

PROGRESSIVE CARDIAC AMYLOIDOSIS FOLLOWING AMYLOID POLYNEUROPATHY

Transplantation

66, 229-233

DOI: [10.1097/00007890-199807270-00016](https://doi.org/10.1097/00007890-199807270-00016)

Citation Report

#	ARTICLE	IF	CITATIONS
1	Cardiac evaluation of patients with familial amyloidotic polyneuropathy proposed for liver transplantation. <i>Transplantation Proceedings</i> , 1999, 31, 2372.	0.3	3
2	Musical Hallucinations After Living-Donor Liver Transplantation. <i>Psychosomatics</i> , 1999, 40, 530-531.	2.5	9
3	Liver transplantation for hereditary transthyretin amyloidosis. <i>Liver Transplantation</i> , 2000, 6, 263-276.	1.3	123
4	Sequential (domino) transplantation of the liver in a transthyretin-50 familial amyloid polyneuropathy. <i>Langenbeck's Archives of Surgery</i> , 2000, 385, 21-26.	0.8	14
5	Heart failure caused by a novel amyloidogenic mutation of the transthyretin gene: ATTR Ala45Ser. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2000, 7, 137-140.	1.4	17
6	Cardiac Amyloid in Patients with Familial Amyloid Polyneuropathy Consists of Abundant Wild-Type Transthyretin. <i>Biochemical and Biophysical Research Communications</i> , 2000, 274, 702-706.	1.0	155
7	An Italian family with Ala-47 transthyretin mutation associated with cardiomyopathy and polyneuropathy. <i>Neuromuscular Disorders</i> , 2000, 10, 52-55.	0.3	6
8	Remission of proteinuria following liver transplantation for familial amyloid polyneuropathy TTR Met30. <i>Transplantation Proceedings</i> , 2000, 32, 2664-2666.	0.3	14
9	Suppression of transthyretin expression by ribozymes: a possible therapy for familial amyloidotic polyneuropathy. <i>Journal of the Neurological Sciences</i> , 2001, 183, 79-84.	0.3	34
10	Cautions after Liver Transplantation for Familial Amyloidotic Polyneuropathy. <i>Internal Medicine</i> , 2001, 40, 193-194.	0.3	0
11	Combined heart and liver transplantation for familial amyloidosis. <i>Internal Medicine Journal</i> , 2001, 31, 66-67.	0.5	19
12	Transthyretin mutations in hyperthyroxinemia and amyloid diseases. <i>Human Mutation</i> , 2001, 17, 493-503.	1.1	200
13	Long term results of heart transplantation in patients with amyloid heart disease. <i>British Heart Journal</i> , 2001, 85, 202-207.	2.2	91
14	Transthyretin-associated neuropathic amyloidosis. <i>Neurology</i> , 2001, 56, 431-435.	1.5	113
15	Electron and immuno-electron microscopy of abdominal fat identifies and characterizes amyloid fibrils in suspected cardiac amyloidosis. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2002, 9, 108-114.	1.4	141
16	Familial transthyretin-type amyloid polyneuropathy in Japan. <i>Neurology</i> , 2002, 58, 1001-1007.	1.5	204
17	Induction of Protein Conformational Change in Mouse Senile Amyloidosis. <i>Journal of Biological Chemistry</i> , 2002, 277, 33164-33169.	1.6	48
18	Transthyretin mutation (TTRGly47Ala) associated with familial amyloid polyneuropathy in a French family. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2002, 9, 272-275.	1.4	5

