Inflammatory Bowel Disease. Cell, 2014, 158, 1000-1010.	28.9	982
11	Deep resequencing of GWAS loci identifies independent rare variants associated with inflammatory bowel disease. Nature Genetics, 2011, 43, 1066-1073.	21.4	698
12	Inherited determinants of Crohn's disease and ulcerative colitis phenotypes: a genetic association study. Lancet, The, 2016, 387, 156-167.	13.7	607
13	Deletion polymorphism upstream of IRGM associated with altered IRGM expression and Crohn's disease. Nature Genetics, 2008, 40, 1107-1112.	21.4	604
14	Genome-wide association identifies multiple ulcerative colitis susceptibility loci. Nature Genetics, 2010, 42, 332-337.	21.4	572
15	Pervasive Sharing of Genetic Effects in Autoimmune Disease. PLoS Genetics, 2011, 7, e1002254.	3.5	540
16	Single-Cell Analysis of Crohn's Disease Lesions Identifies a Pathogenic Cellular Module Associated with Resistance to Anti-TNF Therapy. Cell, 2019, 178, 1493-1508.e20.	28.9	519
17	Prediction of complicated disease course for children newly diagnosed with Crohn's disease: a multicentre inception cohort study. Lancet, The, 2017, 389, 1710-1718.	13.7	482
18	Fine-mapping inflammatory bowel disease loci to single-variant resolution. Nature, 2017, 547, 173-178.	27.8	473

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19	A brief history of human disease genetics. Nature, 2020, 577, 179-189.	27.8	441
20	Crohn's disease-associated NOD2 variants share a signaling defect in response to lipopolysaccharide and peptidoglycan. Gastroenterology, 2003, 124, 140-146.	1.3	382
21	Recent Insights Into the Genetics of Inflammatory Bowel Disease. Gastroenterology, 2011, 140, 1704-1712.e2.	1.3	367
22	Dense genotyping of immune-related disease regions identifies nine new risk loci for primary sclerosing cholangitis. Nature Genetics, 2013, 45, 670-675.	21.4	339
23	Genetics of Inflammatory Bowel Diseases. Gastroenterology, 2015, 149, 1163-1176.e2.	1.3	319
24	Genomics and the Multifactorial Nature of Human Autoimmune Disease. New England Journal of Medicine, 2011, 365, 1612-1623.	27.0	299
25	Functional variants in the <i>LRRK2</i> gene confer shared effects on risk for Crohn's disease and Parkinson's disease. Science Translational Medicine, 2018, 10, .	12.4	273
26	Regulation of IL-8 and IL-1 $\hat{A}$ expression in Crohn's disease associated NOD2/CARD15 mutations. Human Molecular Genetics, 2004, 13, 1715-1725.	2.9	243
27	Defining Complex Contributions of NOD2/CARD15 Gene Mutations, Age at Onset, and Tobacco Use On Crohn's Disease Phenotypes. Inflammatory Bowel Diseases, 2003, 9, 281-289.	1.9	206
28	Heterogeneity of autoimmune diseases: pathophysiologic insights from genetics and implications for new therapies. Nature Medicine, 2015, 21, 730-738.	30.7	189
29	Machine Learning to Predict Mortality and Critical Events in a Cohort of Patients With COVID-19 in New York City: Model Development and Validation. Journal of Medical Internet Research, 2020, 22, e24018.	4.3	174
30	Different tissue phagocytes sample apoptotic cells to direct distinct homeostasis programs. Nature, 2016, 539, 565-569.	27.8	166
31	Sequencing an Ashkenazi reference panel supports population-targeted personal genomics and illuminates Jewish and European origins. Nature Communications, 2014, 5, 4835.	12.8	156
32	Transcriptional risk scores link GWAS to eQTLs and predict complications in Crohn's disease. Nature Genetics, 2017, 49, 1517-1521.	21.4	146
33	A Genome-Wide Scan of Ashkenazi Jewish Crohn's Disease Suggests Novel Susceptibility Loci. PLoS Genetics, 2012, 8, e1002559.	3.5	144
34	Genetic Factors and the Intestinal Microbiome Guide Development of Microbe-Based Therapies for Inflammatory Bowel Diseases. Gastroenterology, 2019, 156, 2174-2189.	1.3	132
35	Defects in Nicotinamide-adenine Dinucleotide Phosphate Oxidase Genes NOX1 and DUOX2 in Very Early Onset Inflammatory Bowel Disease. Cellular and Molecular Gastroenterology and Hepatology, 2015, 1, 489-502.	4.5	127
36	Genome-Wide Association Study Identifies African-Specific Susceptibility Loci in African Americans With Inflammatory Bowel Disease. Gastroenterology, 2017, 152, 206-217.e2.	1.3	120

