

# Maria Duca

## List of Publications by Year in descending order

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Version: 2024-02-01

40  
papers

1,099  
citations

471509

17  
h-index

414414

32  
g-index

43  
all docs

43  
docs citations

43  
times ranked

1143  
citing authors

#	ARTICLE	IF	CITATIONS
1	Development of 2-deoxystreptamine nucleobase conjugates for the inhibition of oncogenic miRNA production. <i>RSC Medicinal Chemistry</i> , 2022, 13, 311-319.	3.9	4
2	Differentiation of Cancer Stem Cells by Using Synthetic Small Molecules: Toward New Therapeutic Strategies against Therapy Resistance. <i>ChemMedChem</i> , 2021, 16, 14-29.	3.2	2
3	Design and Implementation of Synthetic RNA Binders for the Inhibition of miR-21 Biogenesis. <i>ACS Medicinal Chemistry Letters</i> , 2021, 12, 899-906.	2.8	17
4	Potent Tyrosinase Inhibitory Activity of Curcuminoid Analogues and Inhibition Kinetics Studies. <i>Cosmetics</i> , 2021, 8, 35.	3.3	8
5	The Chemical Biology Medicinal Chemistry Continuum: EFMC's Vision. <i>ChemBioChem</i> , 2021, 22, 2823-2825.	2.6	7
6	Unveiling RNA Binding Properties of Verapamil and Preparation of New Derivatives as Inhibitors of HIV TAR Interaction. <i>ChemPlusChem</i> , 2020, 85, 207-216.	2.8	7
7	Frontispiece: Aminoglycoside Conjugation for RNA Targeting: Antimicrobials and Beyond. <i>Chemistry - A European Journal</i> , 2020, 26, .	3.3	0
8	Aminoglycoside Conjugation for RNA Targeting: Antimicrobials and Beyond. <i>Chemistry - A European Journal</i> , 2020, 26, 12273-12309.	3.3	14
9	Inhibition of Patched Drug Efflux Increases Vemurafenib Effectiveness against Resistant BrafV600E Melanoma. <i>Cancers</i> , 2020, 12, 1500.	3.7	9
10	New Chemical Modalities Enabling Specific RNA Targeting and Degradation: Application to SARS-CoV-2 RNA. <i>ACS Central Science</i> , 2020, 6, 1647-1650.	11.3	1
11	New Chemical Modalities Enabling Specific RNA Targeting and Degradation: Application to SARS-CoV-2 RNA. <i>ACS Central Science</i> , 2020, 6, 1647-1650.	11.3	6
12	Functionalized C-nucleosides as remarkable RNA binders: targeting of prokaryotic ribosomal A-site RNA. <i>Chemical Communications</i> , 2019, 55, 10432-10435.	4.1	5
13	Synthetic small-molecule RNA ligands: future prospects as therapeutic agents. <i>MedChemComm</i> , 2019, 10, 1242-1255.	3.4	53
14	Modulation of oncogenic miRNA biogenesis using functionalized polyamines. <i>Scientific Reports</i> , 2018, 8, 1667.	3.3	39
15	Exploring Heterocycle-Spermine Conjugates as Modulators of Oncogenic microRNAs Biogenesis. <i>ACS Omega</i> , 2018, 3, 16500-16508.	3.5	13
16	Building of neomycin nucleobase amino acid conjugates for the inhibition of oncogenic miRNAs biogenesis. <i>Organic and Biomolecular Chemistry</i> , 2018, 16, 6262-6274.	2.8	27
17	Design of Multimodal Small Molecules Targeting miRNAs Biogenesis: Synthesis and In Vitro Evaluation. <i>Methods in Molecular Biology</i> , 2017, 1517, 137-154.	0.9	13
18	Small-molecule approaches toward the targeting of oncogenic miRNAs: roadmap for the discovery of RNA modulators. <i>Future Medicinal Chemistry</i> , 2016, 8, 803-816.	2.3	31