#	ARTICLE	IF	CITATIONS
19	Long-term follow-up of survival of liver transplant recipients with familial amyloid polyneuropathy (Portuguese type). <i>Liver Transplantation</i> , 2002, 8, 787-794.	1.3	76
20	Combined heart and liver transplantation for familial amyloidotic neuropathy: Considerations from the hepatic point of view. <i>Liver Transplantation</i> , 2003, 9, 986-992.	1.3	46
21	Gastrointestinal amyloidosis: Approach to treatment. <i>Current Treatment Options in Gastroenterology</i> , 2003, 6, 17-25.	0.3	16
22	Combined heart and liver transplantation for familial amyloidotic polyneuropathy. <i>Journal of Thoracic and Cardiovascular Surgery</i> , 2003, 125, 1165-1166.	0.4	33
23	Contribution of wild-type transthyretin to hereditary peripheral nerve amyloid. <i>Muscle and Nerve</i> , 2003, 28, 438-442.	1.0	48
24	Outcome of liver transplantation for familial amyloidotic polyneuropathy. <i>Liver Transplantation</i> , 2003, 9, 1273-1280.	1.3	46
25	Hereditary transthyretin amyloidosis from a Scandinavian perspective. <i>Journal of Internal Medicine</i> , 2003, 254, 225-235.	2.7	91
26	Caso clnico Polineuropata en un paciente con ingesta excesiva de alcohol. <i>Medicine</i> , 2003, 8, 5417-5418.	0.0	0
27	Liver transplantation as treatment for neurological disorders. <i>Expert Review of Neurotherapeutics</i> , 2003, 3, 547-555.	1.4	6
28	Clinical and pathological studies of cardiac amyloidosis in transthyretin type familial amyloid polyneuropathy. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2003, 10, 229-239.	1.4	41
29	Orthotopic liver transplantation for hereditary fibrinogen amyloidosis. <i>Transplantation</i> , 2003, 75, 560-561.	0.5	53
30	Clinical and pathological findings of non-Val30Met TTR type familial amyloid polyneuropathy in Japan. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2003, 10, 39-47.	1.4	12
31	Report from the Familial Amyloidotic Polyneuropathy World Transplant Registry (FAPWTR) and the Domino Liver Transplant Registry (DLTR). <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2003, 10, 67-76.	1.4	138
32	Impact of liver transplantation on familial amyloidotic polyneuropathy (FAP) patients' symptoms and complications. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2003, 10, 77-83.	1.4	37
33	Deposition and passage of transthyretin through the blood-nerve barrier in recipients of familial amyloid polyneuropathy livers. <i>Laboratory Investigation</i> , 2004, 84, 865-873.	1.7	64
34	Therapy for immunoglobulin light chain amyloidosis: the new and the old. <i>Blood Reviews</i> , 2004, 18, 17-37.	2.8	59
36	Metabolic storage diseases: amyloidosis. <i>Clinics in Liver Disease</i> , 2004, 8, 915-930.	1.0	17
37	Cardiomyopathies restrictives. <i>EMC - Cardiologie-Angiologie</i> , 2004, 1, 256-270.	0.9	0

#	ARTICLE	IF	CITATIONS
38	Combined heart and liver transplantation in four adults with familial amyloidosis: experience of a single center. <i>Transplantation Proceedings</i> , 2004, 36, 645-647.	0.3	54
39	Familial amyloid polyneuropathy and liver transplantation. <i>Journal of Hepatology</i> , 2004, 41, 188-194.	1.8	39
40	Cardiac transplantation for amyloid heart disease: The United Kingdom experience. <i>Journal of Heart and Lung Transplantation</i> , 2004, 23, 1142-1153.	0.3	165
41	Liver transplantation in transthyretin-related familial amyloid polyneuropathy. <i>Current Opinion in Neurology</i> , 2004, 17, 615-620.	1.8	98
42	Liver Transplantation Does Not Prevent the Development of Life-Threatening Arrhythmia in Familial Amyloidotic Polyneuropathy, Portuguese-Type (ATTR Val30Met) Patients. <i>Transplantation</i> , 2004, 78, 112-116.	0.5	76
43	The systemic amyloidoses. <i>Current Opinion in Rheumatology</i> , 2004, 16, 67-75.	2.0	118
44	Cardiac Amyloidosis: Heterogenous Pathogenic Backgrounds. <i>Internal Medicine</i> , 2004, 43, 1107-1114.	0.3	26
45	Ten Years of Experience with Liver Transplantation for Familial Amyloid Polyneuropathy in Japan: Outcomes of Living Donor Liver Transplantations. <i>Internal Medicine</i> , 2005, 44, 1151-1156.	0.3	45
46	Cardiovascular Magnetic Resonance in Cardiac Amyloidosis. <i>Circulation</i> , 2005, 111, 186-193.	1.6	863
47	Early liver transplantation improves familial amyloidotic polyneuropathy patients' survival. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2005, 12, 233-238.	1.4	62
48	A Swedish family with the rare Phe33Leu transthyretin mutation. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2005, 12, 189-192.	1.4	8
49	Diagnosis and Management of the Cardiac Amyloidoses. <i>Circulation</i> , 2005, 112, 2047-2060.	1.6	696
50	Liver Transplantation for Familial Amyloid Polyneuropathy Non-VAL30MET Variants: Are Cardiac Complications Influenced by Prophylactic Pacing and Immunosuppressive Weaning?. <i>Transplantation Proceedings</i> , 2005, 37, 2214-2220.	0.3	7
51	Cardiomyopathy in Swedish patients with the Gly53Glu and His88Arg transthyretin variants. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2005, 12, 184-188.	1.4	18
52	Amyloidosis. <i>Annual Review of Medicine</i> , 2006, 57, 223-241.	5.0	557
54	Effect of tacrolimus and partial hepatectomy on transthyretin metabolism in rats. <i>Transplant International</i> , 2006, 19, 233-238.	0.8	5
55	Successful Hepatorenal Transplantation in Hereditary Amyloidosis Caused by a Frame-Shift Mutation in Fibrinogen A α -Chain Gene. <i>American Journal of Transplantation</i> , 2006, 6, 632-635.	2.6	32
56	Ventricular late potentials in familial amyloidotic polyneuropathy. <i>Journal of Electrocardiology</i> , 2006, 39, 57-62.	0.4	12

