

Hiroto Hatakeyama

List of Publications by Year in Descending Order

Source: <https://exaly.com/author-pdf/9047758/hiroto-hatakeyama-publications-by-year.pdf>

Version: 2024-04-28

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

53
papers

3,354
citations

33
h-index

57
g-index

61
ext. papers

3,731
ext. citations

8.9
avg, IF

5.24
L-index

#	Paper	IF	Citations
53	Silencing of VEGFR2 by RGD-Modified Lipid Nanoparticles Enhanced the Efficacy of Anti-PD-1 Antibody by Accelerating Vascular Normalization and Infiltration of T Cells in Tumors. <i>Cancers</i> , 2020 , 12,	6.6	5
52	Global Comparison of Changes in the Number of Test-Positive Cases and Deaths by Coronavirus Infection (COVID-19) in the World. <i>Journal of Clinical Medicine</i> , 2020 , 9,	5.1	9
51	Poor outcome with anti-programmed death-ligand 1 (PD-L1) antibody due to poor pharmacokinetic properties in PD-1/PD-L1 blockade-sensitive mouse models 2020 , 8,		9
50	Inhibition Synergistically Enhances the Effects of Magnetic Fluid Hyperthermia in Ovarian Cancer. <i>Molecular Cancer Therapeutics</i> , 2017 , 16, 966-976	6.1	31
49	Determinants of Intestinal Availability for P-glycoprotein Substrate Drugs Estimated by Extensive Simulation With Mathematical Absorption Models. <i>Journal of Pharmaceutical Sciences</i> , 2017 , 106, 2771-2779	3.9	3
48	Antitumor and Antiangiogenic Effects of Aspirin-PC in Ovarian Cancer. <i>Molecular Cancer Therapeutics</i> , 2016 , 15, 2894-2904	6.1	30
47	Role of CTGF in Sensitivity to Hyperthermia in Ovarian and Uterine Cancers. <i>Cell Reports</i> , 2016 , 17, 1621-1631	6.1	17
46	A miR-192-EGR1-HOXB9 regulatory network controls the angiogenic switch in cancer. <i>Nature Communications</i> , 2016 , 7, 11169	17.4	83
45	PEG dilemma- nucleic acids delivery to cancers by controlling biodistribution and intracellular trafficking. <i>Drug Delivery System</i> , 2016 , 31, 293-299	0	0
44	Relationship Between the Physicochemical Properties of Lipid Nanoparticles and the Quality of siRNA Delivery to Liver Cells. <i>Molecular Therapy</i> , 2016 , 24, 788-95	11.7	44
43	Assessment of In Vivo siRNA Delivery in Cancer Mouse Models. <i>Methods in Molecular Biology</i> , 2016 , 1402, 189-197	1.4	8
42	A lipid nanoparticle for the efficient delivery of siRNA to dendritic cells. <i>Journal of Controlled Release</i> , 2016 , 225, 183-91	11.7	73
41	Novel pH-sensitive multifunctional envelope-type nanodevice for siRNA-based treatments for chronic HBV infection. <i>Journal of Hepatology</i> , 2016 , 64, 547-55	13.4	45
40	Anti-tumor effect via passive anti-angiogenesis of PEGylated liposomes encapsulating doxorubicin in drug resistant tumors. <i>International Journal of Pharmaceutics</i> , 2016 , 509, 178-187	6.5	38
39	Molecular Tuning of a Vitamin E-Scaffold pH-Sensitive and Reductive Cleavable Lipid-like Material for Accelerated in Vivo Hepatic siRNA Delivery. <i>ACS Biomaterials Science and Engineering</i> , 2015 , 1, 834-844	5.5	32
38	Size-dependent specific targeting and efficient gene silencing in peritoneal macrophages using a pH-sensitive cationic liposomal siRNA carrier. <i>International Journal of Pharmaceutics</i> , 2015 , 495, 171-178	6.5	21
37	A neutral lipid envelope-type nanoparticle composed of a pH-activated and vitamin E-scaffold lipid-like material as a platform for a gene carrier targeting renal cell carcinoma. <i>Journal of Controlled Release</i> , 2015 , 200, 97-105	11.7	45

