Marvin E Reid

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/9098944/publications.pdf

Version: 2024-02-01

153 papers	3,667 citations	126907 33 h-index	175258 52 g-index
154 all docs	154 docs citations	154 times ranked	4383 citing authors

#	Article	IF	Citations
1	Deficient synthesis of glutathione underlies oxidative stress in aging and can be corrected by dietary cysteine and glycine supplementation. American Journal of Clinical Nutrition, 2011, 94, 847-853.	4.7	246
2	Improvements in haemolysis and indicators of erythrocyte survival do not correlate with acute vasoâ€•occlusive crises in patients with sickle cell disease: a phase III randomized, placeboâ€•controlled, doubleâ€•blind study of the gardos channel blocker senicapoc (ICAâ€•17043). British Journal of Haematology, 2011, 153, 92-104.	2.5	185
3	Cysteine supplementation improves the erythrocyte glutathione synthesis rate in children with severe edematous malnutrition. American Journal of Clinical Nutrition, 2002, 76, 646-652.	4.7	149
4	Glutathione in disease. Current Opinion in Clinical Nutrition and Metabolic Care, 2001, 4, 65-71.	2.5	102
5	Variant-aware saturating mutagenesis using multiple Cas9 nucleases identifies regulatory elements at trait-associated loci. Nature Genetics, 2017, 49, 625-634.	21.4	96
6	Frequency of pain crises in sickle cell anemia and its relationship with the sympatho-vagal balance, blood viscosity and inflammation. Haematologica, 2011, 96, 1589-1594.	3.5	93
7	Albuminuria and Renal Function in Homozygous Sickle Cell Disease. Archives of Internal Medicine, 2007, 167, 701.	3.8	85
8	In vivo rates of erythrocyte glutathione synthesis in children with severe protein-energy malnutrition. American Journal of Physiology - Endocrinology and Metabolism, 2000, 278, E405-E412.	3.5	83
9	Growth, Body Composition, and the Onset of Puberty: Longitudinal Observations in Afro-Caribbean Children. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 3194-3200.	3.6	73
10	Protein metabolism in severe childhood malnutrition. Annals of Tropical Paediatrics, 2008, 28, 87-101.	1.0	72
11	In vivo rates of erythrocyte glutathione synthesis in adults with sickle cell disease. American Journal of Physiology - Endocrinology and Metabolism, 2006, 291, E73-E79.	3.5	65
12	The acute-phase protein response to infection in edematous and nonedematous protein-energy malnutrition. American Journal of Clinical Nutrition, 2002, 76, 1409-1415.	4.7	64
13	Newborn sickle cell disease screening: the Jamaican experience (1995â€“2006). Journal of Medical Screening, 2007, 14, 117-122.	2.3	59
14	Prevention of conversion to abnormal transcranial <scp>D</scp>oppler with hydroxyurea in sickle cell anemia: A <scp>P</scp>hase III international randomized clinical trial. American Journal of Hematology, 2015, 90, 1099-1105.	4.1	59
15	A doubleâ€•blind, placeboâ€•controlled phase II study of the efficacy and safety of 2,2â€•dimethylbutyrate (HQKâ€•1001), an oral fetal globin inducer, in sickle cell disease. American Journal of Hematology, 2014, 89, 709-713.	4.1	54
16	Caesarean myomectomy-a safe procedure. A retrospective case controlled study. Journal of Obstetrics and Gynaecology, 1999, 19, 139-141.	0.9	51
17	Protein kinetic differences between children with edematous and nonedematous severe childhood undernutrition in the fed and postabsorptive states. American Journal of Clinical Nutrition, 2005, 82, 792-800.	4.7	51
18	Venous incompetence, poverty and lactate dehydrogenase in Jamaica are important predictors of leg ulceration in sickle cell anaemia. British Journal of Haematology, 2008, 142, 119-125.	2.5	48

#	Article	IF	Citations
19	Prevention of striae gravidarum with cocoa butter cream. International Journal of Gynecology and Obstetrics, 2010, 108, 65-68.	2.3	47