#	ARTICLE	IF	CITATIONS
57	Severe congestive heart failure with cardiac liver cirrhosis 10 years after orthotopic liver transplantation for familial amyloidotic polyneuropathy. <i>Pathology International</i> , 2006, 56, 408-412.	0.6	14
58	Impact of Liver Transplantation on Cardiac Autonomic Denervation in Familial Amyloid Polyneuropathy. <i>Medicine (United States)</i> , 2006, 85, 229-238.	0.4	63
59	Aging and transthyretin-related amyloidosis: Pathologic examinations in pulmonary amyloidosis. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2006, 13, 24-30.	1.4	31
60	Phenotypic and genotypic heterogeneity in transthyretin-related cardiac amyloidosis: Towards tailoring of therapeutic strategies?. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2006, 13, 143-153.	1.4	57
61	Myocardial hypertrophy and function are related to age at onset in familial amyloidotic polyneuropathy. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2006, 13, 154-159.	1.4	50
62	Chapter 21 Amyloidosis. <i>Handbook of Systemic Autoimmune Diseases</i> , 2007, 7, 383-396.	0.1	1
63	A new transthyretin variant (Glu61Gly) associated with cardiomyopathy. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2007, 14, 65-71.	1.4	9
64	Progression of cardiac amyloid deposition in hereditary transthyretin amyloidosis patients after liver transplantation. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2007, 14, 277-282.	1.4	110
65	Combined heart and liver transplantation for familial amyloidotic neuropathy. <i>European Journal of Cardio-thoracic Surgery</i> , 2007, 32, 180-182.	0.6	49
66	Cardiac Transplantation Using Extended-Donor Criteria Organs for Systemic Amyloidosis Complicated by Heart Failure. <i>Transplantation</i> , 2007, 83, 539-545.	0.5	73
67	Combined Cardiac and Liver Transplantation for the Treatment of Familial Amyloidosis. <i>Baylor University Medical Center Proceedings</i> , 2007, 20, 146-148.	0.2	2
68	The molecular biology and clinical features of amyloid neuropathy. <i>Muscle and Nerve</i> , 2007, 36, 411-423.	1.0	449
69	Familial amyloidosis in a large Spanish kindred resulting from a D38V mutation in the transthyretin gene. <i>European Journal of Clinical Investigation</i> , 2007, 37, 673-678.	1.7	1
70	Progressive Wild-Type Transthyretin Deposition after Liver Transplantation Preferentially Occurs onto Myocardium in FAP Patients. <i>American Journal of Transplantation</i> , 2007, 7, 235-242.	2.6	110
71	An Unusual Case of Cardiac Amyloidosis. <i>Journal of General Internal Medicine</i> , 2007, 22, 1047-1052.	1.3	6
72	Amyloid fibril composition is related to the phenotype of hereditary transthyretin V30M amyloidosis. <i>Journal of Pathology</i> , 2008, 216, 253-261.	2.1	171
73	Marked regression of abdominal fat amyloid in patients with familial amyloid polyneuropathy during long-term follow-up after liver transplantation. <i>Liver Transplantation</i> , 2008, 14, 563-570.	1.3	62
74	Do troponin and Bâ€natriuretic peptide detect cardiomyopathy in transthyretin amyloidosis?. <i>Journal of Internal Medicine</i> , 2008, 263, 294-301.	2.7	53

