# James C Barton

List of Publications by Year
in descending order

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| 262 | 7,091 | 66343 | 76900 |
|---|---|---|---|
| papers | citations | 42 h-index | 74 g-index |

| 267 | 267 | 267 | 3929 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Hemochromatosis and Iron-Overload Screening in a Racially Diverse Population. New England Journal of Medicine, 2005, 352, 1769-1778. | 27.0 | 662 |
| 2 | Hereditary Hemochromatosis. JAMA - Journal of the American Medical Association, 1998, 280, 172. | 7.4 | 253 |
| 3 | Management of Hemochromatosis. Annals of Internal Medicine, 1998, 129, 932. | 3.9 | 223 |
| 4 | A survey of 2,851 patients with hemochromatosis:. American Journal of Medicine, 1999, 106, 619-624. | 1.5 | 207 |
| 5 | Haemochromatosis. Lancet, The, 2007, 370, 1855-1860. | 13.7 | 178 |
| 6 | Serum Ferritin Level Predicts Advanced Hepatic Fibrosis among U.S. Patients with Phenotypic Hemochromatosis. Annals of Internal Medicine, 2003, 138, 627. | 3.9 | 167 |
| 7 | Genetic and Clinical Description of Hemochromatosis Probands and Heterozygotes: Evidence That Multiple Genes Linked to the Major Histocompatibility Complex Are Responsible for Hemochromatosis. Blood Cells, Molecules, and Diseases, 1997, 23, 135-145. | 1.4 | 153 |
| 8 | How I treat hemochromatosis. Blood, 2010, 116, 317-325. | 1.4 | 152 |
| 9 | Two Novel Missense Mutations of the HFE Gene (I105T and G93R) and Identification of the S65C Mutation in Alabama Hemochromatosis Probands. Blood Cells, Molecules, and Diseases, 1999, 25, 147-155. | 1.4 | 150 |
| 10 | Genetic abnormalities and juvenile hemochromatosis: mutations of the HJV gene encoding hemojuvelin. Blood, 2004, 103, 4669-4671. | 1.4 | 133 |
| 11 | Ferroportin 1 (SCL40A1) variant associated with iron overload in African-Americans. Blood Cells, Molecules, and Diseases, 2003, 31, 305-309. | 1.4 | 118 |
| 12 | Hemochromatosis and Iron Overload Screening (HEIRS) Study Design for an Evaluation of 100,000 Primary Care-Based Adults. American Journal of the Medical Sciences, 2003, 325, 53-62. | 1.1 | 118 |
| 13 | Iron overload in African Americans. American Journal of Medicine, 1995, 99, 616-623. | 1.5 | 115 |
| 14 | Calcium Inhibition of Inorganic Iron Absorption in Rats. Gastroenterology, 1983, 84, 90-101. | 1.3 | 99 |
| 15 | HFE gene: Structure, function, mutations, and associated iron abnormalities. Gene, 2015, 574, 179-192. | 2.2 | 97 |
| 16 | Concerns in a primary care population about genetic discrimination by insurers. Genetics in Medicine, 2005, 7, 311-316. | 2.4 | 94 |
| 17 | Acute cryoglobulinemic renal failure after intravenous infusion of gamma globulin. American Journal of Medicine, 1987, 82, 624-629. | 1.5 | 91 |
| 18 | Ultrastructural localization of transferrin, transferrin receptor, and ironâ€binding sites on human placental and duodenal microvilli. British Journal of Haematology, 1985, 60, 81-89. | 2.5 | 88 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | Relationships of Serum Ferritin, Transferrin Saturation, and HFE Mutations and Self-Reported Diabetes in the Hemochromatosis and Iron Overload Screening (HEIRS) Study. Diabetes Care, 2006, 29, 2084-2089. | 8.6 | 85 |
| 20 | A diagnostic approach to hyperferritinemia with a non-elevated transferrin saturation. Journal of Hepatology, 2011, 55, 453-458. | 3.7 | 85 |