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37	Gut microbiota density influences host physiology and is shaped by host and microbial factors. ELife, 2019, 8, .	6.0	118
38	A Pleiotropic Missense Variant in SLC39A8 Is Associated With Crohn's Disease and Human Gut Microbiome Composition. Gastroenterology, 2016, 151, 724-732.	1.3	109
39	Very Early Onset Inflammatory Bowel Disease: A Clinical Approach With a Focus on the Role of Genetics and Underlying Immune Deficiencies. Inflammatory Bowel Diseases, 2020, 26, 820-842.	1.9	100
40	Intestinal Inflammation Modulates the Expression of ACE2 and TMPRSS2 and Potentially Overlaps With the Pathogenesis of SARS-CoV-2–related Disease. Gastroenterology, 2021, 160, 287-301.e20.	1.3	98
41	Association of the V122I Hereditary Transthyretin Amyloidosis Genetic Variant With Heart Failure Among Individuals of African or Hispanic/Latino Ancestry. JAMA - Journal of the American Medical Association, 2019, 322, 2191.	7.4	93
42	Inflammatory Bowel Disease Genetics: Nod2. Annual Review of Medicine, 2007, 58, 401-416.	12.2	91
43	Limitations of Contemporary Guidelines for Managing Patients at High Genetic Risk of Coronary Artery Disease. Journal of the American College of Cardiology, 2020, 75, 2769-2780.	2.8	88
44	Toward a fine-scale population health monitoring system. Cell, 2021, 184, 2068-2083.e11.	28.9	78
45	Exome sequencing reveals a high prevalence of BRCA1 and BRCA2 founder variants in a diverse population-based biobank. Genome Medicine, 2020, 12, 2.	8.2	68
46	Challenges in IBD Research: Precision Medicine. Inflammatory Bowel Diseases, 2019, 25, S31-S39.	1.9	67
47	Insights into the genetic epidemiology of Crohn's and rare diseases in the Ashkenazi Jewish population. PLoS Genetics, 2018, 14, e1007329.	3.5	66
48	Characterization of Genetic Loci That Affect Susceptibility to Inflammatory Bowel Diseases in African Americans. Gastroenterology, 2015, 149, 1575-1586.	1.3	65
49	Genetic identification of a common collagen disease in Puerto Ricans via identity-by-descent mapping in a health system. ELife, 2017, 6, .	6.0	65
50	A myeloid–stromal niche and gp130 rescue in NOD2-driven Crohn's disease. Nature, 2021, 593, 275-281.	27.8	65
51	A Systematic Review of Monogenic Inflammatory Bowel Disease. Clinical Gastroenterology and Hepatology, 2022, 20, e653-e663.	4.4	57
52	Pattern Recognition Receptor Signaling in Human Dendritic Cells is Enhanced by ICOS Ligand and Modulated by the Crohn's Disease ICOSLG Risk Allele. Immunity, 2014, 40, 734-746.	14.3	55
53	Immunoglobulin A Targets a Unique Subset of the Microbiota in Inflammatory Bowel Disease. Cell Host and Microbe, 2021, 29, 83-93.e3.	11.0	53
54	A Frameshift in CSF2RB Predominant Among Ashkenazi Jews Increases Risk for Crohn's Disease and Reduces Monocyte Signaling via GM-CSF. Gastroenterology, 2016, 151, 710-723.e2.	1.3	51

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55	A protein-truncating R179X variant in RNF186 confers protection against ulcerative colitis. Nature Communications, 2016, 7, 12342.	12.8	50
56	Genome-wide polygenic score to predict chronic kidney disease across ancestries. Nature Medicine, 2022, 28, 1412-1420.	30.7	48
57	Magnetic Resonance Imaging Predicts Histopathological Composition of Ileal Crohn's Disease. Journal of Crohn's and Colitis, 2018, 12, 718-729.	1.3	45
58	Exome-wide evaluation of rare coding variants using electronic health records identifies new gene–phenotype associations. Nature Medicine, 2021, 27, 66-72.	30.7	44
59	Common and Rare Variant Prediction and Penetrance of IBD in a Large, Multi-ethnic, Health System-based Biobank Cohort. Gastroenterology, 2021, 160, 1546-1557.	1.3	43
60	Inflamed Ulcerative Colitis Regions Associated With MRGPRX2-Mediated Mast Cell Degranulation and Cell Activation Modules, Defining a New Therapeutic Target. Gastroenterology, 2021, 160, 1709-1724.	1.3	43
61	High-Throughput Characterization of Blood Serum Proteomics of IBD Patients with Respect to Aging and Genetic Factors. PLoS Genetics, 2017, 13, e1006565.	3.5	41