#	ARTICLE	IF	CITATIONS
75	Accelerating restrictive cardiomyopathy after liver transplantation in a patient with familial amyloidotic polyneuropathy: a case report. <i>Journal of Medical Case Reports</i> , 2008, 2, 35.	0.4	5
76	Current State of Combined Heart and Liver Transplantation in the United States. <i>Journal of Heart and Lung Transplantation</i> , 2008, 27, 753-759.	0.3	83
77	A Case of Polyneuropathy and Proteinuria. <i>Clinical Journal of the American Society of Nephrology: CJASN</i> , 2008, 3, 624-636.	2.2	8
78	Evaluation of Myocardial Changes in Familial Amyloid Polyneuropathy after Liver Transplantation. <i>Internal Medicine</i> , 2008, 47, 2133-2137.	0.3	29
79	Tissue Doppler and strain imaging: a new tool for early detection of cardiac amyloidosis. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2009, 16, 63-70.	1.4	50
80	Chain reaction of amyloid fibril formation with induction of basement membrane in familial amyloidotic polyneuropathy. <i>Journal of Pathology</i> , 2009, 219, 481-490.	2.1	37
81	Proportion between wild-type and mutant protein in truncated compared to full-length ATTR: An analysis on transplanted transthyretin T60A amyloidosis patients. <i>Biochemical and Biophysical Research Communications</i> , 2009, 379, 846-850.	1.0	24
82	Combined Heart and Liver Transplantation: A Single-Center Experience. <i>Transplantation</i> , 2009, 88, 219-225.	0.5	118
83	Amyloid Heart Disease. <i>Progress in Cardiovascular Diseases</i> , 2010, 52, 347-361.	1.6	202
85	Poor outcome after liver transplantation for transthyretin amyloid neuropathy in a family with an Ala36Pro transthyretin mutation: Case Report. <i>Liver Transplantation</i> , 2010, 16, NA-NA.	1.3	4
86	Effect of recipient-derived cells on the progression of familial amyloidotic polyneuropathy after liver transplantation: a retrospective study. <i>Annals of Clinical Biochemistry</i> , 2010, 47, 529-534.	0.8	1
87	Progression of transthyretin amyloid neuropathy after liver transplantation. <i>Neurology</i> , 2010, 75, 324-327.	1.5	136
89	Heart complications in familial transthyretin amyloidosis: impact of age and gender. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2010, 17, 63-68.	1.4	72
90	Prognostic value of pre-transplant cardiomyopathy in Swedish liver transplanted patients for familial amyloidotic polyneuropathy. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2011, 18, 171-173.	1.4	12
91	The diflunisal trial: update on study drug tolerance and disease progression. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2011, 18, 196-197.	1.4	20
92	Cardiac Amyloidosis: A Practical Approach to Diagnosis and Management. <i>American Journal of Medicine</i> , 2011, 124, 1006-1015.	0.6	97
93	Wild-type transthyretin significantly contributes to the formation of amyloid fibrils in familial amyloid polyneuropathy patients with amyloidogenic transthyretin Val30Met. <i>Human Pathology</i> , 2011, 42, 236-243.	1.1	20
94	Pre operative cardio pulmonary assessment of the liver transplant candidate. <i>Annals of Hepatology</i> , 2011, 10, 421-433.	0.6	14