| 21 | Peripheral blood erythrocyte parameters in hemochromatosis: Evidence for increased erythrocyte hemoglobin content. Translational Research, 2000, 135, 96-104. | 2.3 | 81 |
| 22 | Biological Variability of Transferrin Saturation and Unsaturated Iron-Binding Capacity. American Journal of Medicine, 2007, 120, 999.e1-999.e7. | 1.5 | 75 |
| 23 | Exome sequencing in HFE C282Y homozygous men with extreme phenotypes identifies a GNPAT variant associated with severe iron overload. Hepatology, 2015, 62, 429-439. | 7.3 | 75 |
| 24 | Initial Screening Transferrin Saturation Values, Serum Ferritin Concentrations, andHFEGenotypes in Whites and Blacks in the Hemochromatosis and Iron Overload Screening Study. Genetic Testing and Molecular Biomarkers, 2005, 9, 231-241. | 1.7 | 69 |
| 25 | Genotypic and phenotypic heterogeneity of African Americans with primary iron overload. Blood Cells, Molecules, and Diseases, 2003, 31, 310-319. | 1.4 | 65 |
| 26 | Effect of Iron Deficiency on Bleomycin-induced Lung Fibrosis in the Hamster. The American Review of Respiratory Disease, 1988, 137, 85-89. | 2.9 | 64 |
| 27 | Ferritin >1000: grand for hemochromatosis screening?. Blood, 2008, 111, 3309-3309. | 1.4 | 63 |
| 28 | Tumor lysis syndrome in nonhematopoietic neoplasms. Cancer, 1989, 64, 738-740. | 4.1 | 61 |
| 29 | Beneficial Effects of Hepatitis in Patients with Acute Myelogenous Leukemia. Annals of Internal Medicine, 1979, 90, 188. | 3.9 | 60 |
| 30 | Hemochromatosis and Vibrio vulnificus Wound Infections. Journal of Clinical Gastroenterology, 2009, 43, 890-893. | 2.2 | 59 |
| 31 | Ultrastructural cytochemistry and radioautography of hemoglobinâ€"iron absorption. Experimental and Molecular Pathology, 1981, 34, 131-144. | 2.1 | 57 |
| 32 | Acute lymphoblastic leukemia in idiopathic refractory sideroblastic anemia: Evidence for a common lymphoid and myeloid progenitor cell. American Journal of Hematology, 1980, 9, 109-115. | 4.1 | 56 |
| 33 | Clinical Manifestations of Hemochromatosis inHFEC282Y Homozygotes Identified by Screening. Canadian Journal of Gastroenterology & Hepatology, 2008, 22, 923-930. | 1.7 | 56 |
| 34 | Factors affecting iron balance. American Journal of Hematology, 1981, 10, 199-225. | 4.1 | 53 |
| 35 | Rheumatoid Arthritis Associated with Expanded Populations of Granular Lymphocytes. Annals of Internal Medicine, 1986, 104, 314. | 3.9 | 52 |
| 36 | Liver Diseases in the Hemochromatosis and Iron Overload Screening Study. Clinical Gastroenterology and Hepatology, 2006, 4, 918-923.e1. | 4.4 | 52 |

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|---|---|---|---|
| 37 | Hemochromatosis: Association of Severity of Iron Overload with Genetic Markers. Blood Cells, Molecules, and Diseases, 1996, 22, 195-204. | 1.4 | 51 |
| 38 | Serum Ferritin and Transferrin Saturation in Asians and Pacific Islanders. Archives of Internal Medicine, 2007, 167, 722. | 3.8 | 51 |
| 39 | Differences in Hepatic Phenotype Between Hemochromatosis Patients With HFE C282Y Homozygosity and Other HFE Genotypes. Journal of Clinical Gastroenterology, 2009, 43, 569-573. | 2.2 | 51 |
| 40 | Hemochromatosis: The genetic disorder of the twentyâ€“first century. Nature Medicine, 1996, 2, 394-395. | 30.7 | 48 |
| 41 | Patient compliance with phlebotomy therapy for iron overload associated with hemochromatosis. American Journal of Gastroenterology, 2003, 98, 2072-2077. | 0.4 | 48 |
