Systemic Amyloidosis Recognition, Prognosis, and Ther

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Citation Report

#	Article	IF	Citations
1	SelecciÃ³n de lo mejor del aÃ±o 2020 en cardiopatÃas familiares y genÃ©tica cardiovascular. REC: CardioClinics, 2021, 56, 9-14.	0.1	0
2	Cardiovascular Toxicities of Proteasome Inhibitors. , 2021, , 59-67.		0
3	Immunoglobulin Light Chain Amyloidosis Presenting as Budd Chiari. Hepatology Forum, 2021, , .	0.5	0
4	Neuromuscular amyloidosis: Unmasking the master of disguise. Muscle and Nerve, 2021, 64, 23-36.	2.2	23
5	Patient and family experience with transthyretin amyloid cardiomyopathy (ATTR-CM) and polyneuropathy (ATTR-PN) amyloidosis: results of two focus groups. Orphanet Journal of Rare Diseases, 2021, 16, 70.	2.7	16
6	The Importance of Multimodality Imaging in the Diagnosis and Management of Patients with Infiltrative Cardiomyopathies: An Update. Diagnostics, 2021, 11, 256.	2.6	4
7	Immunoglobulin light chain amyloidosis. Memo - Magazine of European Medical Oncology, 2021, 14, 103-110.	0.5	1
8	Canadian Guidelines for Hereditary Transthyretin Amyloidosis Polyneuropathy Management. Canadian Journal of Neurological Sciences, 2022, 49, 7-18.	0.5	9
11	Transforming ATTR cardiac amyloidosis into a chronic disease: The enormous potential of quantitative SPECT to improve diagnosis, prognosis, and monitoring of disease progression. Journal of Nuclear Cardiology, 2021, 28, 1846-1850.	2.1	1
12	The Clinical Impact of Proteomics in Amyloid Typing. Mayo Clinic Proceedings, 2021, 96, 1122-1127.	3.0	9
13	Falseâ€positive bone scintigraphy denoting transthyretin amyloid in elderly hypertrophic cardiomyopathy. ESC Heart Failure, 2021, 8, 3387-3391.	3.1	13
14	Utility of abdominal skin punch biopsy for detecting systemic amyloidosis. Journal of Cutaneous Pathology, 2021, 48, 1342-1346.	1.3	7
15	As the Story Unfolds. Journal of Hospital Medicine, 2021, 16, 428-433.	1.4	0
16	Diagnostic delay and characterization of the clinical prodrome in AL amyloidosis among 1523 US adults diagnosed between 2001 and 2019. European Journal of Haematology, 2021, 107, 428-435.	2.2	18
17	Amyloidosis of the Urinary Bladder: A Systematic Review and a Proposed Management Algorithm. Urology, 2021, 156, e12-e19.	1.0	5
18	Improving contemporary outcomes following heart transplantation for cardiac amyloidosis. Journal of Cardiac Surgery, 2021, 36, 3509-3518.	0.7	2
19	Extremely Rare Intraoral Presentation of Localized Amyloidosis. Case Reports in Dentistry, 2021, 2021, 1-7.	0.5	1
20	Cardiac Amyloidosis â€“ An Underdiagnosed Cause of Heart Failure with Preserved Ejection Fraction â€“ Updated Diagnosis and Treatment Options. Revista Romana De Cardiologie, 2021, 31, 283-302.	0.1	1

#	Article	IF	Citations
21	18F-Florbetaben: a potential nuclear protagonist in the diagnosis of cardiac amyloidosis. Clinical and Translational Imaging, 2022, 10, 5-8.	2.1	1
22	Congestive heart failure associated with POEMS syndrome that was adequately distinguished from cardiac amyloidosis: a case report and literature review. Annals of Translational Medicine, 2021, 9, 1266-1266.	1.7	1
23	Differences in the characteristics and contemporary cardiac outcomes of patients with light-chain versus transthyretin cardiac amyloidosis. PLoS ONE, 2021, 16, e0255487.	2.5	8
24	Kidney Carbuncle in a Patient with Primary Systemic Al-Amyloidosis and Nephortic Syndrome. Russian Archives of Internal Medicine, 2021, 11, 303-309.	0.2	0
25	Difficulties in diagnosis of primary AL-amyloidosis. Oncogematologiya, 2021, 16, 74-82.	0.3	2