#	ARTICLE	IF	CITATIONS
95	Successful Heart and Liver Transplantation in a Swiss Patient With Glu89Lys Transthyretin Amyloidosis. <i>Transplantation</i> , 2011, 91, e40-e42.	0.5	4
97	Manifestations of transthyretin-related familial amyloidotic polyneuropathy: Long-term follow-up of Japanese patients after liver transplantation. <i>Surgery Today</i> , 2011, 41, 1211-1218.	0.7	41
98	Variation in amount of wild-type transthyretin in different fibril and tissue types in ATTR amyloidosis. <i>Journal of Molecular Medicine</i> , 2011, 89, 171-180.	1.7	74
99	Development of cardiomyopathy after liver transplantation in Swedish hereditary transthyretin amyloidosis (ATTR) patients. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2011, 18, 200-205.	1.4	57
100	Long-term data from the Familial Amyloidotic Polyneuropathy World Transplant Registry (FAPWTR). <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2011, 18, 193-195.	1.4	81
102	New insights into the clinical evaluation of hereditary transthyretin amyloidosis patients: a single center's experience. <i>Degenerative Neurological and Neuromuscular Disease</i> , 2012, 2, 93.	0.7	1
103	Amyloid Fibril Composition as a Predictor of Development of Cardiomyopathy After Liver Transplantation for Hereditary Transthyretin Amyloidosis. <i>Transplantation</i> , 2012, 93, 1017-1023.	0.5	52
104	Neuropatie amiloidi familiari. <i>EMC - Neurologia</i> , 2012, 12, 1-12.	0.0	0
105	Cardiac phenotype and clinical outcome of familial amyloid polyneuropathy associated with transthyretin alanine 60 variant. <i>European Heart Journal</i> , 2012, 33, 1120-1127.	1.0	143
106	The Diflunisal Trial: Study accrual and drug tolerance. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2012, 19, 37-38.	1.4	39
107	Amyloid diseases of the heart: current and future therapies. <i>QJM - Monthly Journal of the Association of Physicians</i> , 2012, 105, 617-631.	0.2	22
108	Transthyretin familial amyloid polyneuropathy. <i>Handbook of Clinical Neurology / Edited By P J Vinken and G W Bruyn</i> , 2013, 115, 643-658.	1.0	42
109	Systemic amyloidoses: What an internist should know. <i>European Journal of Internal Medicine</i> , 2013, 24, 729-739.	1.0	31
110	Long-term effects of tafamidis for the treatment of transthyretin familial amyloid polyneuropathy. <i>Journal of Neurology</i> , 2013, 260, 2802-2814.	1.8	284
111	Effects of Tafamidis on Transthyretin Stabilization and Clinical Outcomes in Patients with Non-Val30Met Transthyretin Amyloidosis. <i>Journal of Cardiovascular Translational Research</i> , 2013, 6, 1011-1020.	1.1	122
112	Liver transplantation and transthyretin amyloidosis. <i>Muscle and Nerve</i> , 2013, 47, 157-162.	1.0	92
113	Cardiac amyloidosis: A comprehensive review. <i>Cor Et Vasa</i> , 2013, 55, e60-e75.	0.1	38
114	Treatment of End Stage Heart Failure Related to Cardiac Amyloidosis. , 2013, , .		0

#	ARTICLE	IF	CITATIONS
115	Steady turnover of amyloid fibril proteins in gastric mucosa after liver transplantation in familial amyloid polyneuropathy. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2013, 20, 156-163.	1.4	10
116	RETINAL MICROANGIOPATHY AS AN INITIAL MANIFESTATION OF FAMILIAL AMYLOID CARDIOMYOPATHY ASSOCIATED WITH TRANSTHYRETIN E89K MUTATION. <i>Retinal Cases and Brief Reports</i> , 2013, 7, 271-275.	0.3	6
117	Clinical diagnosis and typing of systemic amyloidosis in subcutaneous fat aspirates by mass spectrometry-based proteomics. <i>Haematologica</i> , 2014, 99, 1239-1247.	1.7	140
118	Recent progress in the understanding and treatment of transthyretin amyloidosis. <i>Journal of Clinical Pharmacy and Therapeutics</i> , 2014, 39, 225-233.	0.7	57
119	Efficacy and safety of patisiran for familial amyloidotic polyneuropathy: a phase II multi-dose study. <i>Orphanet Journal of Rare Diseases</i> , 2015, 10, 109.	1.2	246
120	Outcome in patients treated with isolated liver transplantation for familial transthyretin amyloidosis to prevent cardiomyopathy. <i>Clinical Transplantation</i> , 2015, 29, 1098-1104.	0.8	6
121	Liver Transplantation for Hereditary Transthyretin Amyloidosis. <i>Transplantation</i> , 2015, 99, 1847-1854.	0.5	257
122	The transthyretin amyloidoses: advances in therapy. <i>Postgraduate Medical Journal</i> , 2015, 91, 439-448.	0.9	33
123	Cardiac amyloidosis: where are we today?. <i>Journal of Internal Medicine</i> , 2015, 278, 126-144.	2.7	61
124	Evolving landscape in the management of transthyretin amyloidosis. <i>Annals of Medicine</i> , 2015, 47, 625-638.	1.5	181
125	Cardiac Findings and Events Observed in an Open-Label Clinical Trial of Tafamidis in Patients with non-Val30Met and non-Val122Ile Hereditary Transthyretin Amyloidosis. <i>Journal of Cardiovascular Translational Research</i> , 2015, 8, 117-127.	1.1	61
126	Bifunctional crosslinking ligands for transthyretin. <i>Open Biology</i> , 2015, 5, 150105.	1.5	2
127	Novel strategies for the diagnosis and treatment of cardiac amyloidosis. <i>Expert Review of Cardiovascular Therapy</i> , 2015, 13, 1195-1211.	0.6	29
128	Current and future treatment of amyloid diseases. <i>Journal of Internal Medicine</i> , 2016, 280, 177-202.	2.7	73
129	Transthyretin amyloid neuropathy has earlier neural involvement but better prognosis than primary amyloid counterpart: an answer to the paradox?. <i>Annals of Neurology</i> , 2016, 80, 401-411.	2.8	17
130	^{99m} Tc-DPD uptake reflects amyloid fibril composition in hereditary transthyretin amyloidosis. <i>Upsala Journal of Medical Sciences</i> , 2016, 121, 17-24.	0.4	82
131	Survival After Transplantation in Patients With Mutations Other Than Val30Met. <i>Transplantation</i> , 2016, 100, 373-381.	0.5	65
132	Systemic amyloidosis. <i>Lancet, The</i> , 2016, 387, 2641-2654.	6.3	703

