Intissar Anan

List of Publications by Year
in descending order

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28	336	759190 12	888047 17
papers	citations	h-index	g-index

28	28	28	365
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Effect of doxycycline and ursodeoxycholic acid on transthyretin amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2017, 24, 78-79.	3.0	38
2	Enthalpic Forces Correlate with the Selectivity of Transthyretin-Stabilizing Ligands in Human Plasma. Journal of Medicinal Chemistry, 2015, 58, 6507-6515.	6.4	35
3	Colonic enteric nervous system in patients with familial amyloidotic neuropathy. Acta Neuropathologica, 1999, 98, 48-54.	7.7	23
4	Abdominal fat pad biopsies exhibit good diagnostic accuracy in patients with suspected transthyretin amyloidosis. Orphanet Journal of Rare Diseases, 2020, 15, 278.	2.7	22
5	Loss of gastric interstitial cells of Cajal in patients with hereditary transthyretin amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2013, 20, 99-106.	3.0	20
6	Structural basis for transthyretin amyloid formation in vitreous body of the eye. Nature Communications, 2021, 12, 7141.	12.8	20
7	Management of gastrointestinal complications in hereditary transthyretin amyloidosis: a single-center experience over 40 years. Expert Review of Gastroenterology and Hepatology, 2018, 12, 73-81.	3.0	19
8	Endocrine cells in the upper gastrointestinal tract in relation to gastrointestinal dysfunction in patients with familial amyloidotic polyneuropathy. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 1999, 6, 192-198.	3.0	18
9	Comparison of amyloid deposits and infiltration of enteric nervous system in the upper with those in the lower gastrointestinal tract in patients with familial amyloidotic polyneuropathy. Acta Neuropathologica, 2001, 102, 227-232.	7.7	18
10	The Swedish open-label diflunisal trial (DFNS01) on hereditary transthyretin amyloidosis and the impact of amyloid fibril composition. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2019, 26, 39-40.	3.0	16
11	New data on the genetic profile and penetrance of hereditary Val30Met transthyretin amyloidosis in Sweden. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2021, 28, 84-90.	3.0	16
12	Atrial Fibrillation and Central Nervous Complications in Liver Transplanted Hereditary Transthyretin Amyloidosis Patients. Transplantation, 2018, 102, e59-e66.	1.0	14
13	Amyloid fibril composition within hereditary Val30Met (p. Val50Met) transthyretin amyloidosis families. PLoS ONE, 2019, 14, e0211983.	2.5	13
14	Endogenous Human Proteins Interfering with Amyloid Formation. Biomolecules, 2022, 12, 446.	4.0	9
15	Outcome of gastric emptying and gastrointestinal symptoms after liver transplantation for hereditary transthyretin amyloidosis. BMC Gastroenterology, 2015, 15, 51.	2.0	8
16	Combining ECG and echocardiography to identify transthyretin cardiac amyloidosis in heart failure. Clinical Physiology and Functional Imaging, 2021, 41, 408-416.	1.2	8
17	Metabolomics analysis for diagnosis and biomarker discovery of transthyretin amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2021, 28, 234-242.	3.0	8
18	Mechanisms of Transthyretin Inhibition of IAPP Amyloid Formation. Biomolecules, 2021, 11, 411.	4.0	7

#	Article	IF	Citations
19	Detection of a variant protein in hair: new diagnostic method in Portuguese type familial amyloid polyneuropathy Â Commentary: A new hair test for rare antigens. BMJ: British Medical Journal, 1998, 316, 1500-1501.	2.3	5
20	Investigation of AGE, their receptor and NF-kappaB activation and apoptosis in patients with ATTR and Gelsolin amyloidosis. Histology and Histopathology, 2010, 25, 691-9.	0.7	5
21	Transthyretin Glu54Leu â€“ an unknown mutation within the Swedish population associated with amyloid cardiomyopathy and a unique fibril type. Scandinavian Journal of Clinical and Laboratory Investigation, 2019, 79, 372-376.	1.2	3
22	Epidemiology of hereditary transthyretin amyloidosis in the northernmost region of Sweden: a retrospective cohort study. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2022, 29, 120-127.	3.0	3
23	Cerebellar and Cerebral Amyloid Visualized by [18F]flutemetamol PET in Long-Term Hereditary V30M (p.V50M) Transthyretin Amyloidosis Survivors. Frontiers in Neurology, 2022, 13, 816636.	2.4	3
24	The Swedish landscape of hereditary ATTR amyloidosis. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2017, 24, 93-94.	3.0	2
25	Self-reported gastrointestinal symptoms are more common in liver transplanted transthyretin amyloidosis patients than in healthy controls and in patients transplanted for end-stage liver disease. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2019, 26, 47-48.	3.0	2
26	Hereditary transthyretin amyloidosis in Sweden: Comparisons between a non-endemic and an endemic region. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2022, , 1-8.	3.0	1
27	A case report of osteoarthritis associated with hereditary transthyretin amyloidosis ATTRV30M. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2019, 26, 29-30.	3.0	0
28	Amyloid fibril composition type is consistent over time in patients with Val30Met (p.Val50Met) transthyretin amyloidosis. PLoS ONE, 2022, 17, e0266092.	2.5	0