| 42 | Hemojuvelin (HJV) mutations in persons of European, Africanâ€American and Asian ancestry with adult onset haemochromatosis. British Journal of Haematology, 2004, 127, 224-229. | 2.5 | 48 |
| 43 | Association of ferroportin Q248H polymorphism with elevated levels of serum ferritin in African Americans in the Hemochromatosis and Iron Overload Screening (HEIRS) Study. Blood Cells, Molecules, and Diseases, 2007, 38, 247-252. | 1.4 | 44 |
| 44 | Increased Risk of Death From Iron Overload Among 422 Treated Probands With HFE Hemochromatosis and Serum Levels of Ferritin Greater Than 1000 Î¼g/L at Diagnosis. Clinical Gastroenterology and Hepatology, 2012, 10, 412-416. | 4.4 | 43 |
| 45 | HLA haplotype A*03-B*07 in hemochromatosis probands with C282Y homozygosity: frequency disparity in men and women and lack of association with severity of iron overload. Blood Cells, Molecules, and Diseases, 2005, 34, 38-47. | 1.4 | 42 |
| 46 | Diabetes inHFEHemochromatosis. Journal of Diabetes Research, 2017, 2017, 1-16. | 2.3 | 42 |
| 47 | Comparison of the Unsaturated Iron-Binding Capacity with Transferrin Saturation as a Screening Test to Detect C282Y Homozygotes for Hemochromatosis in 101 168 Participants in the Hemochromatosis and Iron Overload Screening (HEIRS) Study. Clinical Chemistry, 2005, 51, 1048-1052. | 3.2 | 41 |
| 48 | Chelation therapy for iron overload. Current Gastroenterology Reports, 2007, 9, 74-82. | 2.5 | 41 |
| 49 | Bone Metastases in Malignant Gastrinoma. Gastroenterology, 1986, 91, 1179-1185. | 1.3 | 40 |
| 50 | HFE, SLC40A1, HAMP, HJV, TFR2, and FTL mutations detected by denaturing highâ€performance liquid chromatography after iron phenotyping and HFE C282Y and H63D genotyping in 785 HEIRS Study participants. American Journal of Hematology, 2009, 84, 710-714. | 4.1 | 39 |
| 51 | Serum ferritin concentrations and body iron stores in a multicenter, multiethnic primaryâ€care population. American Journal of Hematology, 2008, 83, 618-626. | 4.1 | 37 |
| 52 | Screening for Iron Overload: Lessons from the HEmochromatosis and IRon Overload Screening (HEIRS) Study. Canadian Journal of Gastroenterology & Hepatology, 2009, 23, 769-772. | 1.7 | 37 |
| 53 | Prevalence of granual lymphocyte proliferation in patients with rheumatoid arthritis and neutropenia. American Journal of Medicine, 1989, 86, 303-307. | 1.5 | 36 |
| 54 | Diagnosis of Hemochromatosis Probands in a Community Hospital. American Journal of Medicine, 1997, 103, 498-503. | 1.5 | 36 |

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| 55 | Determinants and characteristics of mean corpuscular volume and hemoglobin concentration in whiteHFE C282Y homozygotes in the hemochromatosis and iron overload screening study. American Journal of Hematology, 2007, 82, 898-905. | 4.1 | 36 |
| 56 | Transfusion Iron Overload in Adults with Acute Leukemia: Manifestations and Therapy. American Journal of the Medical Sciences, 2000, 319, 73. | 1.1 | 35 |
| 57 | Hemochromatosis and iron therapy of Restless Legs Syndrome. Sleep Medicine, 2001, 2, 249-251. | 1.6 | 34 |
| 58 | Inheritance of two HFE mutations in African Americans: Cases with hemochromatosis phenotypes and estimates of hemochromatosis phenotype frequency. Genetics in Medicine, 2001, 3, 294-300. | 2.4 | 34 |
