

# Anastasia Khvorova

## List of Publications by Year in descending order

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Version: 2024-02-01

68  
papers

17,441  
citations

117625

34  
h-index

98798

67  
g-index

75  
all docs

75  
docs citations

75  
times ranked

21497  
citing authors

#	ARTICLE	IF	CITATIONS
1	An RNAi therapeutic targeting hepatic DGAT2 in a genetically obese mouse model of nonalcoholic steatohepatitis. <i>Molecular Therapy</i> , 2022, 30, 1329-1342.	8.2	18
2	RNAi-based modulation of IFN- $\beta$ signaling in skin. <i>Molecular Therapy</i> , 2022, 30, 2709-2721.	8.2	13
3	Delivering siRNA Compounds During HOPE to Modulate Organ Function: A Proof-of-concept Study in a Rat Liver Transplant Model. <i>Transplantation</i> , 2022, 106, 1565-1576.	1.0	13
4	Chemical optimization of siRNA for safe and efficient silencing of placental sFLT1. <i>Molecular Therapy - Nucleic Acids</i> , 2022, 29, 135-149.	5.1	15
5	Modulation of DNA transcription: The future of ASO therapeutics?. <i>Cell</i> , 2022, 185, 2011-2013.	28.9	5
6	PK-modifying anchors significantly alter clearance kinetics, tissue distribution, and efficacy of therapeutics siRNAs. <i>Molecular Therapy - Nucleic Acids</i> , 2022, 29, 116-132.	5.1	7
7	Docosanoic acid conjugation to siRNA enables functional and safe delivery to skeletal and cardiac muscles. <i>Molecular Therapy</i> , 2021, 29, 1382-1394.	8.2	37
8	The NIH Somatic Cell Genome Editing program. <i>Nature</i> , 2021, 592, 195-204.	27.8	84
9	ALM2 regulates anti-tumor immunity and is a viable therapeutic target for melanoma. <i>Journal of Experimental Medicine</i> , 2021, 218, .	8.5	34
10	Gene therapy with AR isoform 2 rescues spinal and bulbar muscular atrophy phenotype by modulating AR transcriptional activity. <i>Science Advances</i> , 2021, 7, .	10.3	20
11	Structurally constrained phosphonate internucleotide linkage impacts oligonucleotide-enzyme interaction, and modulates siRNA activity and allele specificity. <i>Nucleic Acids Research</i> , 2021, 49, 12069-12088.	14.5	8
12	Comparative route of administration studies using therapeutic siRNAs show widespread gene modulation in Dorset sheep. <i>JCI Insight</i> , 2021, 6, .	5.0	9
13	Loss of huntingtin function slows synaptic vesicle endocytosis in striatal neurons from the httQ140/Q140 mouse model of Huntington's disease. <i>Neurobiology of Disease</i> , 2020, 134, 104637.	4.4	24
14	Enriched chitosan nanoparticles loaded with siRNA are effective in lowering Huntington's disease gene expression following intranasal administration. <i>Nanomedicine: Nanotechnology, Biology, and Medicine</i> , 2020, 24, 102119.	3.3	55
15	Hydrophobically Modified let-7b miRNA Enhances Biodistribution to NSCLC and Downregulates HMGA2 In Vivo. <i>Molecular Therapy - Nucleic Acids</i> , 2020, 19, 267-277.	5.1	39
16	Cell Type Impacts Accessibility of mRNA to Silencing by RNA Interference. <i>Molecular Therapy - Nucleic Acids</i> , 2020, 21, 384-393.	5.1	20
17	The chemical structure and phosphorothioate content of hydrophobically modified siRNAs impact extrahepatic distribution and efficacy. <i>Nucleic Acids Research</i> , 2020, 48, 7665-7680.	14.5	32
18	Single-Stranded Phosphorothioated Regions Enhance Cellular Uptake of Cholesterol-Conjugated siRNA but Not Silencing Efficacy. <i>Molecular Therapy - Nucleic Acids</i> , 2020, 21, 991-1005.	5.1	22