#	ARTICLE	IF	CITATIONS
133	One mutation, two distinct disease variants: unravelling the impact of transthyretin amyloid fibril composition. <i>Journal of Internal Medicine</i> , 2017, 281, 337-347.	2.7	94
134	Transthyretin amyloidosis: an under-recognized neuropathy and cardiomyopathy. <i>Clinical Science</i> , 2017, 131, 395-409.	1.8	66
135	Progressão da desnervação simpática cardíaca avaliada por cintigrafia com MIBG ¹²³ na polineuropatia amiloideótica familiar e o impacto da transplantação hepática. <i>Revista Portuguesa De Cardiologia</i> , 2017, 36, 333-340.	0.2	15
136	Progression of myocardial sympathetic denervation assessed by 123I-MIBG imaging in familial amyloid polyneuropathy and the effect of liver transplantation. <i>Revista Portuguesa De Cardiologia (English)</i> Tj ETQq1 1 0.784314 rgBI1/Overlo	0.3	14
137	Acquired transthyretin amyloidosis after domino liver transplant: Phenotypic correlation, implication of liver retransplantation. <i>Journal of the Neurological Sciences</i> , 2017, 379, 192-197.	0.3	9
138	Cardiac Amyloidosis: An Updated Review With Emphasis on Diagnosis and Future Directions. <i>Current Problems in Cardiology</i> , 2018, 43, 10-34.	1.1	74
139	Brazilian consensus for diagnosis, management and treatment of transthyretin familial amyloid polyneuropathy. <i>Arquivos De Neuro-Psiquiatria</i> , 2018, 76, 609-621.	0.3	16
140	Amyloid seeding of transthyretin by ex vivo cardiac fibrils and its inhibition. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018, 115, E6741-E6750.	3.3	66
141	History and Current Status of Cardiac Anesthesia in Singapore. <i>Journal of Cardiothoracic and Vascular Anesthesia</i> , 2019, 33, 3394-3401.	0.6	5
142	Diagnostic and Treatment Approaches Involving Transthyretin in Amyloidogenic Diseases. <i>International Journal of Molecular Sciences</i> , 2019, 20, 2982.	1.8	20
143	Systemic Amyloidosis. , 2019, , 267-289.		1
144	A molecular mechanism for transthyretin amyloidogenesis. <i>Nature Communications</i> , 2019, 10, 925.	5.8	92
145	Tafamidis for the treatment of transthyretin amyloidosis. <i>Future Cardiology</i> , 2019, 15, 53-61.	0.5	3
147	Pearls and Oysters: Number, weaker, and dizzier due to transthyretin amyloidosis after two liver transplants. <i>Neurology</i> , 2020, 96, 10.1212/WNL.0000000000011289.	1.5	1
148	Drug Discovery and Development in Rare Diseases: Taking a Closer Look at the Tafamidis Story. <i>Drug Design, Development and Therapy</i> , 2021, Volume 15, 1225-1243.	2.0	29
149	Amyloid Heart Disease. , 2010, , 107-128.		5
150	Liver Transplantation for Transthyretin Amyloidosis. , 2009, , 239-260.		2
151	Hereditary Amyloid Neuropathy. , 2005, , 1921-1935.		5

