Chronic kidney disease and an uncertain diagnosis of Fa diagnosis

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Citation Report

#	Article	IF	CITATIONS
1	Uncertain Diagnosis of Fabry Disease in Patients with Neuropathic Pain, Angiokeratoma or Cornea Verticillata: Consensus on the Approach to Diagnosis and Follow-Up. JIMD Reports, 2014, 17, 83-90.	0.7	42
2	In Patients with an α-Galactosidase A Variant, Small Nerve Fibre Assessment Cannot Confirm a Diagnosis of Fabry Disease. JIMD Reports, 2015, 28, 95-103.	0.7	6
3	NefropatÃas hereditarias y congénitas. Medicine, 2015, 11, 4793-4802.	0.0	2
4	Variations in the GLA gene correlate with globotriaosylceramide and globotriaosylsphingosine analog levels in urine and plasma. Clinica Chimica Acta, 2015, 447, 96-104.	0.5	22
5	Recommendations for initiation and cessation of enzyme replacement therapy in patients with Fabry disease: the European Fabry Working Group consensus document. Orphanet Journal of Rare Diseases, 2015, 10, 36.	1.2	239
6	Enzyme Replacement Therapy for Fabry Disease. FIRE Forum for International Research in Education, 2016, 4, 232640981667942.	0.7	1
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8	Genetic Screening of Anderson-Fabry Disease in Probands Referred From Multispecialty Clinics. Journal of the American College of Cardiology, 2016, 68, 1037-1050.	1.2	50
10	Diagnóstico y tratamiento de la enfermedad de Fabry. Medicina ClÃnica, 2017, 148, 132-138.	0.3	19
11	Diagnosis and treatment of Fabry disease. Medicina ClÃnica (English Edition), 2017, 148, 132-138.	0.1	2
13	Clinical proteomics in kidney disease as an exponential technology: heading towards the disruptive phase. CKJ: Clinical Kidney Journal, 2017, 10, 188-191.	1.4	22
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15	Fabry Nephropathy: An Evidence-Based Narrative Review. Kidney and Blood Pressure Research, 2018, 43, 406-421.	0.9	35
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17	Bedside Stereomicroscopy of Fabry Kidney Biopsies: An Easily Available Method for Diagnosis and Assessment of Sphingolipid Deposits. Nephron, 2018, 138, 13-21.	0.9	6
18	ll dolore nella Malattia di Fabry. Giornale De Techniche Nefrologiche & Dialitiche, 2018, 30, 153-157.	0.1	0
19	Systematic DNA Study for Fabry Disease in the End Stage Renal Disease Patients from a Southern Italy Area. Kidney and Blood Pressure Research, 2018, 43, 1344-1351.	0.9	7
20	Clinical significance of plasma globotriaosylsphingosine levels in Chinese patients with Fabry disease. Experimental and Therapeutic Medicine, 2018, 15, 3733-3742.	0.8	15

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21	Fabry disease caused by the GLA p.Phe113Leu (p.F113L) variant: Natural history in males. European Journal of Medical Genetics, 2020, 63, 103703.	0.7	21
22	Biomarkers of Fabry Nephropathy: Review and Future Perspective. Genes, 2020, 11, 1091.	1.0	16
23	Manifestaciones osteoarticulares de las esfingolipidosis en adultos. EMC - Aparato Locomotor, 2020, 53, 1-29.	0.1	0
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25	Recurrent fever of unknown origin: An overlooked symptom of Fabry disease. Molecular Genetics & Genomic Medicine, 2020, 8, e1454.	0.6	1
26	Early indicators of disease progression in Fabry disease that may indicate the need for disease-specific treatment initiation: findings from the opinion-based PREDICT-FD modified Delphi consensus initiative. BMJ Open, 2020, 10, e035182.	0.8	20
27	<p>Diagnosis and Screening of Patients with Fabry Disease</p> . Therapeutics and Clinical Risk Management, 2020, Volume 16, 551-558.	0.9	37
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33	Renal Manifestations of Fabry Disease: A Narrative Review. Canadian Journal of Kidney Health and Disease, 2021, 8, 205435812098562.	0.6	18
34	Skin Globotriaosylceramide 3 Load Is Increased in Men with Advanced Fabry Disease. PLoS ONE, 2016, 11, e0166484.	1.1	11
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41	Clinical and diagnostic aspects of Fabry disease management: a narrative review with a particular focus on Brazilian experts' perspectives. Journal of Inborn Errors of Metabolism and Screening, 0, 10, .	0.3	1
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46	Diagnostic Flowchart in Fabry Disease. , 2023, , 359-365.		Ο