62	Blood and Intestine eQTLs from an Anti-TNF-Resistant Crohn's Disease Cohort Inform IBD Genetic Association Loci. Clinical and Translational Gastroenterology, 2016, 7, e177.	2.5	40
63	Microbial Engraftment and Efficacy of Fecal Microbiota Transplant for Clostridium Difficile in Patients With and Without Inflammatory Bowel Disease. Inflammatory Bowel Diseases, 2019, 25, 969-979.	1.9	38
64	Implementing genomic screening in diverse populations. Genome Medicine, 2021, 13, 17.	8.2	38
65	An Integrated Taxonomy for Monogenic Inflammatory Bowel Disease. Gastroenterology, 2022, 162, 859-876.	1.3	37
66	A common variant in PNPLA3 is associated with age at diagnosis of NAFLD in patients from a multi-ethnic biobank. Journal of Hepatology, 2020, 72, 1070-1081.	3.7	35
67	Utility of polygenic embryo screening for disease depends on the selection strategy. ELife, 2021, 10, .	6.0	34
68	Population-Based Penetrance of Deleterious Clinical Variants. JAMA - Journal of the American Medical Association, 2022, 327, 350.	7.4	34
69	Collagenous Colitis Is Associated With HLA Signature and Shares Genetic Risks With Other Immune-Mediated Diseases. Gastroenterology, 2020, 159, 549-561.e8.	1.3	31
70	Zebrafish modeling of intestinal injury, bacterial exposures, and medications defines epithelial in vivo responses relevant to human inflammatory bowel disease. DMM Disease Models and Mechanisms, 2019, 12, .	2.4	30
71	Contribution of higher risk genes and European admixture to Crohn $\hat{E}^1\!\!/\!\!4$ s disease in African Americans. Inflammatory Bowel Diseases, 2012, 18, 2277-2287.	1.9	29
72	Augmented intelligence with natural language processing applied to electronic health records for identifying patients with non-alcoholic fatty liver disease at risk for disease progression. International Journal of Medical Informatics, 2019, 129, 334-341.	3.3	29

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73	Natural language processing of electronic health records is superior to billing codes to identify symptom burden in hemodialysis patients. Kidney International, 2020, 97, 383-392.	5.2	27
74	Integrative Analysis of the Inflammatory Bowel Disease Serum Metabolome Improves Our Understanding of Genetic Etiology and Points to Novel Putative Therapeutic Targets. Gastroenterology, 2022, 162, 828-843.e11.	1.3	26
75	Transethnic Transferability of a Genome-Wide Polygenic Score for Coronary Artery Disease. Circulation Genomic and Precision Medicine, 2021, 14, e003092.	3.6	25
76	High-depth whole genome sequencing of an Ashkenazi Jewish reference panel: enhancing sensitivity, accuracy, and imputation. Human Genetics, 2018, 137, 343-355.	3.8	24
77	Singleâ€eell transcriptomics reveals conserved cell identities and fibrogenic phenotypes in zebrafish and human liver. Hepatology Communications, 2022, 6, 1711-1724.	4.3	24
78	Machine learning identifies novel blood protein predictors of penetrating and stricturing complications in newly diagnosed paediatric Crohn's disease. Alimentary Pharmacology and Therapeutics, 2021, 53, 281-290.	3.7	23
79	Whole-genome sequencing of African Americans implicates differential genetic architecture in inflammatory bowel disease. American Journal of Human Genetics, 2021, 108, 431-445.	6.2	21
80	Novel ultra-rare exonic variants identified in a founder population implicate cadherins in schizophrenia. Neuron, 2021, 109, 1465-1478.e4.	8.1	21
81	Institutional profile: translational pharmacogenomics at the Icahn School of Medicine at Mount Sinai. Pharmacogenomics, 2017, 18, 1381-1386.	1.3	20
82	Earlier Anti-Tumor Necrosis Factor Therapy of Crohn's Disease Correlates with Slower Progression of Bowel Damage. Digestive Diseases and Sciences, 2019, 64, 3274-3283.	2.3	20
83	Improved integrative framework combining association data with gene expression features to prioritize Crohn's disease genes. Human Molecular Genetics, 2015, 24, 4147-4157.	2.9	19
84	High-Throughput Identification of the Plasma Proteomic Signature of Inflammatory Bowel Disease. Journal of Crohn's and Colitis, 2019, 13, 462-471.	1.3	18
