Anders Vahlquist

List of Publications by Year in descending order

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361413 377865 1,902 36 20 34 citations h-index g-index papers 36 36 36 1592 docs citations times ranked citing authors all docs

#	Article	IF	Citations
1	Meta-Analysis of Mutations in ALOX12B or ALOXE3 Identified in a Large Cohort of 224 Patients. Genes, 2021, 12, 80.	2.4	20
2	Exploration of novel candidate genes involved in epidermal keratinocyte differentiation and skin barrier repair in man. Differentiation, 2021, 119, 19-27.	1.9	4
3	Ichthyosis: A Road Model for Skin Research. Acta Dermato-Venereologica, 2020, 100, adv00097-206.	1.3	19
4	Patients with congenital ichthyosis and TGM 1 mutations overexpress other ARCI genes in the skin: Part of a barrier repair response?. Experimental Dermatology, 2019, 28, 1164-1171.	2.9	12
5	Inherited Nonsyndromic Ichthyoses: An Update on Pathophysiology, Diagnosis and Treatment. American Journal of Clinical Dermatology, 2018, 19, 51-66.	6.7	133
6	Quantitative image analysis of protein expression and colocalisation in skin sections. Experimental Dermatology, 2018, 27, 196-199.	2.9	9
7	Revertant mosaicism repairs skin lesions in a patient with keratitis-ichthyosis-deafness syndrome by second-site mutations in connexin 26. Human Molecular Genetics, 2017, 26, 1070-1077.	2.9	25
8	Spectrum of Autosomal Recessive Congenital Ichthyosis in Scandinavia: Clinical Characteristics and Novel and Recurrent Mutations in 132 Patients. Acta Dermato-Venereologica, 2016, 96, 932-937.	1.3	58
9	Recurrent Pyoderma Gangrenosum and Cystic Acne Associated with Leucocyte Adhesion Deficiency due to Novel Mutations in ITGB2: Successful Treatment with Infliximab and Adalimumab. Acta Dermato-Venereologica, 2015, 95, 349-351.	1.3	18
10	Interactions between FATP4 and ichthyin in epidermal lipid processing may provide clues to the pathogenesis of autosomal recessive congenital ichthyosis. Journal of Dermatological Science, 2013, 69, 195-201.	1.9	25
11	Generalized and Naevoid Epidermolytic Ichthyosis in Denmark: Clinical and Mutational Findings. Acta Dermato-Venereologica, 2013, 93, 309-313.	1.3	15
12	Oral Alitretinoin in Congenital Ichthyosis: A Pilot Study Shows Variable Effects and a Risk of Central Hypothyroidism. Acta Dermato-Venereologica, 2012, 92, 256-257.	1.3	16
13	The Expression of Epidermal Lipoxygenases and Transglutaminase-1 Is Perturbed by NIPAL4 Mutations: Indications of a Common Metabolic Pathway Essential for Skin Barrier Homeostasis. Journal of Investigative Dermatology, 2012, 132, 2368-2375.	0.7	29
14	Ichthyin/NIPAL4 localizes to keratins and desmosomes in epidermis and Ichthyin mutations affect epidermal lipid metabolism. Archives of Dermatological Research, 2012, 304, 377-386.	1.9	17
15	Treatment of rare keratinization disorders: what's new?. Expert Review of Dermatology, 2011, 6, 211-216.	0.3	2
16	Filaggrin Genotype Determines Functional and Molecular Alterations in Skin of Patients with Atopic Dermatitis and Ichthyosis Vulgaris. PLoS ONE, 2011, 6, e28254.	2.5	58
17	Genotypic and Clinical Spectrum of Self-Improving Collodion Ichthyosis: ALOX12B, ALOXE3, and TGM1 Mutations in Scandinavian Patients. Journal of Investigative Dermatology, 2010, 130, 438-443.	0.7	87
18	Pleomorphic Ichthyosis: Proposed Name for a Heterogeneous Group of Congenital Ichthyoses with Phenotypic Shifting and Mild Residual Scaling. Acta Dermato-Venereologica, 2010, 90, 454-460.	1.3	30

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19	Epidermolysis Bullosa Care in Scandinavia. Dermatologic Clinics, 2010, 28, 425-427.	1.7	12
20	Revised nomenclature and classification of inherited ichthyoses: Results of the First Ichthyosis Consensus Conference in SorÃ"ze 2009. Journal of the American Academy of Dermatology, 2010, 63, 607-641.	1.2	610
21	Expression of Retinoid-regulated Genes in Lamellar Ichthyosis vs. Healthy Control Epidermis: Changes after Oral Treatment with Liarozole. Acta Dermato-Venereologica, 2009, 89, 12-20.	1.3	30
22	Mutations in the Fatty Acid Transport Protein 4 Gene Cause the Ichthyosis Prematurity Syndrome. American Journal of Human Genetics, 2009, 85, 248-253.	6.2	147
23	Congenital Ichthyosis: An Overview of Current and Emerging Therapies. Acta Dermato-Venereologica, 2008, 88, 4-14.	1.3	110
24	Autosomal Recessive Congenital Ichthyosis in Sweden and Estonia: Clinical, Genetic and Ultrastructural Findings in Eighty-three Patients. Acta Dermato-Venereologica, 2003, 83, 24-30.	1.3	30
25	The clinical spectrum of congenital ichthyosis in Sweden: a review of 127 cases. Acta Dermato-venereologica Supplementum, 2003, , 34-47.	0.0	4
26	Keratin 4 Upregulation by Retinoic Acid In Vivo: A Sensitive Marker for Retinoid Bioactivity in Human Epidermis1. Journal of Investigative Dermatology, 2000, 114, 487-493.	0.7	33
27	The vitamin A metabolism and expression of retinoid-binding proteins differ in HaCaT cells and normal human keratinocytes. Archives of Dermatological Research, 1999, 291, 339-345.	1.9	24
28	Positive and negative interaction of 1,25-dihydroxyvitamin D3 and the retinoid CD437 in the induction of human melanoma cell apoptosis., 1999, 81, 467-470.		43
29	Treatment of psoriatic arthritis with extracorporeal photochemotherapy and conventional psoralen–ultraviolet A irradiation. Arthritis and Rheumatism, 1996, 39, 1519-1523.	6.7	46
30	Impaired release of vitamin a from liver in primary biliary cirrhosis. Hepatology, 1988, 8, 136-141.	7.3	39
31	Retinoids and Keratinization International Journal of Dermatology, 1988, 27, 81-95.	1.0	34
32	Clinical use of vitamin A and its derivatives- physiological and pharmacological aspects*. Clinical and Experimental Dermatology, 1985, 10, 133-143.	1.3	9
33	UV treatment of uraemic pruritus reduces the vitamin A content of the skin. European Journal of Clinical Investigation, 1984, 14, 203-206.	3.4	75
34	Hypercarotenemia and hypervitaminosis A. Clinical and Experimental Dermatology, 1981, 6, 225-225.	1.3	0
35	Retinol-binding protein in serum and epidermis of patients with ichthyosis vulgaris. Clinical and Experimental Dermatology, 1979, 4, 445-451.	1.3	5
36	Serum zinc and retinol-binding protein in acne. British Journal of Dermatology, 1977, 96, 283-286.	1.5	74