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	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
2	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
3	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
4	Addressing psychological resilience during the coronavirus disease 2019 pandemic: a rapid review. Current Opinion in Psychiatry, 2021, 34, 29-35.	6.3	41
5	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
6	Edematous severe acute malnutrition is characterized by hypomethylation of DNA. Nature Communications, 2019, 10, 5791.	12.8	23
7	Hydroxyurea for Primary Stroke Prevention: The time draweth nigh. Pediatric Blood and Cancer, 2019, 66, e27536.	1.5	0
8	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
9	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
10	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
11	Variant-aware saturating mutagenesis using multiple Cas9 nucleases identifies regulatory elements at trait-associated loci. Nature Genetics, 2017, 49, 625-634.	21.4	96
12	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
13	An Investigation of the Antioxidant Capacity in Extracts from Moringa oleifera Plants Grown in Jamaica. Plants, 2017, 6, 48.	3.5	37
14	Socio-environmental exposures and health outcomes among persons with sickle cell disease. PLoS ONE, 2017, 12, e0175260.	2.5	25
15	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
16	Prostate Cancer Screening in Jamaica: Results of the Largest National Screening Clinic. Journal of Cancer Epidemiology, 2016, 2016, 1-4.	1.1	6
17	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
18	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

#	Article	IF	Citations
19	Predictors of renal function progression in adults with homozygous sickle cell disease. British Journal of Haematology, 2016, 173, 461-468.	2.5	28
20	Overactive bladder in adults with sickle cell disease. Neurourology and Urodynamics, 2016, 35, 642-646.	1.5	8
21	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
22	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
23	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
24	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
25	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
26	Renal function in adult Jamaicans with homozygous sickle cell disease. Hematology, 2015, 20, 422-428.	1.5	21
27	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
28	Cystatin C: A useful marker of glomerulopathy in sickle cell disease?. Blood Cells, Molecules, and Diseases, 2015, 54, 65-70.	1.4	25
29	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
30	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
31	Risky behaviours of Jamaican adolescents with sickle cell disease. Hematology, 2014, 19, 373-379.	1.5	13
32	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
33	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
34	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
35	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
36	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
37	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
38	Moyamoya syndrome in sickle cell anaemia: a cause of recurrent stroke. BMJ Case Reports, 2014, 2014, bcr2014203727-bcr2014203727.	0.5	9
39	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
40	Pathological factors affecting gastric adenocarcinoma survival in a Caribbean population from 2000-2010. World Journal of Gastrointestinal Surgery, 2014, 6, 94.	1.5	6
41	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
42	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
43	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
44	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
45	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
46	Mortality, Asthma, Smoking and Acute Chest Syndrome in Young Adults with Sickle Cell Disease. Lung, 2013, 191, 95-100.	3.3	43
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	Nutrition and sickle cell disease. Comptes Rendus - Biologies, 2013, 336, 159-163.	0.2	22
49	Splenic enlargement in adults with homozygous sickle cell disease: the Jamaican experience. Hematology, 2013, 18, 46-49.	1.5	2
50	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
51	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
52	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
53	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
54	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

#	Article	IF	Citations
55	Determining Glomerular Filtration Rate in Homozygous Sickle Cell Disease: Utility of Serum Creatinine Based Estimating Equations. PLoS ONE, 2013, 8, e69922.	2.5	44
56	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
57	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
58	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
59	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
60	Acute Pyelonephritis in Pregnancy: A Retrospective Descriptive Hospital Based-Study. ISRN Obstetrics & Gynecology, 2012, 2012, 1-6.	1.2	25
61	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
62	Newborn Screening in Jamaica. American Journal of Preventive Medicine, 2012, 42, e101.	3.0	0
63	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
64	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
65	Higher Rates of Hemolysis Are Not Associated with Albuminuria in Jamaicans with Sickle Cell Disease. PLoS ONE, 2011, 6, e18863.	2.5	29
66	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
67	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
68	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
69	Bone mineral density in Jamaican men on androgen deprivation therapy for prostate cancer. Infectious Agents and Cancer, 2011, 6, S7.	2.6	8
70	Knowledge and health beliefs of Jamaican adolescents with sickle cell disease. Pediatric Blood and Cancer, 2011, 57, 1044-1048.	1.5	26
71	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
72	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