26	Posicionamento sobre DiagnÃ³stico e Tratamento da Amiloidose CardÃaca â€“ 2021. Arquivos Brasileiros De Cardiologia, 2021, 117, 561-598.	0.8	35
27	An unusual localized AL amyloidosis of parotid gland: A case report and literature review. Leukemia Research, 2021, 108, 106594.	0.8	0
28	Gross hematuria: An unusual presenting symptom of systemic wild-type transthyretin amyloidosis. Urology Case Reports, 2021, 39, 101811.	0.3	1
29	Real-World Data and Machine Learning to Predict Cardiac Amyloidosis. International Journal of Environmental Research and Public Health, 2021, 18, 908.	2.6	9
30	Amyloid Cardiomyopathy. JACC: CardioOncology, 2021, 3, 606-610.	4.0	3
31	Epidemiology and clinical manifestations of cardiac amyloidosis. Heart Failure Reviews, 2022, 27, 1471-1484.	3.9	13
32	Initial experience with percutaneous mitral valve repair in patients with cardiac amyloidosis. European Journal of Clinical Investigation, 2021, 51, e13473.	3.4	6
33	Diagnosis of amyloid beyond Congo red. Current Opinion in Nephrology and Hypertension, 2021, 30, 303-309.	2.0	9
34	A case of systemic AL amyloidosis identified by autopsy with suspected epicardial amyloid deposits on echocardiography. Choonpa Igaku, 2022, 49, 39-49.	0.0	0
35	Differences and Similarities in Treatment Paradigms and Goals between AL Amyloidosis and Multiple Myeloma. Hemato, 2021, 2, 680-691.	0.6	1
36	Assessing Cardiac Amyloidosis SubtypesÂby Unsupervised Phenotype Clustering Analysis. Journal of the American College of Cardiology, 2021, 78, 2177-2192.	2.8	11
37	Thromboembolism and bleeding in systemic amyloidosis: a review. ESC Heart Failure, 2022, 9, 11-20.	3.1	27
38	Bilateral carpal tunnel syndrome preceding haemorrhagic bullae in an older adult. Clinical and Experimental Dermatology, 2021, , .	1.3	0

#	Article	IF	Citations
39	Restrictive spirometry pattern and abnormal cardiopulmonary response to exercise in transthyretin cardiac amyloidosis. European Respiratory Journal, 2022, 59, 2102838.	6.7	6
40	The Importance of the DEIC/GEMIC/SBC 2021 Position Statement on Diagnosis and Treatment of Cardiac Amyloidosis to Clinical Practice. , 2021, 1, 80-83.		0
41	Vasculopatia AmiloidÃ³tica Associada a Mieloma MÃºltiplo. Revista De MedicinÄƒ InternÄƒ, Neurologe, Psihiatrie, Neurochirurgie, Dermato-venerologie MedicinÄƒ InternÄƒ, 2021, 28, 48-49.	0.0	0
42	Neuromuscular Complications of Systemic Amyloidosis. American Journal of Medicine, 2022, 135, S13-S19.	1.5	2
43	Sex-related differences in clinical presentation and all-cause mortality in patients with cardiac transthyretin amyloidosis and light chain amyloidosis. International Journal of Cardiology, 2022, 351, 71-77.	1.7	11
44	Mucocutaneous xanthodermatosis as initial presentation of immunoglobulin light chain amyloidosis. American Journal of Medicine, 2022, , .	1.5	0
45	Pearls from the First Gulf Cardiac Amyloidosis Summit 2021. Emirates Medical Journal, 2022, 3, 1-11.	0.3	0
46	Focused review on nutritional status of patients with immunoglobulin light chain amyloidosis. Current Problems in Cancer, 2022, 46, 100833.	2.0	3
47	Association Between Systemic Amyloidosis and Intracranial Hemorrhage. Stroke, 2022, 53, STROKEAHA121038451.	2.0	4
48	Pharmacodynamic evaluation and safety assessment of treatment with antibodies to serum amyloid P component in patients with cardiac amyloidosis: an open-label Phase 2 study and an adjunctive immuno-PET imaging study. BMC Cardiovascular Disorders, 2022, 22, 49.	1.7	14
49	Gastric amyloidosis associated with multiple myeloma: A rare cause of upper gastrointestinal bleeding. Advances in Digestive Medicine, 0, , .	0.2	0