85	Neutralizing Anti-Granulocyte Macrophage-Colony Stimulating Factor Autoantibodies Recognize Post-Translational Glycosylations on Granulocyte Macrophage-Colony Stimulating Factor Years Before Diagnosis and Predict Complicated Crohnâ∈™s Disease. Gastroenterology, 2022, 163, 659-670.	1.3	18
86	Deep Analysis of the Peripheral Immune System in IBD Reveals New Insight in Disease Subtyping and Response to Monotherapy or Combination Therapy. Cellular and Molecular Gastroenterology and Hepatology, 2021, 12, 599-632.	4.5	17
87	Effector CD4+ T Cell Expression Signatures and Immune-Mediated Disease Associated Genes. PLoS ONE, 2012, 7, e38510.	2.5	16
88	Luminally polarized mural and vascular remodeling in ileal strictures of Crohn's disease. Human Pathology, 2018, 79, 42-49.	2.0	16
89	Emergent colectomy rates decreased while elective ileal pouch rates were stable over time: a nationwide inpatient sample study. International Journal of Colorectal Disease, 2019, 34, 1771-1779.	2.2	16
90	Prioritizing Crohn's disease genes by integrating association signals with gene expression implicates monocyte subsets. Genes and Immunity, 2019, 20, 577-588.	4.1	16

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91	Genome-wide polygenic risk score for retinopathy of type 2 diabetes. Human Molecular Genetics, 2021, 30, 952-960.	2.9	14
92	Molecular Characterization of Limited Ulcerative Colitis Reveals Novel Biology and Predictors of Disease Extension. Gastroenterology, 2021, 161, 1953-1968.e15.	1.3	14
93	Coronary Risk Estimation Based on Clinical Data in Electronic Health Records. Journal of the American College of Cardiology, 2022, 79, 1155-1166.	2.8	14
94	Constrictive and Hypertrophic Strictures in Ileal Crohn's Disease. Clinical Gastroenterology and Hepatology, 2022, 20, e1292-e1304.	4.4	13
95	Bridging the Gap Between Host Immune Response and Intestinal Dysbiosis in Inflammatory Bowel Disease: Does Immunoglobulin A Mark the Spot?. Clinical Gastroenterology and Hepatology, 2015, 13, 842-846.	4.4	10
96	A Role for CXCR3 Ligands as Biomarkers of Post-Operative Crohn's Disease Recurrence. Journal of Crohn's and Colitis, 2022, 16, 900-910.	1.3	10
97	Genome-Wide Association Studies: Present Status and Future Directions. Gastroenterology, 2010, 138, 1668-1672.e1.	1.3	9
98	Evaluation of ileal Crohn's disease response to TNF antagonists: Validation of MR enterography for assessing response. Initial results. European Journal of Radiology Open, 2020, 7, 100217.	1.6	9
99	Luminal Narrowing Alone Allows an Accurate Diagnosis of Crohn's Disease Small Bowel Strictures at Cross-Sectional Imaging. Journal of Crohn's and Colitis, 2021, 15, 1009-1018.	1.3	8
100	Stratification of risk of progression to colectomy in ulcerative colitis via measured and predicted gene expression. American Journal of Human Genetics, 2021, 108, 1765-1779.	6.2	6
101	The heritable immune system. Nature Biotechnology, 2015, 33, 608-609.	17.5	5
102	The Promise of Epigenetics. Has It Delivered New Insights?. Digestive Diseases, 2016, 34, 12-19.	1.9	4
103	Genetic pleiotropy of <i>ERCC6</i> lossâ€ofâ€function and deleterious missense variants links retinal dystrophy, arrhythmia, and immunodeficiency in diverse ancestries. Human Mutation, 2021, 42, 969-977.	2.5	3
104	From single-target to cellular niche targeting in Crohn's disease: intercepting bad communications. EBioMedicine, 2021, 74, 103690.	6.1	3
105	Polygenic risk score for alcohol drinking behavior improves prediction of inflammatory bowel disease risk. Human Molecular Genetics, 2021, 30, 514-523.	2.9	2
106	Genome-First Recall of Healthy Individuals by Polygenic Risk Score Reveals Differences in Coronary Artery Calcium. American Heart Journal, 2022, 250, 29-29.	2.7	1
107	P-175â€fPleiotropic Effects of Novel Functional LRRK2 Variation on Crohn's Disease and Parkinson's Disease Risk. Inflammatory Bowel Diseases, 2016, 22, S62-S63.	1.9	0
108	Risk Alleles for Drug Targets: Genomic Markers of Drug Response. , 2019, , 333-341.		0

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109	SUN-032 Exome Sequencing Reveals that Pathogenic RET Variants Occur at Higher Prevalence Than Previously Recognized: Data from a US Health System Biobank. Journal of the Endocrine Society, 2019, 3, .	0.2	0