20	Utility of WHOQOL-BREF in measuring quality of life in Sickle Cell Disease. Health and Quality of Life Outcomes, 2009, 7, 75.	2.4	46
21	Determining Glomerular Filtration Rate in Homozygous Sickle Cell Disease: Utility of Serum Creatinine Based Estimating Equations. PLoS ONE, 2013, 8, e69922.	2.5	44
22	Relation between liver fat content and the rate of VLDL apolipoprotein B-100 synthesis in children with protein-energy malnutrition1â€“3. American Journal of Clinical Nutrition, 2005, 81, 1126-1132.	4.7	43
23	Lipid kinetic differences between children with kwashiorkor and those with marasmus. American Journal of Clinical Nutrition, 2006, 83, 1283-1288.	4.7	43
24	Depression and loneliness in Jamaicans with sickle cell disease. BMC Psychiatry, 2010, 10, 40.	2.6	43
25	Stroke recurrence in children with sickle cell disease treated with hydroxyurea following first clinical stroke. American Journal of Hematology, 2011, 86, 846-850.	4.1	43
26	Mortality, Asthma, Smoking and Acute Chest Syndrome in Young Adults with Sickle Cell Disease. Lung, 2013, 191, 95-100.	3.3	43
27	Addressing psychological resilience during the coronavirus disease 2019 pandemic: a rapid review. Current Opinion in Psychiatry, 2021, 34, 29-35.	6.3	41
28	Excess Risk of Maternal Death from Sickle Cell Disease in Jamaica: 1998â€“2007. PLoS ONE, 2011, 6, e26281.	2.5	40
29	Operative morbidity and reproductive outcome in secondary myomectomy: a prospective cohort study. Human Reproduction, 2002, 17, 2967-2971.	0.9	37
30	An Investigation of the Antioxidant Capacity in Extracts from Moringa oleifera Plants Grown in Jamaica. Plants, 2017, 6, 48.	3.5	37
31	Plasma Concentration of Platelet-Derived Microparticles Is Related to Painful Vaso-Occlusive Phenotype Severity in Sickle Cell Anemia. PLoS ONE, 2014, 9, e87243.	2.5	36
32	Blood Viscosity and the Expression of Inflammatory and Adhesion Markers in Homozygous Sickle Cell Disease Subjects with Chronic Leg Ulcers. PLoS ONE, 2013, 8, e68929.	2.5	34
33	Dietary Protein, Growth and Urea Kinetics in Severely Malnourished Children and During Recovery. Journal of Nutrition, 1999, 129, 969-979.	2.9	33
34	Prevalence and predictors of microalbuminuria in Jamaican children with sickle cell disease. Archives of Disease in Childhood, 2011, 96, 1135-1139.	1.9	33
35	Group I mGlu receptors potentiate synaptosomal [3H]glutamate release independently of exogenously applied arachidonic acid. Neuropharmacology, 1999, 38, 477-485.	4.1	32
36	Testosterone replacement therapy does not promote priapism in hypogonadal men with sickle cell disease: 12â€•month safety report. Andrology, 2013, 1, 576-582.	3.5	32

#	Article	IF	Citations
37	Discrepancies between clinical and postmortem diagnoses in Jamaica: a study from the University Hospital of the West Indies. Journal of Clinical Pathology, 2004, 57, 980-985.	2.0	31
38	The impact of induced labour on postpartum blood loss. Journal of Obstetrics and Gynaecology, 2004, 24, 12-15.	0.9	31
39	A doseâ€escalation phase IIa study of 2,2â€dimethylbutyrate (HQKâ€1001), an oral fetal globin inducer, in sickle cell disease. American Journal of Hematology, 2013, 88, E255-60.	4.1	31
40	Sulfur amino acid metabolism in children with severe childhood undernutrition: methionine kinetics. American Journal of Clinical Nutrition, 2006, 84, 1400-1405.	4.7	30
41	A phase 1/2 trial of HQKâ€1001, an oral fetal globin inducer, in sickle cell disease. American Journal of Hematology, 2012, 87, 1017-1021.	4.1	30
42	Increased rates of body dissatisfaction, depressive symptoms, and suicide attempts in Jamaican teens with sickle cell disease. Pediatric Blood and Cancer, 2016, 63, 2159-2166.	1.5	30
43	Higher Rates of Hemolysis Are Not Associated with Albuminuria in Jamaicans with Sickle Cell Disease. PLoS ONE, 2011, 6, e18863.	2.5	29