50	Proposed Cardiac End Points for Clinical Trials in Immunoglobulin Light Chain Amyloidosis: Report From the Amyloidosis Forum Cardiac Working Group. Circulation: Heart Failure, 2022, 15, CIRCHEARTFAILURE121009038.	3.9	6
51	Value of repeat renal biopsy in the evaluation of AL amyloidosis patients lacking renal response despite of complete hematologic remission: a case report and literature review. BMC Nephrology, 2022, 23, 127.	1.8	0
52	Pulmonary 99mTc-HMDP uptake correlates with restrictive ventilatory defects and abnormal lung reactance in transthyretin cardiac amyloidosis patients. Respiratory Research, 2022, 23, 72.	3.6	2
53	AfectaciÃ³n cutÃ¡nea en las discrasias de las cÃ©lulas plasmÃ¡ticas. Piel, 2022, , .	0.0	0
54	Synthesis and Evaluation of Monoaryl Derivatives as Transthyretin Fibril Formation Inhibitors. Pharmaceutical Chemistry Journal, 2022, 56, 38-47.	0.8	2
55	Proteins Do Not Replicate, They Precipitate: Phase Transition and Loss of Function Toxicity in Amyloid Pathologies. Biology, 2022, 11, 535.	2.8	14
56	Novel approaches to diagnosis and management of hereditary transthyretin amyloidosis. Journal of Neurology, Neurosurgery and Psychiatry, 2022, 93, 668-678.	1.9	35

#	Article	IF	Citations
57	From Biology to Treatment of Monoclonal Gammopathies of Neurological Significance. Cancers, 2022, 14, 1562.	3.7	9
58	Navigating the Complex Web of Prescribing Amyloidosis Therapeutics: A Primer. Journal of the American Heart Association, 2022, 11, e023895.	3.7	1
59	Nomogram Model for Dynamic and Individual Prediction of Cardiac Response and Survival for Light Chain Amyloidosis in 737 Patients With Cardiac Involvement. Frontiers in Oncology, 2021, 11, 758502.	2.8	1
61	New Advanced Imaging Parameters and Biomarkersâ€"A Step Forward in the Diagnosis and Prognosis of TTR Cardiomyopathy. Journal of Clinical Medicine, 2022, 11, 2360.	2.4	4
62	Kidney involvement in hereditary transthyretin amyloidosis: a cohort study of 103 patients. CKJ: Clinical Kidney Journal, 2022, 15, 1747-1754.	2.9	10
63	â€œHere comes the story of the Hurricaneâ€ a case report of AL cardiac amyloidosis and myocardial bridging. European Heart Journal - Case Reports, 0, , .	0.6	3
64	Amyotrophic lateral sclerosis mimics. Muscle and Nerve, 2022, 66, 240-252.	2.2	5
65	Phase 2 trial of ixazomib, cyclophosphamide, and dexamethasone for previously untreated light chain amyloidosis. Blood Advances, 2022, 6, 5429-5435.	5.2	3
66	Clinical recommendations to diagnose and monitor patients with transthyretin amyloid cardiomyopathy in Asia. Clinical Cardiology, 2022, 45, 898-907.	1.8	3
67	Renal Amyloidosis: Epidemiological, Clinical, and Laboratory Profile in Adults from One Nephrology Center. International Journal of Nephrology, 2022, 2022, 1-9.	1.3	2
68	Oral, periorbital, and inguinal purpura in a patient with paraproteinemia. JAAD Case Reports, 2022, 27, 79-81.	0.8	0
69	Cardiac Amyloidosis with Normal Wall Thickness: Prevalence, Clinical Characteristics and Outcome in a Retrospective Analysis. Biomedicines, 2022, 10, 1765.	3.2	3
70	Cardiac amyloidosis: a review of the literature. Acta Cardiologica, 2022, 77, 683-692.	0.9	8
71	Cutaneous Ossifying Amyloidoma. American Journal of Dermatopathology, 2022, 44, 760-763.	0.6	1
72	Beyond Sarcomeric Hypertrophic Cardiomyopathy: How to Diagnose and Manage Phenocopies. Current Cardiology Reports, 2022, 24, 1567-1585.	2.9	14
73	Micro- and Nanoplasticsâ€™ Effects on Protein Folding and Amyloidosis. International Journal of Molecular Sciences, 2022, 23, 10329.	4.1	11