#	Article	IF	Citations
73	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
74	Prevalence and predictors of microalbuminuria in Jamaican children with sickle cell disease. Archives of Disease in Childhood, 2011, 96, 1135-1139.	1.9	33
75	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
76	Excess Risk of Maternal Death from Sickle Cell Disease in Jamaica: 1998â€“2007. PLoS ONE, 2011, 6, e26281.	2.5	40
77	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
78	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
79	Female urinary retention at the University Hospital of the West Indies challenging diagnosis. West Indian Medical Journal, 2011, 60, 102.	0.4	1
80	Compliance with intramuscular penicillin prophylaxis in children with sickle cell disease in Jamaica. West Indian Medical Journal, 2011, 60, 177-80.	0.4	8
81	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
82	Depression and loneliness in Jamaicans with sickle cell disease. BMC Psychiatry, 2010, 10, 40.	2.6	43
83	Seizures in the Jamaica cohort study of sickle cell disease. British Journal of Haematology, 2010, 151, 265-272.	2.5	22
84	Bilateral psoas abscesses and vertebral osteomyelitis in a patient with sickle cell disease. BMJ Case Reports, 2010, 2010, bcr0120102642-bcr0120102642.	0.5	2
85	Developmental origins of cardiovascular risk in Jamaican children: The Vulnerable Windows Cohort Study. British Journal of Nutrition, 2010, 104, 1026-1033.	2.3	26
86	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
87	Prevention of striae gravidarum with cocoa butter cream. International Journal of Gynecology and Obstetrics, 2010, 108, 65-68.	2.3	47
88	Elements in human placentae in Jamaica. West Indian Medical Journal, 2010, 59, 479-85.	0.4	8
89	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
90	Validation of the SF-36 in Jamaicans with sickle-cell disease. Psychology, Health and Medicine, 2009, 14, 606-618.	2.4	14

#	Article	IF	Citations
91	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
92	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
93	Sociocultural deterrents to mammographic screening in Jamaica. West Indian Medical Journal, 2009, 58, 28-32.	0.4	9
94	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
95	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
96	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
97	Protein metabolism in severe childhood malnutrition. Annals of Tropical Paediatrics, 2008, 28, 87-101.	1.0	72
98	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
99	Quality of life in patients with sickle cell disease in Jamaica: rural-urban differences. Rural and Remote Health, 2008, 8, 890.	0.5	27
100	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
101	Newborn sickle cell disease screening: the Jamaican experience (1995â€“2006). Journal of Medical Screening, 2007, 14, 117-122.	2.3	59
102	Albuminuria and Renal Function in Homozygous Sickle Cell Disease. Archives of Internal Medicine, 2007, 167, 701.	3.8	85
103	Arginine flux and intravascular nitric oxide synthesis in severe childhood undernutrition. American Journal of Clinical Nutrition, 2007, 86, 1024-1031.	4.7	6
104	Mammographic referral patterns for two breast imaging units in Jamaica. West Indian Medical Journal, 2007, 56, 159-62.	0.4	9
105	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
106	Component structure of the SF-36 in Jamaicans with sickle cell disease. West Indian Medical Journal, 2007, 56, 491-7.	0.4	13
107	Intrathoracic and pelvic extramedullary haematopoiesis in sickle cell disease: a case report. West Indian Medical Journal, 2007, 56, 540-3.	0.4	5
108	Sulfur amino acid metabolism in children with severe childhood undernutrition: cysteine kinetics. American Journal of Clinical Nutrition, 2006, 84, 1393-1399.	4.7	21

