Fabry disease â€˜The New Great Imposterâ€: results of Medicine Departments (FIMeD)

Clinical Genetics

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Citation Report

#	Article	IF	Citations
2	Cerebral hemodynamics and endothelial function in patients with Fabry disease. BMC Neurology, 2013, 13, 170.	0.8	16
3	Difficulties and barriers in diagnosing Fabry disease: what can be learnt from the literature?. Expert Opinion on Medical Diagnostics, 2013, 7, 589-599.	1.6	26
4	Misdiagnosis of familial Mediterranean fever in patients with Andersonâ€“Fabry disease. Clinical Genetics, 2013, 83, 576-581.	1.0	18
5	Analysis of the landscape of biologically-derived pharmaceuticals in Europe: Dominant production systems, molecule types on the rise and approval trends. European Journal of Pharmaceutical Sciences, 2013, 48, 428-441.	1.9	31
6	Mutation identification of Fabry disease in families with other lysosomal storage disorders. Clinical Genetics, 2013, 84, 281-285.	1.0	1
7	Angiokeratomas â€“ When is a few too many?. International Journal of STD and AIDS, 2014, 25, 378-379.	0.5	3
8	Prevalence of Raynaud Phenomenon and Nailfold Capillaroscopic Abnormalities in Fabry Disease. Medicine (United States), 2015, 94, e780.	0.4	5
9	High Variability of Fabry Disease Manifestations in an Extended Italian Family. BioMed Research International, 2015, 2015, 1-5.	0.9	23
10	Diagnosing Fabry disease--delays and difficulties within discordant siblings. QJM - Monthly Journal of the Association of Physicians, 2015, 108, 585-590.	0.2	4
11	Diagnostic dilemma and delay in <scp>F</scp>abry disease: Insights from a case series of young female patients. Journal of Paediatrics and Child Health, 2015, 51, 369-372.	0.4	3
12	Fabry disease in infancy and early childhood: a systematic literature review. Genetics in Medicine, 2015, 17, 323-330.	1.1	82
13	Sudoscan as a noninvasive tool to assess sudomotor dysfunction in patients with Fabry disease: results from a case–control study. Therapeutics and Clinical Risk Management, 2016, 12, 135.	0.9	17
14	The impact of fever/hyperthermia in the diagnosis of Fabry: A retrospective analysis. European Journal of Internal Medicine, 2016, 32, 26-30.	1.0	9
15	Sudden death following AV node ablation in a man with Fabry disease mimicking hypertrophic cardiomyopathy. Journal of Clinical Forensic and Legal Medicine, 2016, 42, 8-10.	0.5	0
17	Molecular and clinical studies in five index cases with novel mutations in the GLA gene. Gene, 2016, 578, 100-104.	1.0	20
18	Musculoskeletal manifestations of Fabry disease: A retrospective study. Joint Bone Spine, 2016, 83, 421-426.	0.8	21
19	Manifestations rhumatologiques de la maladie de FabryÂ: Ã©tude rÃ©trospective. Revue Du Rhumatisme (Edition Francaise), 2016, 83, 56-61.	0.0	1
20	A painful diagnosis. Internal and Emergency Medicine, 2017, 12, 341-347.	1.0	0

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21	Disease Progression Modeling to Evaluate the Effects of Enzyme Replacement Therapy on Kidney Function in Adult Patients with the Classic Phenotype of Fabry Disease. Kidney and Blood Pressure Research, 2017, 42, 1-15.	0.9	12
22	Recommendations for the inclusion of Fabry disease as a rare febrile condition in existing algorithms for fever of unknown origin. Internal and Emergency Medicine, 2017, 12, 1059-1067.	1.0	7
24	Fabry Disease in Internal Medicine: The Role of Fever and Hyperthermia in Diagnosis. Giornale De Techniche Nefrologiche & Dialitiche, 2017, 29, S12-S15.	0.1	0
25	Misdiagnosis. Giornale De Techniche Nefrologiche & Dialitiche, 2017, 29, S3-S4.	0.1	0
26	Identification of a novel loss-of-function mutation of the GLA gene in a Chinese Han family with Fabry disease. BMC Medical Genetics, 2018, 19, 219.	2.1	1
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36	AA amyloidosis associated with Fabry disease. International Journal of Clinical Practice, 2020, 74, e13577.	0.8	1
37	Prevalence of GLA gene mutations and polymorphisms in patients with multiple sclerosis: A cross-sectional study. Journal of the Neurological Sciences, 2020, 412, 116782.	0.3	2
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48	A 39Âyear-old man with acroparesthesia and uncommon renal arterial lesions. What is theÂdiagnosis?. Journal of Nephrology, 2022, , 1.	0.9	0
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