44	Examination of the management of traumatic brain injury in the developing and developed world: focus on resource utilization, protocols, and practices that alter outcome. Journal of Neurosurgery, 2008, 109, 433-438.	1.6	28
45	Dietary cysteine is used more efficiently by children with severe acute malnutrition with edema compared with those without edema. American Journal of Clinical Nutrition, 2012, 95, 84-90.	4.7	28
46	Predictors of renal function progression in adults with homozygous sickle cell disease. British Journal of Haematology, 2016, 173, 461-468.	2.5	28
47	A trial comparing the use of rectal misoprostol plus perivascular vasopressin with perivascular vasopressin alone to decrease myometrial bleeding at the time of abdominal myomectomy. Fertility and Sterility, 2013, 100, 1044-1049.	1.0	27
48	Quality of life in patients with sickle cell disease in Jamaica: rural-urban differences. Rural and Remote Health, 2008, 8, 890.	0.5	27
49	Developmental origins of cardiovascular risk in Jamaican children: The Vulnerable Windows Cohort Study. British Journal of Nutrition, 2010, 104, 1026-1033.	2.3	26
50	Knowledge and health beliefs of Jamaican adolescents with sickle cell disease. Pediatric Blood and Cancer, 2011, 57, 1044-1048.	1.5	26
51	Management of diabetes mellitus in three settings in Jamaica. Revista Panamericana De Salud Publica/Pan American Journal of Public Health, 2001, 9, 65-72.	1.1	26
52	Correlation between transvaginal ultrasound measured endometrial thickness and histopathological findings in Afro-Caribbean Jamaican women with postmenopausal bleeding. Journal of Obstetrics and Gynaecology, 2004, 24, 568-572.	0.9	25
53	Acute Pyelonephritis in Pregnancy: A Retrospective Descriptive Hospital Based-Study. ISRN Obstetrics & Gynecology, 2012, 2012, 1-6.	1.2	25
54	Cystatin C: A useful marker of glomerulopathy in sickle cell disease?. Blood Cells, Molecules, and Diseases, 2015, 54, 65-70.	1.4	25

#	Article	IF	Citations
55	Socio-environmental exposures and health outcomes among persons with sickle cell disease. PLoS ONE, 2017, 12, e0175260.	2.5	25
56	Microalbuminuria, renal function and waist:hip ratio in black hypertensive Jamaicans. Journal of Human Hypertension, 1998, 12, 221-227.	2.2	23
57	Edematous severe acute malnutrition is characterized by hypomethylation of DNA. Nature Communications, 2019, 10, 5791.	12.8	23
58	Seizures in the Jamaica cohort study of sickle cell disease. British Journal of Haematology, 2010, 151, 265-272.	2.5	22
59	Nutrition and sickle cell disease. Comptes Rendus - Biologies, 2013, 336, 159-163.	0.2	22
60	Impact of a Comprehensive Sickle Cell Center on Early Childhood Mortality in a Developing Country: The Jamaican Experience. Journal of Pediatrics, 2015, 167, 702-705.e1.	1.8	22
61	Methods for measuring glutathione concentration and rate of synthesis. Current Opinion in Clinical Nutrition and Metabolic Care, 2000, 3, 385-390.	2.5	21
62	Sulfur amino acid metabolism in children with severe childhood undernutrition: cysteine kinetics. American Journal of Clinical Nutrition, 2006, 84, 1393-1399.	4.7	21
63	Hydroxyurea use in prevention of stroke recurrence in children with sickle cell disease in a developing country: A cost effectiveness analysis. Pediatric Blood and Cancer, 2015, 62, 1862-1864.	1.5	21
64	Renal function in adult Jamaicans with homozygous sickle cell disease. Hematology, 2015, 20, 422-428.	1.5	21
65	A Retrospective Analysis of the Significance of Haemoglobin SS and SC in Disease Outcome in Patients With Sickle Cell Disease and Dengue Fever. EBioMedicine, 2015, 2, 937-941.	6.1	18
66	Response of splanchnic and whole-body leucine kinetics to treatment of children with edematous protein-energy malnutrition accompanied by infection. American Journal of Clinical Nutrition, 2002, 76, 633-640.	4.7	17
