

Joanne T Douglas

List of Publications by Year in descending order

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Version: 2024-02-01

40
papers

377
citations

1683354

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1719596

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g-index

43
all docs

43
docs citations

43
times ranked

420
citing authors

#	ARTICLE	IF	CITATIONS
1	My experience of living with nonfluent/agrammatic variant primary progressive aphasia: Adaptations and strategies to improve quality of life. <i>Dementia</i> , 2021, 20, 936-951.	1.0	5
2	Development of a Palliative Care Approach for Primary Progressive Aphasia: My Experience as a Person Living With This Rare Disorder. <i>Journal of Palliative Care</i> , 2021, , 082585972110267.	0.4	2
3	Adaptation to Early-Stage Nonfluent/Agrammatic Variant Primary Progressive Aphasia. <i>American Journal of Alzheimer's Disease and Other Dementias</i> , 2014, 29, 289-292.	0.9	14
4	Ex Vivo Transfer of the Hoxc-8-interacting Domain of Smad1 by a Tropism-modified Adenoviral Vector Results in Efficient Bone Formation in a Rabbit Model of Spinal Fusion. <i>Journal of Spinal Disorders and Techniques</i> , 2010, 23, 63-73.	1.8	18
5	Adenoviral vectors for gene therapy. <i>Molecular Biotechnology</i> , 2007, 36, 71-80.	1.3	150
6	Adenovirus-Mediated Gene Delivery: An Overview. , 2004, 246, 3-14.		7
7	Adenovirus-Mediated Gene Delivery to Skeletal Muscle. , 2004, 246, 29-36.		7
8	Promoter Optimization and Artificial Promoters for Transcriptional Targeting in Gene Therapy. , 2003, , 481-503.		0
9	Cancer Gene Therapy. <i>Technology in Cancer Research and Treatment</i> , 2003, 2, 51-63.	0.8	11
10	Engineering Targeted Bacteriophage as Evolvable Vectors for Therapeutic Gene Delivery. , 2003, , 405-428.		0
11	Alternative Strategies for Targeted Delivery of Nucleic Acid-Liposome Complexes. , 2003, , 1-16.		0
12	Selection of Peptides on Phage. , 2003, , 547-579.		3
13	Single-Chain Fv Fragments from Phage Display Libraries. , 2003, , 597-620.		1
14	Strategies to Alter the Tropism of Adenoviral Vectors via Genetic Capsid Modification. , 2003, , 171-200.		1
15	Mechanisms of Retroviral Particle Maturation and Attachment. , 2003, , 241-251.		0
16	Targeting Retroviral Vectors Using Molecular Bridges. , 2003, , 253-266.		0
17	Genetic Engineering of Targeted Retroviral Vectors. , 2003, , 293-320.		5
18	Tumor/Tissue-Selective Promoters. , 2003, , 457-479.		0

#	ARTICLE	IF	CITATIONS
19	Physiological Targeting. , 2003, , 505-525.		0
20	Genetic Targeting of Retroviral Vectors. , 2003, , 267-291.		4
21	Generation of Safe, Targetable Sindbis Vectors That Have the Potential for Direct in vivo Gene Therapy. , 2003, , 353-375.		1
22	Targeted Gene Delivery via Lipidic Vectors. , 2003, , 17-32.		1
23	Immunoliposomes: A Targeted Delivery Tool for Cancer Treatment. , 2003, , 33-62.		3
24	Receptor-Directed Gene Delivery Using Molecular Conjugates. , 2003, , 63-86.		2
25	Conjugate-Based Targeting of Adeno-Associated Virus Vectors. , 2003, , 201-219.		0
26	Receptor Targeting of Adeno-Associated Virus Vectors. , 2003, , 221-239.		0
27	Targeting Bacteriophage Vectors. , 2003, , 429-455.		0
28	Retroviral Particle Display for Complex Glycosylated and Disulfide-Bonded Protein Domains. , 2003, , 621-634.		0
29	Cell Surface Display and Cytometric Screening for Protein Ligand Isolation and Engineering. , 2003, , 635-657.		0
30	Monitoring Gene Therapy by Positron Emission Tomography. , 2003, , 659-685.		3
31	Clostridium-Mediated Transfer of Therapeutic Proteins to Solid Tumors. , 2003, , 527-546.		1
32	Targeting of Adenoviral Gene Therapy Vectors: The Flexibility of Chemical and Molecular Conjugation. , 2003, , 123-141.		1
33	Targeting of Poliovirus Replicons to Neurons in the Central Nervous System. , 2003, , 337-352.		0
34	Pseudotyping of Adenoviral Vectors. , 2003, , 87-121.		0
35	Targeting Measles Virus Entry. , 2003, , 321-336.		0
36	Genetic Targeting of Adenoviral Vectors. , 2003, , 143-170.		3

#	ARTICLE	IF	CITATIONS
37	Redirecting the Tropism of HSV-1 for Gene Therapy Applications. , 2003, , 377-403.		0
38	Antibody Phage Display Libraries for Use in Therapeutic Gene Targeting. , 2003, , 581-596.		0
39	Targeted adenoviral vectors. Molecular Physics, 2002, 100, 3075-3091.	0.8	2
40	A system for the propagation of adenoviral vectors with genetically modified receptor specificities. Nature Biotechnology, 1999, 17, 470-475.	9.4	132