Anna M Rose

List of Publications by Year in descending order

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ANNA M ROSE

#	Article	IF	CITATIONS
1	Risk and Lag-time for Development of Lacrimal Gland Carcinoma Expleomorphic Adenoma: Observations and Conjectural Study. Ophthalmic Plastic and Reconstructive Surgery, 2022, Publish Ahead of Print, .	0.8	0
2	A remarkable case of HbH disease illustrates the relative contributions of the α-globin enhancers to gene expression. Blood, 2021, 137, 572-575.	1.4	6
3	Anti-neutrophil cytoplasmic antibody–positive vasculitis secondary to therapeutic levamisole. Rheumatology, 2020, 59, 2680-2680.	1.9	1
4	Multiple primary malignancies and prolonged survival in a patient with widespread metastatic cutaneous melanoma. Melanoma Research, 2018, 28, 163-166.	1.2	0
5	Gene of the month: <i>PRPF31</i> . Journal of Clinical Pathology, 2017, 70, 729-732.	2.0	6
6	Presentation, Treatment, and Prognosis of Secondary Melanoma within the Orbit. Frontiers in Oncology, 2017, 7, 125.	2.8	20
7	Transcriptional regulation of PRPF31 gene expression by MSR1 repeat elements causes incomplete penetrance in retinitis pigmentosa. Scientific Reports, 2016, 6, 19450. Book ReviewsOxford Handbook of Anaesthesia (4th edn) Edited by Keith G Allman, Iain H Wilson	3.3	42
8	Oxford University Press 2016 Price £34.99. Pp 1266 ISBN 978 0 1998719410ECG from Basics to Essentials: Step by Step Roland X Stroobandt, S Serge Barold, Alfons F Sinnaeve Wiley-Blackwell 2016 Price £43.99. Pp 431 ISBN 978 1 119 06641 5Strategic Scientific and Medical Writing Pieter H Joubert, Silvia M Rogers Springer 2015 Price £53.99. Pp 140 ISBN 978 3 662 48315 2. British Journal of Hospital Medicine (London,) Tj	0.5 ETQq0 0 0	0) rgBT /Overlo
9	Aetiology and management of malnutrition in HIV-positive children. Archives of Disease in Childhood, 2014, 99, 546-551.	1.9	47
10	Dominant <i>PRPF31</i> Mutations Are Hypostatic to a Recessive <i>CNOT3</i> Polymorphism in Retinitis Pigmentosa: A Novel Phenomenon of "Linked <i>Trans</i> Acting Epistasis― Annals of Human Genetics, 2014, 78, 62-71.	0.8	28
11	A Rare Case of Orbital Haemangiopericytoma Arising in Childhood. Orbit, 2013, 32, 384-386.	0.8	6
12	A Study into the Evolutionary Divergence of the Core Promoter Elements of PRPF31 and TFPT. Journal of Molecular and Genetic Medicine: an International Journal of Biomedical Research, 2013, 07, .	0.1	1
13	CNOT3 Is a Modifier of PRPF31 Mutations in Retinitis Pigmentosa with Incomplete Penetrance. PLoS Genetics, 2012, 8, e1003040.	3.5	109
14	Expression of PRPF31 and TFPT: regulation in health and retinal disease. Human Molecular Genetics, 2012, 21, 4126-4137.	2.9	15
15	Epistasis and immunity: the role of genetic interactions in autoimmune diseases. Immunology, 2012, 137, 131-138.	4.4	32
16	A 112 kb Deletion in Chromosome 19q13.42 Leads to Retinitis Pigmentosa. , 2011, 52, 6597.		22