67	Prevalence of pneumococcal polysaccharide vaccine administration and incidence of invasive pneumococcal disease in children in Jamaica aged over 4 years with sickle cell disease diagnosed by newborn screening. Annals of Tropical Paediatrics, 2009, 29, 197-202.	1.0	17
68	Morinda citrifolia (Noni) as an Anti-Inflammatory Treatment in Women with Primary Dysmenorrhoea: A Randomised Double-Blind Placebo-Controlled Trial. Obstetrics and Gynecology International, 2013, 2013, 1-6.	1.3	17
69	Asymptomatic bacteriuria in sickle cell disease: a cross-sectional study. BMC Infectious Diseases, 2006, 6, 46.	2.9	16
70	Urea kinetics varies in Jamaican women and men in relation to adiposity, lean body mass and protein intake. European Journal of Clinical Nutrition, 1997, 51, 107-115.	2.9	15
71	Contribution of pulmonary diseases to COVID-19 mortality in a diverse urban community of New York. Chronic Respiratory Disease, 2021, 18, 147997312098680.	2.4	15
72	Iron deficiency anaemia in Jamaican children, aged 1-5 years, with sickle cell disease. West Indian Medical Journal, 2005, 54, 292-6.	0.4	15

#	Article	IF	Citations
73	Laparoscopic cholecystectomy for chronic cholecystitis in Jamaican patients with sickle cell disease: preliminary experience. West Indian Medical Journal, 2006, 55, 22-4.	0.4	15
74	Validation of the SF-36 in Jamaicans with sickle-cell disease. Psychology, Health and Medicine, 2009, 14, 606-618.	2.4	14
75	Effects of randomized supplementation of methionine or alanine on cysteine and glutathione production during the early phase of treatment of children with edematous malnutrition. American Journal of Clinical Nutrition, 2014, 99, 1052-1058.	4.7	14
76	Is testosterone deficiency a possible risk factor for priapism associated with sickle-cell disease?. International Urology and Nephrology, 2015, 47, 47-52.	1.4	14
77	Transcranial Doppler velocity among Jamaican children with sickle cell anaemia: determining the significance of haematological values and nutrition. British Journal of Haematology, 2018, 181, 242-251.	2.5	14
78	Glutathione S-transferase polymorphisms may be associated with risk of oedematous severe childhood malnutrition. British Journal of Nutrition, 2006, 96, 243-248.	2.3	13
79	Risky behaviours of Jamaican adolescents with sickle cell disease. Hematology, 2014, 19, 373-379.	1.5	13
80	The transfer of 15N from urea to lysine in the human infant. British Journal of Nutrition, 2000, 83, 505-12.	2.3	13
81	Component structure of the SF-36 in Jamaicans with sickle cell disease. West Indian Medical Journal, 2007, 56, 491-7.	0.4	13
82	Vasopressin as a hemostatic and dissection aid at vaginal hysterectomy. International Journal of Gynecology and Obstetrics, 2004, 86, 65-66.	2.3	12
83	Glycine production in severe childhood undernutrition. American Journal of Clinical Nutrition, 2006, 84, 143-149.	4.7	12
84	Locus of control, depression and quality of life among persons with sickle cell disease in Jamaica. Psychology, Health and Medicine, 2013, 18, 451-460.	2.4	12
85	Newborn Screening for Sickle Cell Disease in Jamaica: A Review â€“ Past, Present and Future. West Indian Medical Journal, 2014, 63, 147-50.	0.4	11
86	Age as a Predictive Factor of Mammographic Breast Density in Jamaican Women. Clinical Radiology, 2002, 57, 472-476.	1.1	10
87	Hydroxycarbamide treatment reduces transcranial Doppler velocity in the absence of transfusion support in children with sickle cell anaemia, elevated transcranial Doppler velocity, and cerebral vasculopathy: the EXTEND trial. British Journal of Haematology, 2021, 195, 612-620.	2.5	10
88	A clinical audit of the quality of care of hypertension in general practice. West Indian Medical Journal, 2005, 54, 176-80.	0.4	10
89	Blood pressure in Jamaican children: relationship to body size and composition. West Indian Medical Journal, 1999, 48, 61-8.	0.4	10
