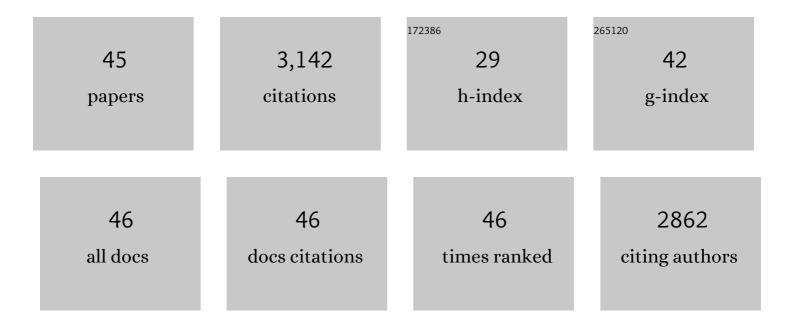
Thazah P Prakash

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Glucagon Like Peptide 1 Receptor Agonists for Targeted Delivery of Antisense Oligonucleotides to Pancreatic Beta Cell. Journal of the American Chemical Society, 2021, 143, 3416-3429.	6.6	39
2	Site-specific incorporation of 5â€2-methyl DNA enhances the therapeutic profile of gapmer ASOs. Nucleic Acids Research, 2021, 49, 1828-1839.	6.5	26
3	Site-specific Incorporation of 2′,5′-Linked Nucleic Acids Enhances Therapeutic Profile of Antisense Oligonucleotides. ACS Medicinal Chemistry Letters, 2021, 12, 922-927.	1.3	13
4	Overcoming the challenges of tissue delivery for oligonucleotide therapeutics. Trends in Pharmacological Sciences, 2021, 42, 588-604.	4.0	47
5	<i>Chop</i> / <i>Ddit3</i> depletion in β cells alleviates ER stress and corrects hepatic steatosis in mice. Science Translational Medicine, 2021, 13, .	5.8	38
6	Targeted Delivery of Antisense Oligonucleotides Using Neurotensin Peptides. Journal of Medicinal Chemistry, 2020, 63, 8471-8484.	2.9	27
7	Mechanisms of palmitic acid-conjugated antisense oligonucleotide distribution in mice. Nucleic Acids Research, 2020, 48, 4382-4395.	6.5	33
8	Fatty acid conjugation enhances potency of antisense oligonucleotides in muscle. Nucleic Acids Research, 2019, 47, 6029-6044.	6.5	93
9	Lipid Conjugates Enhance Endosomal Release of Antisense Oligonucleotides Into Cells. Nucleic Acid Therapeutics, 2019, 29, 245-255.	2.0	48
10	Conjugation of hydrophobic moieties enhances potency of antisense oligonucleotides in the muscle of rodents and non-human primates. Nucleic Acids Research, 2019, 47, 6045-6058.	6.5	48
11	S-Acyl-2-Thioethyl: A Convenient Base-Labile Protecting Group for the Synthesis of siRNAs Containing 5′-Vinylphosphonate. Molecules, 2019, 24, 225.	1.7	0
12	Receptor-Mediated Uptake of Phosphorothioate Antisense Oligonucleotides in Different Cell Types of the Liver. Nucleic Acid Therapeutics, 2018, 28, 119-127.	2.0	49
13	Evaluation of the effect of 2′-O-methyl, fluoro hexitol, bicyclo and Morpholino nucleic acid modifications on potency of GalNAc conjugated antisense oligonucleotides in mice. Bioorganic and Medicinal Chemistry Letters, 2018, 28, 3774-3779.	1.0	16
14	Targeted delivery of antisense oligonucleotides to pancreatic β-cells. Science Advances, 2018, 4, eaat3386.	4.7	132
15	Antisense oligonucleotides targeting mutant Ataxin-7 restore visual function in a mouse model of spinocerebellar ataxia type 7. Science Translational Medicine, 2018, 10, .	5.8	63
16	Subâ€organ Fractionation of Hepatic Cells after Antisense Oligonucleotide Treatment in Mice. FASEB Journal, 2018, 32, 760.11.	0.2	0
17	Asialoglycoprotein receptor 1 mediates productive uptake of N-acetylgalactosamine-conjugated and unconjugated phosphorothioate antisense oligonucleotides into liver hepatocytes. Nucleic Acids Research, 2017, 45, 12388-12400.	6.5	111
18	Characterizing the effect of GalNAc and phosphorothioate backbone on binding of antisense oligonucleotides to the asialoglycoprotein receptor. Nucleic Acids Research, 2017, 45, 2294-2306.	6.5	72