36	Advances in an active and passive targeting to tumor and adipose tissues. <i>Expert Opinion on Drug Delivery</i> , 2015 , 12, 41-52	8	34
35	Multifunctional Envelope-Type Nano Device: Evolution from Nonselective to Active Targeting System. <i>Bioconjugate Chemistry</i> , 2015 , 26, 1266-76	6.3	12
34	RNAi-mediated gene knockdown and anti-angiogenic therapy of RCCs using a cyclic RGD-modified liposomal-siRNA system. <i>Journal of Controlled Release</i> , 2014 , 173, 110-8	11.7	87
33	The systemic administration of an anti-miRNA oligonucleotide encapsulated pH-sensitive liposome results in reduced level of hepatic microRNA-122 in mice. <i>Journal of Controlled Release</i> , 2014 , 173, 43-50	11.7	56
32	2VOMe-phosphorodithioate-modified siRNAs show increased loading into the RISC complex and enhanced anti-tumour activity. <i>Nature Communications</i> , 2014 , 5, 3459	17.4	81
31	Improvement of doxorubicin efficacy using liposomal anti-polo-like kinase 1 siRNA in human renal cell carcinomas. <i>Molecular Pharmaceutics</i> , 2014 , 11, 2713-9	5.6	34
30	An apolipoprotein E modified liposomal nanoparticle: ligand dependent efficiency as a siRNA delivery carrier for mouse-derived brain endothelial cells. <i>International Journal of Pharmaceutics</i> , 2014 , 465, 77-82	6.5	33
29	An aptamer ligand based liposomal nanocarrier system that targets tumor endothelial cells. <i>Biomaterials</i> , 2014 , 35, 7110-20	15.6	50
28	Comparative study of the sensitivities of cancer cells to doxorubicin, and relationships between the effect of the drug-efflux pump P-gp. <i>Biological and Pharmaceutical Bulletin</i> , 2014 , 37, 1926-35	2.3	32
27	In vivo therapeutic potential of Dicer-hunting siRNAs targeting infectious hepatitis C virus. <i>Scientific Reports</i> , 2014 , 4, 4750	4.9	40
26	Application of apolipoprotein E-modified liposomal nanoparticles as a carrier for delivering DNA and nucleic acid in the brain. <i>International Journal of Nanomedicine</i> , 2014 , 9, 4267-76	7.3	18
25	Hepatic Monoacylglycerol O-acyltransferase 1 as a Promising Therapeutic Target for Steatosis, Obesity, and Type 2 Diabetes. <i>Molecular Therapy - Nucleic Acids</i> , 2014 , 3, e154	10.7	35
24	Cancer multidrug resistance: mechanisms involved and strategies for circumvention using a drug delivery system. <i>Archives of Pharmacal Research</i> , 2014 , 37, 4-15	6.1	126
23	The effect of liposomal size on the targeted delivery of doxorubicin to Integrin $\alpha\beta$ -expressing tumor endothelial cells. <i>Biomaterials</i> , 2013 , 34, 5617-27	15.6	77
22	Lipid envelope-type nanoparticle incorporating a multifunctional peptide for systemic siRNA delivery to the pulmonary endothelium. <i>ACS Nano</i> , 2013 , 7, 7534-41	16.7	65
21	A neutral envelope-type nanoparticle containing pH-responsive and SS-cleavable lipid-like material as a carrier for plasmid DNA. <i>Advanced Healthcare Materials</i> , 2013 , 2, 1120-5	10.1	50
20	Gene silencing via RNAi and siRNA quantification in tumor tissue using MEND, a liposomal siRNA delivery system. <i>Molecular Therapy</i> , 2013 , 21, 1195-203	11.7	97
19	In vitro optimization of 2VOMe-4Vthioribonucleoside-modified anti-microRNA oligonucleotides and its targeting delivery to mouse liver using a liposomal nanoparticle. <i>Nucleic Acids Research</i> , 2013 , 41, 10659-67	20.1	43