90	Moyamoya syndrome in sickle cell anaemia: a cause of recurrent stroke. BMJ Case Reports, 2014, 2014, bcr2014203727-bcr2014203727.	0.5	9

#	Article	IF	Citations
91	Mammographic referral patterns for two breast imaging units in Jamaica. West Indian Medical Journal, 2007, 56, 159-62.	0.4	9
92	EXpanding Treatment for Existing Neurological Disease (EXTEND): An Open-Label Phase II Clinical Trial of Hydroxyurea Treatment in Sickle Cell Anemia. JMIR Research Protocols, 2016, 5, e185.	1.0	9
93	Clinical Factors Associated With Morbidity and Mortality in Patients Admitted with Sickle Cell Disease. West Indian Medical Journal, 2014, 63, 711-6.	0.4	9
94	Sociocultural deterrents to mammographic screening in Jamaica. West Indian Medical Journal, 2009, 58, 28-32.	0.4	9
95	A comparative study of the quality and availability of health information used to facilitate cost burden analysis of diabetes and hypertension in the Caribbean. West Indian Medical Journal, 2008, 57, 383-92.	0.4	9
96	The Possible Impact of Teachers and School Nurses on the Lives of Children Living With Sickle Cell Disease. Journal of School Health, 2011, 81, 219-222.	1.6	8
97	Bone mineral density in Jamaican men on androgen deprivation therapy for prostate cancer. Infectious Agents and Cancer, 2011, 6, S7.	2.6	8
98	Overactive bladder in adults with sickle cell disease. Neurourology and Urodynamics, 2016, 35, 642-646.	1.5	8
99	Elements in human placentae in Jamaica. West Indian Medical Journal, 2010, 59, 479-85.	0.4	8
100	Compliance with intramuscular penicillin prophylaxis in children with sickle cell disease in Jamaica. West Indian Medical Journal, 2011, 60, 177-80.	0.4	8
101	Polymorphisms in genes involved in folate metabolism as risk factors for oedematous severe childhood malnutrition: a hypothesis-generating study. Annals of Tropical Paediatrics, 2006, 26, 107-114.	1.0	7
102	Relationship between acute chest syndrome and the sympatho-vagal balance in adults with hemoglobin SS disease; a case control study. Clinical Hemorheology and Microcirculation, 2013, 53, 231-238.	1.7	7
103	Vitamin D levels are low in adult patients with sickle cell disease in Jamaica and West Africa. BMC Hematology, 2014, 14, 12.	2.6	7
104	Supplementation with Aromatic Amino Acids Improves Leucine Kinetics but Not Aromatic Amino Acid Kinetics in Infants with Infection, Severe Malnutrition, and Edema. Journal of Nutrition, 2004, 134, 3004-3010.	2.9	6
105	Depo medroxyprogesterone acetate (DMPA) therapy for uterine myomata prior to surgery. International Journal of Gynecology and Obstetrics, 2004, 85, 174-176.	2.3	6
106	Arginine flux and intravascular nitric oxide synthesis in severe childhood undernutrition. American Journal of Clinical Nutrition, 2007, 86, 1024-1031.	4.7	6
107	Associations amongst disease severity, religious coping and depression in a cohort of Jamaicans with sickle-cell disease. Mental Health, Religion and Culture, 2014, 17, 937-945.	0.9	6
108	Dietary Supplementation with Aromatic Amino Acids Increases Protein Synthesis in Children with Severe Acute Malnutrition. Journal of Nutrition, 2014, 144, 660-666.	2.9	6

#	Article	IF	Citations
109	Prostate Cancer Screening in Jamaica: Results of the Largest National Screening Clinic. Journal of Cancer Epidemiology, 2016, 2016, 1-4.	1.1	6
110	Pathological factors affecting gastric adenocarcinoma survival in a Caribbean population from 2000-2010. World Journal of Gastrointestinal Surgery, 2014, 6, 94.	1.5	6
111	Association between exposure/non-exposure to the mandatory seat belt law with regards to compliance in vehicle accident victims: a hospital review. West Indian Medical Journal, 2007, 56, 236-9.	0.4	6
112	Tyrosine requirement during the rapid catch-up growth phase of recovery from severe childhood undernutrition. British Journal of Nutrition, 2010, 104, 1174-1180.	2.3	5