74	Prevalence of Amyloid Deposition in Patients Undergoing Surgical Repair of Traumatic Distal Biceps Tendon Ruptures. Journal of Hand Surgery Global Online, 2022, 4, 344-347.	0.8	4
75	Feasibility of 68Ga-Labeled Fibroblast Activation Protein Inhibitor PET/CT in Light-Chain CardiacÂAmyloidosis. JACC: Cardiovascular Imaging, 2022, 15, 1960-1970.	5.3	13

#	Article	IF	Citations
76	The Role of Echocardiogram and Cardiac Rhythm Analysis for Early Detection of Cardiac Amyloidosis. Journal of the Advanced Practitioner in Oncology, 2022, 13, 695-704.	0.4	1
77	Independent prognostic value of left ventricular mass index in patients with light-chain amyloidosis. Acta Cardiologica, 0, , 1-7.	0.9	0
78	Clinical characteristics and prognostic value of renal immune complex deposition in patients with light chain amyloidosis. Frontiers in Oncology, 0, 12, .	2.8	1
79	Graded Cardiac Response Criteria for Patients With Systemic Light Chain Amyloidosis. Journal of Clinical Oncology, 2023, 41, 1393-1403.	1.6	11
80	Systemic and Cardiac Amyloidosis in Captive Celebes Crested Macaques (Macaca nigra). Journal of Comparative Pathology, 2022, 199, 12-16.	0.4	0
81	Noncardiac Manifestations of Hereditary Amyloidosis. American Journal of Cardiology, 2022, , .	1.6	0
82	Monoclonal gammopathy of increasing significance: time to screen?. Haematologica, 0, , .	3.5	2
83	AL Amyloidosis for Cardiologists. JACC: CardioOncology, 2022, 4, 427-441.	4.0	19
84	Diagnosis and classification of vascular liver disorders. Diagnostic Histopathology, 2023, 29, 36-51.	0.4	1
85	Cardiac amyloidosisâ€"An underdiagnosed cause of heart failure: A case report and review of literature. Clinical Case Reports (discontinued), 2022, 10, .	0.5	0
86	Current Understanding of Systemic Amyloidosis and Underlying Disease Mechanisms. American Journal of Cardiology, 2022, 185, S2-S10.	1.6	1
87	Systemic amyloidosis journey from diagnosis to outcomes: a twelve-year real-world experience of a single center in a middle-income country. Orphanet Journal of Rare Diseases, 2022, 17, .	2.7	3
88	Collagen inhibits phagocytosis of amyloid inÂvitro and inÂvivo and may act as a â€˜donâ€™t eat meâ€™ signal. Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis, 2023, 30, 249-260.	3.0	3
89	Immunoglobulin Light Chain Amyloidosis: Diagnosis and Risk Assessment. Journal of the National Comprehensive Cancer Network: JNCCN, 2023, 21, 83-90.	4.9	1
91	Concurrent Presentation of Light Chain Amyloidosis and Renal Cell Carcinoma in an Elderly Male : A Case Report and Review of the Literature. , 2023, 2, .		0
92	The Clinical Characteristics of Immunoglobulin Light Chain Amyloidosis in the Chinese Population: A Systematic Scoping Review. Hemato, 2023, 4, 12-25.	0.6	0
93	Amyloidosis and autonomic failure. , 2023, , 575-583.		0
94	Hematological Associations in Amyloidosis. , 2023, , 119-128.		0

#	Article	IF	Citations
95	Cerebral Infarction as the First Manifestation of Multiple Myeloma and AL Amyloidosis: a Case Report. SN Comprehensive Clinical Medicine, 2023, 5, .	0.6	1
96	State-of-the-art review on AL amyloidosis in Western Countries: Epidemiology, health economics, risk assessment and therapeutic management of a rare disease. Blood Reviews, 2023, 59, 101040.	5.7	4
97	Neurological Manifestations in AL and Wild-Type ATTR Amyloidosis. , 2023, , 19-24.		0
98	Hepatic amyloidosis accompanied with massive ascites and kidney impairment: A case report. , 2023, 1, 24-26.		0
99	Artificial intelligenceâ€“enabled classification of hypertrophic heart diseases using electrocardiograms. Cardiovascular Digital Health Journal, 2023, 4, 48-59.	1.3	6