#	Article	IF	Citations
109	Sulfur amino acid metabolism in children with severe childhood undernutrition: methionine kinetics. American Journal of Clinical Nutrition, 2006, 84, 1400-1405.	4.7	30
110	Glycine production in severe childhood undernutrition. American Journal of Clinical Nutrition, 2006, 84, 143-149.	4.7	12
111	Lipid kinetic differences between children with kwashiorkor and those with marasmus. American Journal of Clinical Nutrition, 2006, 83, 1283-1288.	4.7	43
112	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
113	Asymptomatic bacteriuria in sickle cell disease: a cross-sectional study. BMC Infectious Diseases, 2006, 6, 46.	2.9	16
114	Relationship between birth weight and urea kinetics in children. European Journal of Clinical Nutrition, 2006, 60, 197-202.	2.9	4
115	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
116	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
117	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
118	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
119	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
120	A clinical audit of the quality of care of hypertension in general practice. West Indian Medical Journal, 2005, 54, 176-80.	0.4	10
121	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
122	Oesophageal foreign bodies at the University Hospital of the West Indies. West Indian Medical Journal, 2005, 54, 47-50.	0.4	0
123	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
124	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
125	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
126	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

#	Article	IF	Citations
127	The impact of induced labour on postpartum blood loss. Journal of Obstetrics and Gynaecology, 2004, 24, 12-15.	0.9	31
128	Depo medroxyprogesterone acetate (DMPA) therapy for uterine myomata prior to surgery. International Journal of Gynecology and Obstetrics, 2004, 85, 174-176.	2.3	6
129	Vasopressin as a hemostatic and dissection aid at vaginal hysterectomy. International Journal of Gynecology and Obstetrics, 2004, 86, 65-66.	2.3	12
130	Effect of the oral contraceptive pill on patients undergoing controlled ovarian hyperstimulation. West Indian Medical Journal, 2004, 53, 39-43.	0.4	3
131	A comparison of two methods of labor induction with vaginal misoprostol. International Journal of Gynecology and Obstetrics, 2003, 80, 271-277.	2.3	1
132	Anticardiolipin, other antiphospholipid antibody tests and diagnosis of the antiphospholipid syndrome. Human Antibodies, 2003, 12, 63-66.	1.5	1
133	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
134	Operative morbidity and reproductive outcome in secondary myomectomy: a prospective cohort study. Human Reproduction, 2002, 17, 2967-2971.	0.9	37
135	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
136	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
137	Age as a Predictive Factor of Mammographic Breast Density in Jamaican Women. Clinical Radiology, 2002, 57, 472-476.	1.1	10
138	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
139	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
140	Glutathione in disease. Current Opinion in Clinical Nutrition and Metabolic Care, 2001, 4, 65-71.	2.5	102
141	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
142	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
143	Methods for measuring glutathione concentration and rate of synthesis. Current Opinion in Clinical Nutrition and Metabolic Care, 2000, 3, 385-390.	2.5	21
144	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

#	Article	IF	Citations
145	The transfer of 15N from urea to lysine in the human infant. British Journal of Nutrition, 2000, 83, 505-12.	2.3	13
146	Quality of care of hypertension in three clinical settings in Jamaica. West Indian Medical Journal, 2000, 49, 220-5.	0.4	4
147	Dietary Protein, Growth and Urea Kinetics in Severely Malnourished Children and During Recovery. Journal of Nutrition, 1999, 129, 969-979.	2.9	33
148	Caesarean myomectomy-a safe procedure. A retrospective case controlled study. Journal of Obstetrics and Gynaecology, 1999, 19, 139-141.	0.9	51
149	Group I mGlu receptors potentiate synaptosomal [3H]glutamate release independently of exogenously applied arachidonic acid. Neuropharmacology, 1999, 38, 477-485.	4.1	32
150	Blood pressure in Jamaican children: relationship to body size and composition. West Indian Medical Journal, 1999, 48, 61-8.	0.4	10
151	Lupus anticoagulant in Jamaican primiparae and the clinical significance in asymptomatic patients. West Indian Medical Journal, 1999, 48, 126-8.	0.4	2
152	Microalbuminuria, renal function and waist:hip ratio in black hypertensive Jamaicans. Journal of Human Hypertension, 1998, 12, 221-227.	2.2	23
153	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