

# Hereditary transthyretin amyloidosis: a model of medicine

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Cryo-EM structure of a transthyretin-derived amyloid fibril from a patient with hereditary ATTR amyloidosis. <i>Nature Communications</i> , 2019, 10, 5008.	5.8	127
3	Expanding the spectrum of transthyretin amyloidosis. <i>Muscle and Nerve</i> , 2020, 61, 3-4.	1.0	2
4	Leading RNA Interference Therapeutics Part 1: Silencing Hereditary Transthyretin Amyloidosis, with a Focus on Patisiran. <i>Molecular Diagnosis and Therapy</i> , 2020, 24, 49-59.	1.6	35
5	Analysis of autonomic outcomes in APOLLO, a phase III trial of the RNAi therapeutic patisiran in patients with hereditary transthyretin-mediated amyloidosis. <i>Journal of Neurology</i> , 2020, 267, 703-712.	1.8	35
6	Kidney Transplantation in Systemic Amyloidosis. <i>Transplantation</i> , 2020, 104, 2035-2047.	0.5	6
7	Transthyretin amyloid cardiomyopathy: An uncharted territory awaiting discovery. <i>European Journal of Internal Medicine</i> , 2020, 82, 7-15.	1.0	32
8	Repurposing Benzbromarone for Familial Amyloid Polyneuropathy: A New Transthyretin Tetramer Stabilizer. <i>International Journal of Molecular Sciences</i> , 2020, 21, 7166.	1.8	15
9	Clinical 3-D Gait Assessment of Patients With Polyneuropathy Associated With Hereditary Transthyretin Amyloidosis. <i>Frontiers in Neurology</i> , 2020, 11, 605282.	1.1	6
10	Hereditary transthyretin amyloidosis overview. <i>Neurological Sciences</i> , 2022, 43, 595-604.	0.9	39
11	SERPINA1 modulates expression of amyloidogenic transthyretin. <i>Experimental Cell Research</i> , 2020, 395, 112217.	1.2	7
12	ATTRv amyloidosis Italian Registry: clinical and epidemiological data. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2020, 27, 259-265.	1.4	51
13	A Review of Patisiran (ONPATTRO®) for the Treatment of Polyneuropathy in People with Hereditary Transthyretin Amyloidosis. <i>Neurology and Therapy</i> , 2020, 9, 301-315.	1.4	96
14	Amyloid Typing by Mass Spectrometry in Clinical Practice: a Comprehensive Review of 16,175 Samples. <i>Mayo Clinic Proceedings</i> , 2020, 95, 1852-1864.	1.4	105
15	Hereditary transthyretin amyloidosis: current treatment. <i>Current Opinion in Neurology</i> , 2020, 33, 553-561.	1.8	11
16	Avoiding misdiagnosis: expert consensus recommendations for the suspicion and diagnosis of transthyretin amyloidosis for the general practitioner. <i>BMC Family Practice</i> , 2020, 21, 198.	2.9	60
17	Transthyretin Amyloidosis: Update on the Clinical Spectrum, Pathogenesis, and Disease-Modifying Therapies. <i>Neurology and Therapy</i> , 2020, 9, 317-333.	1.4	59
18	Conventional Molecular Dynamics and Metadynamics Simulation Studies of the Binding and Unbinding Mechanism of TTR Stabilizers AG10 and Tafamidis. <i>ACS Chemical Neuroscience</i> , 2020, 11, 3025-3035.	1.7	7
19	RNA Interference Nanotherapeutics for Treatment of Glioblastoma Multiforme. <i>Molecular Pharmaceutics</i> , 2020, 17, 4040-4066.	2.3	22

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21	Recommendations for pre-symptomatic genetic testing for hereditary transthyretin amyloidosis in the era of effective therapy: a multicenter Italian consensus. <i>Orphanet Journal of Rare Diseases</i> , 2020, 15, 348.	1.2	22
22	Advances in Treatment of ATTRv Amyloidosis: State of the Art and Future Prospects. <i>Brain Sciences</i> , 2020, 10, 952.	1.1	9
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27	Quantification of cardiac amyloid with [ <sup>18</sup> F]Flutemetamol in patients with V30M hereditary transthyretin amyloidosis. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2020, 27, 191-199.	1.4	14
28	Advances in the diagnosis of inherited neuromuscular diseases and implications for therapy development. <i>Lancet Neurology</i> , The, 2020, 19, 522-532.	4.9	36
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34	&lt;p&gt;Diagnosis and Treatment of Hereditary Transthyretin Amyloidosis (hATTR) Polyneuropathy: Current Perspectives on Improving Patient Care&lt;/p&gt;. <i>Therapeutics and Clinical Risk Management</i> , 2020, Volume 16, 109-123.	0.9	78
35	Structural Analysis of the Effect of a Dual-FLAG Tag on Transthyretin. <i>Biochemistry</i> , 2020, 59, 1013-1022.	1.2	7
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80	miRNA regulation of G protein-coupled receptor mediated angiogenic pathways in cancer. <i>Nucleus (India)</i> , 0, , 1.	0.9	4
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114	Proposing a minimal set of metrics and methods to predict probabilities of amyloidosis disease and onset age in individuals. <i>Aging</i> , 2020, 12, 22356-22369.	1.4	3
115	CNS Involvement in Hereditary Transthyretin Amyloidosis. <i>Neurology</i> , 2021, 97, 1111-1119.	1.5	30
116	Salivary Proteomics Identifies Transthyretin as a Biomarker of Early Dementia Conversion. <i>Journal of Alzheimer's Disease Reports</i> , 2022, 6, 31-41.	1.2	4
117	Unique Phenotypes With Corresponding Pathology in Late-Onset Hereditary Transthyretin Amyloidosis of A97S vs. V30M. <i>Frontiers in Aging Neuroscience</i> , 2021, 13, 786322.	1.7	8
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148	<sc>Long-term</sc> treatment effects of inotersen on <sc>health-related</sc> quality of life in patients with <sc>hATTR</sc> amyloidosis with polyneuropathy: Analysis of the <sc>open-label</sc> extension of the <sc>NEURO-ATTR</sc> trial. <i>Muscle and Nerve</i> , 2022, 66, 438-446.	1.0	3
149	A Study of Familial Amyloid Polyneuropathy Induced by the TTR Val30Leu Mutation in China. <i>European Neurology</i> , 0, , 1-6.	0.6	0
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