100	The exploration of B cell maturation antigen expression in plasma cell dyscrasias beyond multiple myeloma. BMC Cancer, 2023, 23, .	2.6	1
101	Chronic Pain and Quality of Life. , 2023, , 467-473.		0
102	Spatial resolution of renal amyloid deposits through MALDI-MSI: a combined digital and molecular approach to monoclonal gammopathies. Journal of Clinical Pathology, 0, , jcp-2023-208790.	2.0	3
103	Diagnostic Challenges and Solutions in Systemic Amyloidosis. International Journal of Molecular Sciences, 2023, 24, 4655.	4.1	2
104	Cardiac Amyloidosis. Annals of Internal Medicine, 2023, 176, ITC33-ITC48.	3.9	8
105	Machine Learning Approaches in Diagnosis, Prognosis and Treatment Selection of Cardiac Amyloidosis. International Journal of Molecular Sciences, 2023, 24, 5680.	4.1	6
106	The role of trichoscopy beyond hair and scalp diseases: A review. Journal of the European Academy of Dermatology and Venereology, 2023, 37, 1473-1479.	2.4	1
107	Comprehensive Heart Failure History. , 2023, , 23-34.		0
108	Systemic Amyloidosis: Is It ATTR or AL?. American Journal of Medicine, 2023, , .	1.5	0
109	Brazilian consensus for diagnosis, management and treatment of hereditary transthyretin amyloidosis with peripheral neuropathy: second edition. Arquivos De Neuro-Psiquiatria, 2023, 81, 308-321.	0.8	0
110	Amyloidosis: a case series and review of theÂliterature. Journal of Medical Case Reports, 2023, 17, .	0.8	1
111	Prognostic Factors and Survival of Chinese Patients with Cardiac Amyloidosis. Disease Markers, 2023, 2023, 1-8.	1.3	0
112	Systemic AL amyloidosis: current approach and future direction. Oncotarget, 2023, 14, 384-394.	1.8	3

#	Article	IF	Citations
113	Gastroparesis: an under-recognised manifestation of systemic amyloidosis. BMJ Case Reports, 2023, 16, e254634.	0.5	0
114	SFTSV infection is associated with transient overproliferation of monoclonal lambda-type plasma cells. IScience, 2023, 26, 106799.	4.1	0
115	Yellowish papules and plaques in the periocular area. Clinical and Experimental Dermatology, 0, , .	1.3	0
116	Heart Failure, Neuropathy, and Spinal Stenosis. JAMA - Journal of the American Medical Association, 0, , .	7.4	0
117	Risk Stratification in Transthyretin Cardiac Amyloidosis: The Added Value of Lung Spirometry. Journal of Clinical Medicine, 2023, 12, 3684.	2.4	0
118	Amyloid fibrils degradation: the pathway to recovery or aggravation of the disease?. Frontiers in Molecular Biosciences, 0, 10, .	3.5	1
119	Diagnosis for Chinese patients with light chain amyloidosis: a scoping review. Annals of Medicine, 2023, 55, 1144-1155.	3.8	1
120	Cardiovascular toxicity from therapies for light chain amyloidosis. Frontiers in Cardiovascular Medicine, 0, 10, .	2.4	0
121	Refractory Arrhythmias as a Potential Indicator of Underlying Cardiac Amyloidosis: A Case Report. Cureus, 2023, , .	0.5	0
122	Epidemiology and clinical outcomes of lightâ€chain amyloidosis in Sweden: A nationwide populationâ€based study. European Journal of Haematology, 2023, 111, 697-705.	2.2	0
123	Strong positive light chain immunostaining in a patient with transthyretin amyloidosis. Hematology, 2023, 28, .	1.5	0
124	A Complex Case of Extensive Systemic Amyloidosis With Underlying Monoclonal Gammopathy. Cureus, 2023, , .	0.5	0
125	Development of small molecules for disrupting pathological amyloid aggregation in neurodegenerative diseases. , 0, 3, .		0
126	Iatrogenic Acute Ascending Aortic Dissection During Combined Heart/Liver Transplant for Amyloidosis. Transplantation Proceedings, 2023, , .	0.6	0
