## Ignaty Leshchiner

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Combined tumor and immune signals from genomes or transcriptomes predict outcomes of checkpoint inhibition in melanoma. Cell Reports Medicine, 2022, 3, 100500.	6.5	13
2	Characterizing genetic intra-tumor heterogeneity across 2,658 human cancer genomes. Cell, 2021, 184, 2239-2254.e39.	28.9	260
3	Longitudinal Single-Cell Dynamics of Chromatin Accessibility and Mitochondrial Mutations in Chronic Lymphocytic Leukemia Mirror Disease History. Cancer Discovery, 2021, 11, 3048-3063.	9.4	31
4	Parallel Genomic Alterations of Antigen and Payload Targets Mediate Polyclonal Acquired Clinical Resistance to Sacituzumab Govitecan in Triple-Negative Breast Cancer. Cancer Discovery, 2021, 11, 2436-2445.	9.4	69
5	Molecular features of exceptional response to neoadjuvant anti-androgen therapy in high-risk localized prostate cancer. Cell Reports, 2021, 36, 109665.	6.4	24
6	The RNA helicase Ddx21 controls Vegfc-driven developmental lymphangiogenesis by balancing endothelial cell ribosome biogenesis and p53 function. Nature Cell Biology, 2021, 23, 1136-1147.	10.3	17
7	The cationic amino acid exporter Slc7a7 is induced and vital in tissue macrophages with sustained efferocytic activity. Journal of Cell Science, 2020, 133, .	2.0	8
8	Distinct evolutionary paths in chronic lymphocytic leukemia during resistance to the graft-versus-leukemia effect. Science Translational Medicine, 2020, 12, .	12.4	17
9	The evolutionary history of 2,658 cancers. Nature, 2020, 578, 122-128.	27.8	690
10	The Human Tumor Atlas Network: Charting Tumor Transitions across Space and Time at Single-Cell Resolution. Cell, 2020, 181, 236-249.	28.9	334
11	Butler enables rapid cloud-based analysis of thousands of human genomes. Nature Biotechnology, 2020, 38, 288-292.	17.5	11
12	Genomic Profiling of Smoldering Multiple Myeloma Identifies Patients at a High Risk of Disease Progression. Journal of Clinical Oncology, 2020, 38, 2380-2389.	1.6	110
13	Mutations in RABL3 alter KRAS prenylation and are associated with hereditary pancreatic cancer. Nature Genetics, 2019, 51, 1308-1314.	21.4	47
14	Liquid versus tissue biopsy for detecting acquired resistance and tumor heterogeneity in gastrointestinal cancers. Nature Medicine, 2019, 25, 1415-1421.	30.7	359
15	Mitochondrial Reprogramming Underlies Resistance to BCL-2 Inhibition in Lymphoid Malignancies. Cancer Cell, 2019, 36, 369-384.e13.	16.8	224
16	A Murine Model of Chronic Lymphocytic Leukemia Based on B Cell-Restricted Expression of Sf3b1 Mutation and Atm Deletion. Cancer Cell, 2019, 35, 283-296.e5.	16.8	71
17	Tfap2a is a novel gatekeeper of nephron differentiation during kidney development. Development (Cambridge), 2019, 146, .	2.5	41
18	Growth dynamics in naturally progressing chronic lymphocytic leukaemia. Nature, 2019, 570, 474-479.	27.8	86

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19	T Cell Determinants of Response and Resistance to PD-1 Blockade in Richter's Transformation. Blood, 2019, 134, 680-680.	1.4	1
20	MCL-1 and PKA/AMPK Axis Fuel Venetoclax Resistance in Lymphoid Cancers. Blood, 2019, 134, 1284-1284.	1.4	3
21	Distinct Evolutionary Patterns in Chronic Lymphocytic Leukemia (CLL) during Resistance to Graft-Versus-Leukemia (GvL). Blood, 2019, 134, 516-516.	1.4	0
22	Distinct mutational signatures characterize concurrent loss of polymerase proofreading and mismatch repair. Nature Communications, 2018, 9, 1746.	12.8	142
23	Molecular subtypes of diffuse large B cell lymphoma are associated with distinct pathogenic mechanisms and outcomes. Nature Medicine, 2018, 24, 679-690.	30.7	1,224
24	Heterogeneity and Coexistence of T790M and T790 Wild-Type Resistant Subclones Drive Mixed Response to Third-Generation Epidermal Growth Factor Receptor Inhibitors in Lung Cancer. JCO Precision Oncology, 2018, 2018, 1-15.	3.0	17
25	Comment on "DNA damage is a pervasive cause of sequencing errors, directly confounding variant identification― Science, 2018, 361, .	12.6	2
26	Widespread Chromosomal Losses and Mitochondrial DNA Alterations as Genetic Drivers in Hürthle Cell Carcinoma. Cancer Cell, 2018, 34, 242-255.e5.	16.8	185
27	An integrated clinical program and crowdsourcing strategy for genomic sequencing and Mendelian disease gene discovery. Npj Genomic Medicine, 2018, 3, 21.	3.8	24
28	Activating MAPK Pathway Mutations Mediate Primary Resistance to PI3K Inhibitors in Chronic Lymphocytic Leukemia (CLL). Blood, 2018, 132, 587-587.	1.4	43
29	Genetic Determinants of Venetoclax Resistance in Lymphoid Malignancies. Blood, 2018, 132, 893-893.	1.4	4
30	Clonal and Single Cell Dynamics of Resistance to Graft-Versus-Leukemia (GvL) in Chronic Lymphocytic Leukemia (CLL). Blood, 2018, 132, 820-820.	1.4	0
31	Comprehensive Molecular Characterization of Pheochromocytoma and Paraganglioma. Cancer Cell, 2017, 31, 181-193.	16.8	532
32	The zebrafish kidney mutant zeppelin reveals that brca2/fancd1 is essential for pronephros development. Developmental Biology, 2017, 428, 148-163.	2.0	38
33	Integrated Molecular Characterization of Uterine Carcinosarcoma. Cancer Cell, 2017, 31, 411-423.	16.8	309
34	Polyclonal Secondary <i>FGFR2</i> Mutations Drive Acquired Resistance to FGFR Inhibition in Patients with FGFR2 Fusion–Positive Cholangiocarcinoma. Cancer Discovery, 2017, 7, 252-263.	9.4	384
35	A mutational signature reveals alterations underlying deficient homologous recombination repair in breast cancer. Nature Genetics, 2017, 49, 1476-1486.	21.4	427
36	Integrated Genomic Characterization of Pancreatic Ductal Adenocarcinoma. Cancer Cell, 2017, 32, 185-203.e13.	16.8	1,428