113	The effect of feto-maternal size and childhood growth on left ventricular mass and arterial stiffness in Afro-Caribbean children. Journal of Human Hypertension, 2011, 25, 457-464.	2.2	5
114	Gangrene of the Digits of the Right Lower Limb in a Patient with Homozygous Sickle Cell Disease and Ulcerative Colitis. Clinics and Practice, 2014, 4, 7-9.	1.4	5
115	Population Reference Values and Prevalence Rates following Universal Screening for Subclinical Hypothyroidism during Pregnancy of an Afro-Caribbean Cohort. European Thyroid Journal, 2014, 3, 234-9.	2.4	5
116	Intrathoracic and pelvic extramedullary haematopoiesis in sickle cell disease: a case report. West Indian Medical Journal, 2007, 56, 540-3.	0.4	5
117	Cardiovascular disease risk factors in menopausal Jamaican black women after hysterectomy and bilateral oophorectomy: an observational study. West Indian Medical Journal, 2010, 59, 625-32.	0.4	5
118	Relationship between birth weight and urea kinetics in children. European Journal of Clinical Nutrition, 2006, 60, 197-202.	2.9	4
119	CARESTâ€"Multilingual Regional Integration for Health Promotion and Research on Sickle Cell Disease and Thalassemia. American Journal of Public Health, 2016, 106, 851-853.	2.7	4
120	Impact of the National Health Fund policy on hormone treatment for prostate cancer in Jamaica. Revista Panamericana De Salud Publica/Pan American Journal of Public Health, 2011, 29, 404-408.	1.1	4
121	Quality of care of hypertension in three clinical settings in Jamaica. West Indian Medical Journal, 2000, 49, 220-5.	0.4	4
122	A preliminary examination of the effects of genetic variants of redox enzymes on susceptibility to oedematous malnutrition and on percentage cytotoxicity in response to oxidative stressin vitro. Annals of Tropical Paediatrics, 2011, 31, 27-36.	1.0	3
123	Nutritional Repletion of Children with Severe Acute Malnutrition Does Not Affect VLDL Apolipoprotein B-100 Synthesis Rate. Journal of Nutrition, 2012, 142, 931-935.	2.9	3
124	A case series of cholecystectomy in Jamaican sickle cell disease patients - The need for a new strategy. Annals of Medicine and Surgery, 2017, 15, 37-42.	1.1	3
125	Exploring putative genetic determinants of inter-individual phenotypic heterogeneity in sickle cell disease: A cross-sectional Jamaican cohort-based study. Blood Cells, Molecules, and Diseases, 2018, 73, 1-8.	1.4	3
126	A Survey of the Pain Management of Acute Painful Crisis among Patients with Sickle Cell Disease at Two Centres in Jamaica. West Indian Medical Journal, 2014, 63, 252-7.	0.4	3

#	Article	IF	Citations
127	HIV infection seroprevalence and risk factor study among pregnant women attending the Antenatal Clinic at the University Hospital of the West Indies, Kingston, Jamaica. West Indian Medical Journal, 2002, 51, 80-3.	0.4	3
128	Utility of rapid staining of fine needle aspiration smears at the University Hospital of the West Indies. West Indian Medical Journal, 2003, 52, 34-6.	0.4	3
129	Effect of the oral contraceptive pill on patients undergoing controlled ovarian hyperstimulation. West Indian Medical Journal, 2004, 53, 39-43.	0.4	3
130	The prevalence of positive anticardiolipin antibody in Jamaican women with spontaneous abortion and the correlation with past clinical history. Journal of Obstetrics and Gynaecology, 2004, 24, 452-454.	0.9	2
131	Bilateral psoas abscesses and vertebral osteomyelitis in a patient with sickle cell disease. BMJ Case Reports, 2010, 2010, bcr0120102642-bcr0120102642.	0.5	2
132	Splenic enlargement in adults with homozygous sickle cell disease: the Jamaican experience. Hematology, 2013, 18, 46-49.	1.5	2