127	Clinical effects of tranexamic acid on bleeding tendency due to fibrinolytic activation of AL amyloidosis. BMJ Case Reports, 2023, 16, e254483.	0.5	0
128	Peripheral Nervous, Hepatic, and Gastrointestinal Endpoints for AL Amyloidosis Clinical Trials: Report from the Amyloidosis Forum Multi-organ System Working Group. Advances in Therapy, 2023, 40, 4695-4710.	2.9	1
129	Monitoring cardiac amyloidosis with multimodality imaging. Revista Espanola De Cardiologia (English) Tj ETQq0 0 0 rgBT /Overlock 10 Tf	0.8	0
130	Amiloidosis sistÃ©mica con Ã©nfasis en amiloidosis de cadenas livianas de inmunoglobulinas. Medicina Y Laboratorio, 2023, 27, 395-409.	0.1	0

#	Article	IF	Citations
131	Development and characterization of a prototypic pan-amyloid clearing agent â€“ a novel murine peptide-immunoglobulin fusion. Frontiers in Immunology, 0, 14, .	4.8	0
132	Pathomorphological Features of the Novel Coronavirus Disease in Patients with Systemic Amyloidosis. Biomedicines, 2023, 11, 2811.	3.2	0
133	Amyloid detection and typing yield of skin biopsy in systemic amyloidosis and polyneuropathy. Annals of Clinical and Translational Neurology, 2023, 10, 2347-2359.	3.7	1
134	Emerging therapeutic avenues in cardiac amyloidosis. European Journal of Pharmacology, 2023, 960, 176142.	3.5	0
136	Caregiverâ€™s psychological well-being and quality of relationship with cardiac amyloidosis patients. Psychology, Health and Medicine, 2024, 29, 66-78.	2.4	0
137	Chitosan oligosaccharides inhibit the fibrillation of insulin and disassemble its preformed fibrils. International Journal of Biological Macromolecules, 2024, 254, 127857.	7.5	0
138	Dermatoscopy of primary localised cutaneous amyloidosis â€•A crossâ€sectional study in a setting of South Asian public dermatology department. Skin Health and Disease, 0, , .	1.5	1
139	Perturbational phenotyping of human blood cells reveals genetically determined latent traits associated with subsets of common diseases. Nature Genetics, 0, , .	21.4	1
141	Characterization of Amyloidosis in the Gastrointestinal Tract With an Emphasis on Histologically Distinct Interstitial Patterns of Deposition and Misinterpretations. American Journal of Surgical Pathology, 2024, 48, 302-308.	3.7	0
142	Fineâ€needle aspirationÂof amyloidoma: A critical analysis. Cancer Cytopathology, 2024, 132, 179-185.	2.4	0
144	Early diagnosis and management of cardiac amyloidosis: A clinical perspective. European Journal of Clinical Investigation, 0, , .	3.4	0
145	Prediction of the Feasibility of Using the â‰ªGold Standardâ‰« Thioflavin T to Detect Amyloid Fibril in Acidic Media. Analytical Chemistry, 2024, 96, 2158-2164.	6.5	0
146	Update on Amyloid Polyneuropathy and Treatment. Current Treatment Options in Neurology, 2024, 26, 51-66.	1.8	0
147	A Comparison of Amyloid Deposition inÂEndoscopic and Open Carpal TunnelÂRelease. Journal of Hand Surgery, 2024, 49, 301-309.	1.6	0
148	Being a Better Starfish Thrower, CarpalÂTunnel Syndrome, and CardiacÂAmyloidosis: The Hand Surgeonâ€™sÂOpportunity to Make aÂProfoundÂDifference. Journal of Hand Surgery, 2024, 49, 373-376.	1.6	0
149	Cardiac Amyloidosis Due to Transthyretin Protein. JAMA - Journal of the American Medical Association, 2024, 331, 778.	7.4	0
150	AmyKoSâ€”a German real-world prospective cohort study of patients with suspected and proven amyloidosis. , 2024, 3, .		0
151	Acute Exacerbation of Idiopathic Pulmonary Fibrosis With Concurrent Cardiac Amyloidosis: A Technetium Pyrophosphate Study. Cureus, 2024, , .	0.5	0

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