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19	A convenient synthesis of 5′-triantennary N-acetyl-galactosamine clusters based on nitromethanetrispropionic acid. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 2194-2197.	1.0	9
20	Elucidation of the Biotransformation Pathways of a Galnac3-conjugated Antisense Oligonucleotide in Rats and Monkeys. Molecular Therapy - Nucleic Acids, 2016, 5, e319.	2.3	46
21	Conjugation of mono and di-GalNAc sugars enhances the potency of antisense oligonucleotides via ASGR mediated delivery to hepatocytes. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 3690-3693.	1.0	36
22	Argonaute 2-dependent Regulation of Gene Expression by Single-stranded miRNA Mimics. Molecular Therapy, 2016, 24, 946-955.	3.7	51
23	Comprehensive Structure–Activity Relationship of Triantennary <i>N</i> -Acetylgalactosamine Conjugated Antisense Oligonucleotides for Targeted Delivery to Hepatocytes. Journal of Medicinal Chemistry, 2016, 59, 2718-2733.	2.9	107
24	Hsp90 protein interacts with phosphorothioate oligonucleotides containing hydrophobic 2′-modifications and enhances antisense activity. Nucleic Acids Research, 2016, 44, 3892-3907.	6.5	65
25	Efficient Synthesis and Biological Evaluation of 5′-GalNAc Conjugated Antisense Oligonucleotides. Bioconjugate Chemistry, 2015, 26, 1451-1455.	1.8	68
26	Effect of 2′- O -[2-[2-(N , N -dimethylamino)ethoxy]ethyl] modification on activity of gapmer antisense oligonucleotides containing 2′,4′-constrained 2′- O -ethyl nucleic acid. Bioorganic and Medicinal Chemistry Letters, 2015, 25, 1688-1691.	1.0	3
27	Solid-phase synthesis of 5′-triantennary N-acetylgalactosamine conjugated antisense oligonucleotides using phosphoramidite chemistry. Bioorganic and Medicinal Chemistry Letters, 2015, 25, 4127-4130.	1.0	21
28	Targeted delivery of antisense oligonucleotides to hepatocytes using triantennary <i>N</i> -acetyl galactosamine improves potency 10-fold in mice. Nucleic Acids Research, 2014, 42, 8796-8807.	6.5	465
29	TCP1 complex proteins interact with phosphorothioate oligonucleotides and can co-localize in oligonucleotide-induced nuclear bodies in mammalian cells. Nucleic Acids Research, 2014, 42, 7819-7832.	6.5	80
30	Lipid Nanoparticles Improve Activity of Single-Stranded siRNA and Gapmer Antisense Oligonucleotides in Animals. ACS Chemical Biology, 2013, 8, 1402-1406.	1.6	41
31	Single-Stranded siRNAs Activate RNAi in Animals. Cell, 2012, 150, 883-894.	13.5	239
32	An Overview of Sugarâ€Modified Oligonucleotides for Antisense Therapeutics. Chemistry and Biodiversity, 2011, 8, 1616-1641.	1.0	170
33	Synthesis and Biophysical Evaluation of 2′,4′-Constrained 2′ <i>O</i> -Methoxyethyl and 2′,4′-Constr 2′ <i>O</i> -Ethyl Nucleic Acid Analogues. Journal of Organic Chemistry, 2010, 75, 1569-1581.	rained	182
34	Short Antisense Oligonucleotides with Novel 2′â~'4′ Conformationaly Restricted Nucleoside Analogues Show Improved Potency without Increased Toxicity in Animals. Journal of Medicinal Chemistry, 2009, 52, 10-13.	2.9	236
35	RNA interference by 2′,5′-linked nucleic acid duplexes in mammalian cells. Bioorganic and Medicinal Chemistry Letters, 2006, 16, 3238-3240.	1.0	26
36	Synthesis and Evaluation ofS-Acyl-2-thioethyl Esters of Modified Nucleoside 5â€~-Monophosphates as Inhibitors of Hepatitis C Virus RNA Replication. Journal of Medicinal Chemistry, 2005, 48, 1199-1210.	2.9	34

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37	Positional Effect of Chemical Modifications on Short Interference RNA Activity in Mammalian Cells. Journal of Medicinal Chemistry, 2005, 48, 4247-4253.	2.9	259
38	2â€~-O-[2-(Guanidinium)ethyl]-Modified Oligonucleotides:  Stabilizing Effect on Duplex and Triplex Structuresâ€. Organic Letters, 2004, 6, 1971-1974.	2.4	55
39	2â€~-O-[2-(Methylthio)ethyl]-Modified Oligonucleotide: An Analogue of 2â€~-O-[2-(Methoxy)-ethyl]-Modified Oligonucleotide with Improved Protein Binding Properties and High Binding Affinity to Target RNAâ€. Biochemistry, 2002, 41, 11642-11648.	1.2	33
40	2′-DMAOE RNA: Emerging Oligonucleotides with Promising Antisense Properties. Nucleosides & Nucleotides, 1999, 18, 1381-1382.	0.5	8
41	Synthesis, Hybridization, and Nuclease Resistance Properties of 2′-O-Aminooxyethyl Modified Oligonucleotides. Nucleosides & Nucleotides, 1999, 18, 1419-1420.	0.5	3
42	N-(2-Cyanoethoxycarbonyloxy)succinimide:Â A New Reagent for Protection of Amino Groups in Oligonucleotides. Journal of Organic Chemistry, 1999, 64, 6468-6472.	1.7	38
43	A New Protecting Group Strategy for Amino Groups in Oligonucleotide Chemistry: CEOC Group. Nucleosides & Nucleotides, 1999, 18, 1199-1201.	0.5	3
44	Carbohydrate Modifications in Antisense Oligonucleotide Therapy: New Kids on the Block. Nucleosides & Nucleotides, 1999, 18, 1737-1746.	0.5	6
45	Targeted Delivery of Antisense Oligonucleotides Through Angiotensin Type 1 Receptor. Nucleic Acid Therapeutics, 0, , .	2.0	2