#	ARTICLE	IF	CITATIONS
152	Disorders of Peripheral Nerves. , 2008, , 2249-2355.		18
153	Disorders of Peripheral Nerves. , 2012, , 1915-2015.		11
154	Progression of cardiomyopathy after liver transplantation in patients with familial amyloidotic polyneuropathy, portuguese type1. Transplantation, 2002, 73, 745-751.	0.5	139
155	Hereditary Amyloidosis with Recurrent Lung Infiltrates. American Journal of Case Reports, 2016, 17, 874-879.	0.3	3
156	Contemporary Treatment of Amyloid Heart Disease. Current Pharmaceutical Design, 2014, 21, 491-506.	0.9	7
158	Cardiovascular Involvement in Transthyretin Cardiac Amyloidosis. Heart Failure Clinics, 2022, 18, 73-87.	1.0	12
159	Immunoglobulin Light Chain Amyloidosis (Primary Amyloidosis, AL). , 2004, , 157-195.		1
160	Continued Deposition of Wild-Type Transthyretin in Cardiac Amyloid after Liver Transplantation. , 2004, , 354-356.		1
161	Effect of Tacrolimus and Cyclosporine on Transthyretin (Ttr) Metabolism in Rats. , 2007, , 214-216.		0
163	ATTR: Diagnosis, Prognosis, and Treatment. , 2010, , 191-204.		3
164	Hereditary Neuromuscular Diseases and Cardiac Involvement. , 2011, , 385-400.		0
165	Cardiac and Multi-Organ Transplantation in Patients with Amyloidosis. , 0, , .		0
166	Clinical and Pathologic Issues in Patients with Amyloidosis: Practical Comments Regarding Diagnosis, Therapy, and Solid Organ Transplantation. , 2012, , 377-391.		0
167	Peripheral Nerve Amyloidosis. , 2012, , 361-374.		1
169	Clinical and Pathologic Issues in Patients with Amyloidosis: Summary and Practical Comments Regarding Diagnosis, Therapy, and Solid Organ Transplantation. Current Clinical Pathology, 2015, , 489-505.	0.0	1
170	Peripheral Nerve Amyloidosis. Current Clinical Pathology, 2015, , 437-450.	0.0	1
171	Hereditary Neuromuscular Diseases and Cardiac Involvement. , 2016, , 373-388.		0
173	CRISPR-Cas9 In Vivo Gene Editing for Transthyretin Amyloidosis. New England Journal of Medicine, 2021, 385, 1721-1723.	13.9	13

#	ARTICLE	IF	CITATIONS
174	Amyloid heart disease. New frontiers and insights in pathophysiology, diagnosis, and management. Texas Heart Institute Journal, 2005, 32, 178-84.	0.1	73
175	Hipertrofia Ventricular Esquerda: Um FenÃ³tipo, Duas HipÃ³teses, TrÃªs LiÃ§Ãµes. Arquivos Brasileiros De Cardiologia, 2021, 117, 1056-1059.	0.3	0
176	Development of amyloid beta gold nanorod aggregates as optoacoustic probes. PLoS ONE, 2022, 17, e0259608.	1.1	1
177	Amyloid neuropathy and autonomic dysfunction. Neurology and Clinical Neuroscience, 0, , .	0.2	3
178	High-avidity binding drives nucleation of amyloidogenic transthyretin monomer. JCI Insight, 2022, 7, .	2.3	2
179	Molecular Mechanisms of Cardiac Amyloidosis. International Journal of Molecular Sciences, 2022, 23, 25.	1.8	20
180	Analysis of amyloidogenic transthyretin mutations using continuum solvent free energy calculations. Proteins: Structure, Function and Bioinformatics, 2022, 90, 2080-2090.	1.5	2
181	Protein Aggregation in Neurodegenerative Diseases. , 2022, , 26-58.		0
182	A decade of approved first-in-class small molecule orphan drugs: Achievements, challenges and perspectives. European Journal of Medicinal Chemistry, 2022, 243, 114742.	2.6	1
183	Tissue-based diagnosis of systemic amyloidosis: Experience of the informal diagnostic center at Uppsala University Hospital. Upsala Journal of Medical Sciences, 0, 127, .	0.4	1