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#	Article	IF	CITATIONS
37	Scalable whole-exome sequencing of cell-free DNA reveals high concordance with metastatic tumors. Nature Communications, 2017, 8, 1324.	12.8	584
38	Resolving the phylogenetic origin of glioblastoma via multifocal genomic analysis of pre-treatment and treatment-resistant autopsy specimens. Npj Precision Oncology, 2017, 1, 33.	5.4	27
39	The evolutionary landscape of chronic lymphocytic leukemia treated with ibrutinib targeted therapy. Nature Communications, 2017, 8, 2185.	12.8	148
40	An argument for early genomic sequencing in atypical cases: a <i>WISP3</i> variant leads to diagnosis of progressive pseudorheumatoid arthropathy of childhood. Rheumatology, 2016, 55, kev367.	1.9	6
41	Loss of function mutation in <i>LOX</i> causes thoracic aortic aneurysm and dissection in humans. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, 8759-8764.	7.1	144
42	Molecular Mechanisms of Resistance to First- and Second-Generation ALK Inhibitors in <i>ALK</i>	9.4	919
43	Resensitization to Crizotinib by the Lorlatinib <i>ALK</i> Resistance Mutation L1198F. New England Journal of Medicine, 2016, 374, 54-61.	27.0	433
44	The Landscape of Dynamic Genetic Changes in Ibrutinib-Treated CLL. Blood, 2016, 128, 188-188.	1.4	3
45	Inherited <i>CHST11/MIR3922</i> deletion is associated with a novel recessive syndrome presenting with skeletal malformation and malignant lymphoproliferative disease. Molecular Genetics & amp; Genomic Medicine, 2015, 3, 413-423.	1.2	11
46	Paired exome analysis of Barrett's esophagus and adenocarcinoma. Nature Genetics, 2015, 47, 1047-1055.	21.4	310
47	The Molecular Taxonomy of Primary Prostate Cancer. Cell, 2015, 163, 1011-1025.	28.9	2,435
48	Comprehensive assessment of cancer missense mutation clustering in protein structures. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, E5486-95.	7.1	195
49	Summarizing polygenic risks for complex diseases in a clinical whole-genome report. Genetics in Medicine, 2015, 17, 536-544.	2.4	34
50	An international effort towards developing standards for best practices in analysis, interpretation and reporting of clinical genome sequencing results in the CLARITY Challenge. Genome Biology, 2014, 15, R53.	9.6	101
51	Synthesis and physical behavior of amphiphilic dendrimers with layered organization of hydrophilic and hydrophobic blocks. Colloid and Polymer Science, 2013, 291, 927-936.	2.1	5
52	Rapid identification of kidney cyst mutations by whole exome sequencing in zebrafish. Development (Cambridge), 2013, 140, 4445-4451.	2.5	43
53	Specific temperature-induced perturbations of secondary mRNA structures are associated with the cold-adapted temperature-sensitive phenotype of influenza A virus. RNA Biology, 2012, 9, 1266-1274.	3.1	17
54	Bithiophenesilane-Based Dendronized Polymers: Facile Synthesis and Properties of Novel Highly Branched Organosilicon Macromolecular Structures. Macromolecules, 2012, 45, 2014-2024.	4.8	35

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55	Synthesis of Carbosilane Dendrimers with Variable Distance between Branching Nodes. Macromolecules, 2012, 45, 8796-8804.	4.8	18
56	Mutation mapping and identification by whole-genome sequencing. Genome Research, 2012, 22, 1541-1548.	5.5	126
57	Combinatorial Approach to Determine Functional Group Effects on Lipidoid-Mediated siRNA Delivery. Bioconjugate Chemistry, 2010, 21, 1448-1454.	3.6	64
58	Structure of carbosilane amphiphilic liquid-crystalline codendrimers in bulk and in thin (Langmuir) films. Russian Chemical Bulletin, 2008, 57, 2101-2110.	1.5	2
59	Liquid Crystal Codendrimers with a Statistical Distribution of Phenolic and Mesogenic Groups: Behavior as Langmuir and Langmuirâ^'Blodgett Films. Langmuir, 2008, 24, 11082-11088.	3.5	12
60	Detection of Explosives using nanofibrous membranes. , 2008, , .		1
61	Organized Monolayers of Carbosilane Dendrimers with Mesogenic Terminal Groups. Macromolecules, 2005, 38, 8028-8035.	4.8	23