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19	2'-O-Methyl at 20-mer Guide Strand 3' Termini May Negatively Affect Target Silencing Activity of Fully Chemically Modified siRNA. <i>Molecular Therapy - Nucleic Acids</i> , 2020, 21, 266-277.	5.1	10
20	Data on enrichment of chitosan nanoparticles for intranasal delivery of oligonucleotides to the brain. <i>Data in Brief</i> , 2020, 28, 105093.	1.0	3
21	A divalent siRNA chemical scaffold for potent and sustained modulation of gene expression throughout the central nervous system. <i>Nature Biotechnology</i> , 2019, 37, 884-894.	17.5	126
22	Hydrophobicity drives the systemic distribution of lipid-conjugated siRNAs via lipid transport pathways. <i>Nucleic Acids Research</i> , 2019, 47, 1070-1081.	14.5	87
23	Serum Deprivation of Mesenchymal Stem Cells Improves Exosome Activity and Alters Lipid and Protein Composition. <i>Science</i> , 2019, 16, 230-241.	4.1	61
24	Nucleic Acid Therapeutics for Neurological Diseases. <i>Neurotherapeutics</i> , 2019, 16, 245-247.	4.4	16
25	The valency of fatty acid conjugates impacts siRNA pharmacokinetics, distribution, and efficacy in vivo. <i>Journal of Controlled Release</i> , 2019, 302, 116-125.	9.9	48
26	Gene Silencing With siRNA (RNA Interference): A New Therapeutic Option During Ex Vivo Machine Liver Perfusion Preservation. <i>Liver Transplantation</i> , 2019, 25, 140-151.	2.4	47
27	Diverse lipid conjugates for functional extra-hepatic siRNA delivery in vivo. <i>Nucleic Acids Research</i> , 2019, 47, 1082-1096.	14.5	122
28	Rac1 Activity Is Modulated by Huntingtin and Dysregulated in Models of Huntington's Disease. <i>Journal of Huntington's Disease</i> , 2019, 8, 53-69.	1.9	23
29	Editorial: <i>Nucleic Acids Research and Nucleic Acid Therapeutics</i> . <i>Nucleic Acids Research</i> , 2018, 46, 1563-1564.	14.5	8
30	Efficient Gene Silencing in Brain Tumors with Hydrophobically Modified siRNAs. <i>Molecular Cancer Therapeutics</i> , 2018, 17, 1251-1258.	4.1	14
31	Comparison of partially and fully chemically-modified siRNA in conjugate-mediated delivery in vivo. <i>Nucleic Acids Research</i> , 2018, 46, 2185-2196.	14.5	125
32	Loading of Extracellular Vesicles with Hydrophobically Modified siRNAs. <i>Methods in Molecular Biology</i> , 2018, 1740, 199-214.	0.9	13
33	Hydrophobicity of Lipid-Conjugated siRNAs Predicts Productive Loading to Small Extracellular Vesicles. <i>Molecular Therapy</i> , 2018, 26, 1520-1528.	8.2	31
34	Chitosan-Mangafodipir nanoparticles designed for intranasal delivery of siRNA and DNA to brain. <i>Journal of Drug Delivery Science and Technology</i> , 2018, 43, 453-460.	3.0	41
35	Functional features defining the efficacy of cholesterol-conjugated, self-deliverable, chemically modified siRNAs. <i>Nucleic Acids Research</i> , 2018, 46, 10905-10916.	14.5	48
36	RNAi modulation of placental sFLT1 for the treatment of preeclampsia. <i>Nature Biotechnology</i> , 2018, 36, 1164-1173.	17.5	126