18	The polyethyleneglycol dilemma: advantage and disadvantage of PEGylation of liposomes for systemic genes and nucleic acids delivery to tumors. <i>Biological and Pharmaceutical Bulletin</i> , 2013 , 36, 892-9	2.3	301
17	Size-controlled, dual-ligand modified liposomes that target the tumor vasculature show promise for use in drug-resistant cancer therapy. <i>Journal of Controlled Release</i> , 2012 , 162, 225-32	11.7	80
16	Intracellular stability of 2'-OMe-4'-thioribonucleoside modified siRNA leads to long-term RNAi effect. <i>Nucleic Acids Research</i> , 2012 , 40, 5787-93	20.1	37
15	Delivery of Nucleic Acids and Gene Delivery 2011 , 411-444		6
14	A new peptide motif present in the protective antigen of anthrax toxin exerts its efficiency on the cellular uptake of liposomes and applications for a dual-ligand system. <i>International Journal of Pharmaceutics</i> , 2011 , 412, 106-14	6.5	14
13	Dual-ligand modification of PEGylated liposomes shows better cell selectivity and efficient gene delivery. <i>Journal of Controlled Release</i> , 2011 , 153, 141-8	11.7	162
12	A multifunctional envelope type nano device (MEND) for gene delivery to tumours based on the EPR effect: a strategy for overcoming the PEG dilemma. <i>Advanced Drug Delivery Reviews</i> , 2011 , 63, 152-60	18.5	506
11	Synthesis, structure, and biological activity of dumbbell-shaped nanocircular RNAs for RNA interference. <i>Bioconjugate Chemistry</i> , 2011 , 22, 2082-92	6.3	37
10	Systemic delivery of siRNA to tumors using a lipid nanoparticle containing a tumor-specific cleavable PEG-lipid. <i>Biomaterials</i> , 2011 , 32, 4306-16	15.6	168
9	Endosomal escape and the knockdown efficiency of liposomal-siRNA by the fusogenic peptide shGALA. <i>Biomaterials</i> , 2011 , 32, 5733-42	15.6	93
8	A DNA microarray-based analysis of the host response to a nonviral gene carrier: a strategy for improving the immune response. <i>Molecular Therapy</i> , 2011 , 19, 1487-98	11.7	20
7	siRNA delivery by multifunctional envelope-type nano device (MEND). <i>Drug Delivery System</i> , 2010 , 25, 590-597	0	1
6	A novel nonviral gene delivery system: multifunctional envelope-type nano device. <i>Advances in Biochemical Engineering/Biotechnology</i> , 2010 , 119, 197-230	1.7	4
5	Ornithine and tryptophan analogs as efficient polycations for short interference RNA delivery to tumor cells. <i>Biological and Pharmaceutical Bulletin</i> , 2010 , 33, 1246-9	2.3	6
4	Design of a dual-ligand system using a specific ligand and cell penetrating peptide, resulting in a synergistic effect on selectivity and cellular uptake. <i>International Journal of Pharmaceutics</i> , 2010 , 396, 143-8	6.5	50
3	A pH-sensitive fusogenic peptide facilitates endosomal escape and greatly enhances the gene silencing of siRNA-containing nanoparticles in vitro and in vivo. <i>Journal of Controlled Release</i> , 2009 , 139, 127-32	11.7	209
2	Efficient short interference RNA delivery to tumor cells using a combination of octaarginine, GALA and tumor-specific, cleavable polyethylene glycol system. <i>Biological and Pharmaceutical Bulletin</i> , 2009 , 32, 928-32	2.3	39
1	Factors governing the in vivo tissue uptake of transferrin-coupled polyethylene glycol liposomes in vivo. <i>International Journal of Pharmaceutics</i> , 2004 , 281, 25-33	6.5	140