| 59 | Insurance, Employment, and Psychosocial Consequences of A Diagnosis of Hereditary Hemochromatosis in Subjects Without End Organ Damage. American Journal of Gastroenterology, 2003, 98, 1175-1180. | 0.4 | 34 |
| 60 | Iron overload and prolonged ingestion of iron supplements: Clinical features and mutation analysis of hemochromatosis-associated genes in four cases. American Journal of Hematology, 2006, 81, 760-767. | 4.1 | 33 |
| 61 | African-centric TP53 variant increases iron accumulation and bacterial pathogenesis but improves response to malaria toxin. Nature Communications, 2020, 11, 473. | 12.8 | 33 |
| 62 | Transferrin Receptor-2 (TFR2) Mutation Y250X in Alabama Caucasian and African American Subjects with and without Primary Iron Overload. Blood Cells, Molecules, and Diseases, 2001, 27, 279-284. | 1.4 | 32 |
| 63 | HLA-A and -B alleles and haplotypes in hemochromatosis probands with HFEC282Y homozygosity in central Alabama. BMC Medical Genetics, 2002, 3, 9. | 2.1 | 31 |
| 64 | SLC40A1 Q248H allele frequencies and Q248H-associated risk of non-HFE iron overload in persons of sub-Saharan African descent. Blood Cells, Molecules, and Diseases, 2007, 39, 206-211. | 1.4 | 31 |
| 65 | Pica associated with iron deficiency or depletion: clinical and laboratory correlates in 262 non-pregnant adult outpatients. BMC Hematology, 2010, 10, 9. | 2.6 | 31 |
| 66 | Hepcidin, iron, and bacterial infection. Vitamins and Hormones, 2019, 110, 223-242. | 1.7 | 31 |
| 67 | Coinheritance of Alleles Associated With Hemochromatosis and Hereditary Hyperferritinemia-Cataract Syndrome. Blood, 1998, 92, 4480-4481. | 1.4 | 30 |
| 68 | Severity of iron overload in hemochromatosis: effect of volunteer blood donation before diagnosis. Transfusion, 2001, 41, 123-129. | 1.6 | 29 |
| 69 | HFE C282Y Homozygosity Is Associated With Lower Total and Low-Density Lipoprotein Cholesterol. Circulation: Cardiovascular Genetics, 2009, 2, 34-37. | 5.1 | 29 |
| 70 | Iron deficiency due to excessive therapeutic phlebotomy in hemochromatosis. American Journal of Hematology, 2000, 65, 223-226. | 4.1 | 27 |
| 71 | Transfusion Iron Overload in Adults with Acute Leukemia: Manifestations and Therapy. American Journal of the Medical Sciences, 2000, 319, 73-78. | 1.1 | 27 |
| 72 | HLA-A and -B alleles and haplotypes in 240 index patients with common variable immunodeficiency and selective IgG subclass deficiency in central Alabama. BMC Medical Genetics, 2003, 4, 3. | 2.1 | 27 |

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|---|---|---|---|
| 73 | Geographic and racial/ethnic differences in HFE mutation frequencies in the Hemochromatosis and Iron Overload Screening (HEIRS) Study. Ethnicity and Disease, 2006, 16, 815-21. | 2.3 | 27 |
| 74 | Genetic screening for iron overload: No evidence of discrimination at 1 year. Journal of Family Practice, 2007, 56, 829-34. | 0.2 | 27 |
| 75 | Hemochromatosis and Severe Iron Overload Associated with Compound Heterozygosity for TFR2 R455Q and Two Novel Mutations TFR2 R396X and G792R. Acta Haematologica, 2006, 115, 102-105. | 1.4 | 25 |
| 76 | Cirrhosis in Hemochromatosis: Independent Risk Factors in 368 HFE p.C282Y Homozygotes. Annals of Hepatology, 2018, 17, 871-879. | 1.5 | 25 |
| 77 | Current status of blastic transformation in chronic myelogenous leukemia. American Journal of Hematology, 1978, 4, 281-291. | 4.1 | 24 |