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37	Minimal information for studies of extracellular vesicles 2018 (MISEV2018): a position statement of the International Society for Extracellular Vesicles and update of the MISEV2014 guidelines. <i>Journal of Extracellular Vesicles</i> , 2018, 7, 1535750.	12.2	6,961
38	Exosomes Produced from 3D Cultures of MSCs by Tangential Flow Filtration Show Higher Yield and Improved Activity. <i>Molecular Therapy</i> , 2018, 26, 2838-2847.	8.2	309
39	Nuclear Localization of Huntingtin mRNA Is Specific to Cells of Neuronal Origin. <i>Cell Reports</i> , 2018, 24, 2553-2560.e5.	6.4	34
40	Optimized Cholesterol-siRNA Chemistry Improves Productive Loading onto Extracellular Vesicles. <i>Molecular Therapy</i> , 2018, 26, 1973-1982.	8.2	65
41	Heavily and fully modified RNAs guide efficient SpyCas9-mediated genome editing. <i>Nature Communications</i> , 2018, 9, 2641.	12.8	83
42	Improving siRNA Delivery <i>In Vivo</i> Through Lipid Conjugation. <i>Nucleic Acid Therapeutics</i> , 2018, 28, 128-136.	3.6	90
43	Transvascular Delivery of Hydrophobically Modified siRNAs: Gene Silencing in the Rat Brain upon Disruption of the Blood-Brain Barrier. <i>Molecular Therapy</i> , 2018, 26, 2580-2591.	8.2	36
44	Novel Cluster and Monomer-Based GalNAc Structures Induce Effective Uptake of siRNAs <i>In Vitro</i> and <i>In Vivo</i> . <i>Bioconjugate Chemistry</i> , 2018, 29, 2478-2488.	3.6	32
45	Disrupting The Brain Keeper To Allow Silencing Of Deleterious Genes In The Nervous System. , 2018, , .		0
46	The chemical evolution of oligonucleotide therapies of clinical utility. <i>Nature Biotechnology</i> , 2017, 35, 238-248.	17.5	816
47	Synthesis and Evaluation of Parenchymal Retention and Efficacy of a Metabolically Stable <i>N</i> -Phosphocholine- <i>N</i> -docosahexaenoyl-serine siRNA Conjugate in Mouse Brain. <i>Bioconjugate Chemistry</i> , 2017, 28, 1758-1766.	3.6	33
48	5′, -Vinylphosphonate improves tissue accumulation and efficacy of conjugated siRNAs <i>in vivo</i> . <i>Nucleic Acids Research</i> , 2017, 45, 7581-7592.	14.5	83
49	Oligonucleotide Therapeutics – A New Class of Cholesterol-Lowering Drugs. <i>New England Journal of Medicine</i> , 2017, 376, 4-7.	27.0	128
50	Pharmacokinetic Profiling of Conjugated Therapeutic Oligonucleotides: A High-Throughput Method Based Upon Serial Blood Microsampling Coupled to Peptide Nucleic Acid Hybridization Assay. <i>Nucleic Acid Therapeutics</i> , 2017, 27, 323-334.	3.6	37
51	Visualization of self-delivering hydrophobically modified siRNA cellular internalization. <i>Nucleic Acids Research</i> , 2017, 45, 15-25.	14.5	119
52	Loading of Extracellular Vesicles with Chemically Stabilized Hydrophobic siRNAs for the Treatment of Disease in the Central Nervous System. <i>Bio-protocol</i> , 2017, 7, .	0.4	9
53	A High-throughput Assay for mRNA Silencing in Primary Cortical Neurons <i>in vitro</i> with Oligonucleotide Therapeutics. <i>Bio-protocol</i> , 2017, 7, .	0.4	6
54	High-resolution proteomic and lipidomic analysis of exosomes and microvesicles from different cell sources. <i>Journal of Extracellular Vesicles</i> , 2016, 5, 32570.	12.2	503

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55	Exosome-mediated Delivery of Hydrophobically Modified siRNA for Huntingtin mRNA Silencing. <i>Molecular Therapy</i> , 2016, 24, 1836-1847.	8.2	351
56	Docosahexaenoic Acid Conjugation Enhances Distribution and Safety of siRNA upon Local Administration in Mouse Brain. <i>Molecular Therapy - Nucleic Acids</i> , 2016, 5, e344.	5.1	67
57	A High-Throughput Method for Direct Detection of Therapeutic Oligonucleotide-Induced Gene Silencing <i>In Vivo</i> . <i>Nucleic Acid Therapeutics</i> , 2016, 26, 86-92.	3.6	38
58	Guanabenz (Wytensin <sup>®</sup> , $\Phi$ ) selectively enhances uptake and efficacy of hydrophobically modified siRNAs. <i>Nucleic Acids Research</i> , 2015, 43, 8664-8672.	14.5	39
59	Hydrophobically Modified siRNAs Silence Huntingtin mRNA in Primary Neurons and Mouse Brain. <i>Molecular Therapy - Nucleic Acids</i> , 2015, 4, e266.	5.1	115
60	Taking charge of siRNA delivery. <i>Nature Biotechnology</i> , 2014, 32, 1197-1198.	17.5	17
61	Novel Hydrophobically Modified Asymmetric RNAi Compounds (sd-rxRNA) Demonstrate Robust Efficacy in the Eye. <i>Journal of Ocular Pharmacology and Therapeutics</i> , 2013, 29, 855-864.	1.4	67
62	Identifying siRNA-Induced Off-Targets by Microarray Analysis. <i>Methods in Molecular Biology</i> , 2008, 442, 45-63.	0.9	28
63	Experimental validation of the importance of seed complement frequency to siRNA specificity. <i>Rna</i> , 2008, 14, 853-861.	3.5	122
64	A protocol for designing siRNAs with high functionality and specificity. <i>Nature Protocols</i> , 2007, 2, 2068-2078.	12.0	197
65	3' UTR seed matches, but not overall identity, are associated with RNAi off-targets. <i>Nature Methods</i> , 2006, 3, 199-204.	19.0	782
66	Position-specific chemical modification of siRNAs reduces "off-target" transcript silencing. <i>Rna</i> , 2006, 12, 1197-1205.	3.5	686
67	Rational siRNA design for RNA interference. <i>Nature Biotechnology</i> , 2004, 22, 326-330.	17.5	1,856
68	Functional siRNAs and miRNAs Exhibit Strand Bias. <i>Cell</i> , 2003, 115, 209-216.	28.9	2,320