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19	Oncogenic MicroRNAs Biogenesis as a Drug Target: Structure–Activity Relationship Studies on New Aminoglycoside Conjugates. <i>Chemistry - A European Journal</i> , 2016, 22, 5350-5362.	3.3	41
20	Ribosome-targeting antibiotics as inhibitors of oncogenic microRNAs biogenesis: Old scaffolds for new perspectives in RNA targeting. <i>Bioorganic and Medicinal Chemistry</i> , 2015, 23, 5334-5344.	3.0	18
21	Artificial Nucleobase–Amino Acid Conjugates: A New Class of TAR RNA Binding Agents. <i>Chemistry - A European Journal</i> , 2014, 20, 2071-2079.	3.3	33
22	Targeting the Production of Oncogenic MicroRNAs with Multimodal Synthetic Small Molecules. <i>ACS Chemical Biology</i> , 2014, 9, 711-721.	3.4	99
23	Sequence-Specific Base Pair Mimics Are Efficient Topoisomerase IB Inhibitors. <i>Biochemistry</i> , 2012, 51, 43-51.	2.5	3
24	Contributions of the D-Ring to the Activity of Etoposide against Human Topoisomerase III $\beta$ : Potential Interactions with DNA in the Ternary Enzyme–Drug–DNA Complex. <i>Biochemistry</i> , 2011, 50, 5058-5066.	2.5	16
25	Structural Basis for the Exceptional Stability of Bisaminoacylated Nucleotides and Transfer RNAs. <i>Journal of the American Chemical Society</i> , 2011, 133, 11368-11377.	13.7	5
26	Targeting DNA base pair mismatch with artificial nucleobases. Advances and perspectives in triple helix strategy. <i>Organic and Biomolecular Chemistry</i> , 2011, 9, 326-336.	2.8	60
27	Design of novel RNA ligands that bind stem–bulge HIV-1 TAR RNA. <i>Chemical Communications</i> , 2010, 46, 6162.	4.1	33
28	Fluorescent labeling of human mesenchymal stem cells by thiophene fluorophores conjugated to a lipophilic carrier. <i>Chemical Communications</i> , 2010, 46, 7948.	4.1	23
29	Tandem Azide-Alkyne 1,3-Dipolar Cycloaddition/Electrophilic Addition: A Concise Three-Component Route to 4,5-Disubstituted Triazolyl-Nucleosides. <i>Synlett</i> , 2009, 2009, 2123-2126.	1.8	17
30	Aminoacylation of transfer RNAs with one and two amino acids. <i>Methods</i> , 2008, 44, 87-99.	3.8	9
31	Modeling the reactive properties of tandemly activated tRNAs. <i>Organic and Biomolecular Chemistry</i> , 2008, 6, 3292.	2.8	8
32	Targeting <i>MDR1</i> Gene: Synthesis and Cellular Study of Modified Daunomycin-Triplex-Forming Oligonucleotide Conjugates Able to Inhibit Gene Expression in Resistant Cell Lines. <i>Molecular Pharmacology</i> , 2008, 73, 1568-1577.	2.3	12
33	The triple helix: 50 years later, the outcome. <i>Nucleic Acids Research</i> , 2008, 36, 5123-5138.	14.5	302
34	Synthesis of bisaminoacylated pdCpAs and tandemly activated transfer RNAs. <i>Bioorganic and Medicinal Chemistry</i> , 2007, 15, 4629-4642.	3.0	7
35	Molecular basis of the targeting of topoisomerase II-mediated DNA cleavage by VP16 derivatives conjugated to triplex-forming oligonucleotides. <i>Nucleic Acids Research</i> , 2006, 34, 1900-1911.	14.5	27
36	Novel carbamate derivatives of 4- $\beta$ -amino-4-O-demethyl-4-desoxypodophyllotoxin as inhibitors of topoisomerase II: synthesis and biological evaluation. <i>Organic and Biomolecular Chemistry</i> , 2005, 3, 1074-1080.	2.8	11

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37	Triple Helix-Forming Oligonucleotides Conjugated to New Inhibitors of Topoisomerase II: Synthesis and Binding Properties. <i>Bioconjugate Chemistry</i> , 2005, 16, 873-884.	3.6	7
38	Synthesis and Biological Study of a New Series of 4-Demethylepipodophyllotoxin Derivatives. <i>Journal of Medicinal Chemistry</i> , 2005, 48, 593-603.	6.4	42
39	Selective Generation and Reactivity of 5'-Adenosinyl and 2'-Adenosinyl Radicals. <i>Chemistry - A European Journal</i> , 2004, 10, 1249-1255.	3.3	28
40	Synthesis and Biological Activity of Sulfonamide Derivatives of Epipodophyllotoxin. <i>Journal of Medicinal Chemistry</i> , 2004, 47, 2365-2374.	6.4	42