133	A cross-sectional clinic-based study exploring whether variants within the glutathione S-transferase, haptoglobin and uridine 5â€²-diphospho-glucuronosyltransferase 1A1 genes are associated with interindividual phenotypic variation in sickle cell anaemia in. European Journal of Haematology, 2018, 100, 147-153.	2.2	2
134	Improving patient outcomes with inguinal hernioplastyâ€”local anaesthesia versus local anaesthesia and conscious sedation: a randomized controlled trial. Hernia: the Journal of Hernias and Abdominal Wall Surgery, 2019, 23, 561-567.	2.0	2
135	The Moderating Effect of Physical Activity on the Relationship between Sleep and Emotional Distress and the Difference between Blacks and Whites: A Secondary Data Analysis Using the National Health Interview Survey from 2005â€“2015. International Journal of Environmental Research and Public Health, 2021, 18, 1718.	2.6	2
136	Local Anesthesia Versus Local Anesthesia and Conscious Sedation for Inguinal Hernioplasty: Protocol of a Randomized Controlled Trial. JMIR Research Protocols, 2017, 6, e20.	1.0	2
137	Diagnostic Accuracy of Spot and Timed Measurements of Urinary Albumin Concentration to Determine Microalbuminuria in Sickle Cell Disease. West Indian Medical Journal, 2013, 62, 808-16.	0.4	2
138	Bone Mineral Density in Jamaican Black Women after Hysterectomy and Bilateral Oophorectomy: An Observational Study. West Indian Medical Journal, 2014, 62, 593-8.	0.4	2
139	Lupus anticoagulant in Jamaican primiparae and the clinical significance in asymptomatic patients. West Indian Medical Journal, 1999, 48, 126-8.	0.4	2
140	Preliminary report on the validation of a questionnaire measuring patient satisfaction with services at the sickle cell unit in Jamaica. West Indian Medical Journal, 2009, 58, 331-40.	0.4	2
141	A comparison of two methods of labor induction with vaginal misoprostol. International Journal of Gynecology and Obstetrics, 2003, 80, 271-277.	2.3	1
142	Anticardiolipin, other antiphospholipid antibody tests and diagnosis of the antiphospholipid syndrome. Human Antibodies, 2003, 12, 63-66.	1.5	1
143	Congenital abnormalities at a tertiary center in Jamaica: An 18-month maternalâ€“fetal medicine experience. Journal of Maternal-Fetal and Neonatal Medicine, 2012, 25, 687-691.	1.5	1
144	Effectiveness of antenatal screening for sickle cell trait: the impact on womenâ€™s self-report of sickle cell trait status. Pathogens and Global Health, 2012, 106, 55-59.	2.3	1

#	Article	IF	Citations
145	Computed Tomography in the evaluation of seizures at the University Hospital of the West Indies. West Indian Medical Journal, 2001, 50, 151-4.	0.4	1
146	Female urinary retention at the University Hospital of the West Indies challenging diagnosis. West Indian Medical Journal, 2011, 60, 102.	0.4	1
147	Newborn Screening in Jamaica. American Journal of Preventive Medicine, 2012, 42, e101.	3.0	0
148	Hydroxyurea for Primary Stroke Prevention: The time draweth nigh. Pediatric Blood and Cancer, 2019, 66, e27536.	1.5	0
149	Oesophageal foreign bodies at the University Hospital of the West Indies. West Indian Medical Journal, 2005, 54, 47-50.	0.4	0
150	Dietary cysteine is utilized more efficiently by children with edematous severe childhood undernutrition compared to those with nonâ€edematous severe childhood undernutrition during nutritional rehabilitation. FASEB Journal, 2011, 25, 983.1.	0.5	0
151	Dietary supplementation with aromatic amino acids improves net protein synthesis in children with severe acute malnutrition during hospitalization. FASEB Journal, 2012, 26, 42.2.	0.5	0
152	Obstetric Outcomes of an Afro - Caribbean Cohort Following Universal Screening and Treatment of Subclinical Hypothyroidism. West Indian Medical Journal, 2015, 65, 78-82.	0.4	0
153	Delays in presentations of stroke patients at the University Hospital of the West Indies. West Indian Medical Journal, 2009, 58, 341-6.	0.4	0