Sara MarcÃ³ Costa

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	AAV-mediated BMP7 gene therapy counteracts insulin resistance and obesity. Molecular Therapy - Methods and Clinical Development, 2022, 25, 190-204.	4.1	6
2	Treatment of skeletal and non-skeletal alterations of Mucopolysaccharidosis type IVA by AAV-mediated gene therapy. Nature Communications, 2021, 12, 5343.	12.8	15
3	Seven-year follow-up of durability and safety of AAV CNS gene therapy for a lysosomal storage disorder in a large animal. Molecular Therapy - Methods and Clinical Development, 2021, 23, 370-389.	4.1	16
4	<i>In Vivo</i> Gene Therapy for Mucopolysaccharidosis Type III (Sanfilippo Syndrome): A New Treatment Horizon. Human Gene Therapy, 2019, 30, 1211-1221.	2.7	25
5	FGF21 gene therapy as treatment for obesity and insulin resistance. EMBO Molecular Medicine, 2018, 10,	6.9	176
6	Disease correction by AAV-mediated gene therapy in a new mouse model of mucopolysaccharidosis type IIID. Human Molecular Genetics, 2017, 26, 1535-1551.	2.9	39
7	Progressive neurologic and somatic disease in a novel model of human Mucopolysaccharidosis type IIIC. DMM Disease Models and Mechanisms, 2016, 9, 999-1013.	2.4	14
8	348. Correction of CNS and Somatic Pathology by Intra-Cerebrospinal Fluid Gene Therapy for Mucopolysaccharidosis Type II. Molecular Therapy, 2016, 24, S139.	8.2	0
9	CNS-directed gene therapy for the treatment of neurologic and somatic mucopolysaccharidosis type II (Hunter syndrome). JCI Insight, 2016, 1, e86696.	5.0	56
10	Biochemical, histological and functional correction of mucopolysaccharidosis Type IIIB by intra-cerebrospinal fluid gene therapy. Human Molecular Genetics, 2015, 24, 2078-2095.	2.9	48
11	Whole body correction of mucopolysaccharidosis IIIA by intracerebrospinal fluid gene therapy. Journal of Clinical Investigation, 2013, 123, 3254-3271.	8.2	176
12	Liver Production of Sulfamidase Reverses Peripheral and Ameliorates CNS Pathology in Mucopolysaccharidosis IIIA Mice. Molecular Therapy, 2012, 20, 254-266.	8.2	51
13	Correction of Pathological Accumulation of Glycosaminoglycans in Central Nervous System and Peripheral Tissues of MPSIIIA Mice Through Systemic AAV9 Gene Transfer. Human Gene Therapy, 2012, 23, 1237-1246.	2.7	102