| 78 | Optimal Management Strategies for Chronic Iron Overload. Drugs, 2007, 67, 685-700. | 10.9 | 24 |
| 79 | Selective subnormal IgG3 in 121 adult index patients with frequent or severe bacterial respiratory tract infections. Cellular Immunology, 2016, 299, 50-57. | 3.0 | 24 |
| 80 | Prevalence of iron deficiency in 62,685 women of seven race/ethnicity groups: The HEIRS Study. PLoS ONE, 2020, 15, e0232125. | 2.5 | 24 |
| 81 | Radioimmunometric Quantification of Surface Lactoferrin in Blood Mononuclear Cells. American Journal of the Medical Sciences, 1994, 307, 102-107. | 1.1 | 22 |
| 82 | Dietary Iron Intake and Serum Ferritin Concentration in 213 Patients Homozygous for theHFE<sup>C282Y</sup>Hemochromatosis Mutation. Canadian Journal of Gastroenterology & Hepatology, 2012, 26, 345-349. | 1.7 | 22 |
| 83 | Comparisons of CVID and IgGSD: Referring Physicians, Autoimmune Conditions, Pneumovax Reactivity, Immunoglobulin Levels, Blood Lymphocyte Subsets, and HLA-A and -B Typing in 432 Adult Index Patients. Journal of Immunology Research, 2014, 2014, 1-10. | 2.2 | 22 |
| 84 | HFEMutations in Caucasian Participants of the Hemochromatosis and Iron Overload Screening Study with Serum Ferritin Level <1000 Ï¼g/L. Canadian Journal of Gastroenterology & Hepatology, 2013, 27, 390-392. | 1.7 | 21 |
| 85 | Iron, Lead, and Cobalt Absorption: Similarities and Dissimilarities. Experimental Biology and Medicine, 1981, 166, 64-69. | 2.4 | 20 |
| 86 | HFE C282Y and H63D in adults with malignancies in a community medical oncology practice. BMC Cancer, 2004, 4, 6. | 2.6 | 20 |
| 87 | Iron overload in an African American woman with SS hemoglobinopathy and a promoter mutation in the X-linked erythroid-specific 5-aminolevulinate synthase (ALAS2) gene. Blood Cells, Molecules, and Diseases, 2005, 34, 226-228. | 1.4 | 20 |
| 88 | Case Report: Neoplastic Meningitis with Eosinophilic Pleocytosis in Hodgkinâ€™s Disease: A Case with Cerebellar Dysfunction and a Review of the Literature. American Journal of the Medical Sciences, 1988, 296, 322-326. | 1.1 | 20 |
| 89 | Pseudochloroma: Extramedullar Hematopoietic Nodules in Chronic Myelogenous Leukemia. Annals of Internal Medicine, 1979, 91, 735. | 3.9 | 20 |
| 90 | Multi-Organ Iron Overload in an African-American Man with ALAS2 R452S and SLC40A1 R561G. Acta Haematologica, 2008, 120, 168-173. | 1.4 | 19 |

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|---|---|---|---|
| 91 | Hemochromatosis and Iron Overload: From Bench to Clinic. American Journal of the Medical Sciences, 2013, 346, 403-412. | 1.1 | 19 |
| 92 | Diagnosis of hemochromatosis in family members of probands: A comparison of phenotyping and HFE genotyping. Genetics in Medicine, 1999, 1, 89-93. | 2.4 | 18 |
| 93 | Juvenile Hemochromatosis in the Southeastern United States: A Report of Seven Cases in Two Kinships. Blood Cells, Molecules, and Diseases, 2002, 29, 104-115. | 1.4 | 18 |
| 94 | Attitudes about and Psychosocial Outcomes of HFE Genotyping for Hemochromatosis. Genetic Testing and Molecular Biomarkers, 2004, 8, 90-97. | 1.7 | 18 |
| 95 | Isolated marrow lymphoma: An entity of possible T-cell derivation. Cancer, 1980, 46, 1767-1774. | 4.1 | 17 |
| 96 | 15-hydroxyprostaglandin dehydrogenase and Î”13 reductase content of gastrointestinal organs of rabbits and rats. Prostaglandins, 1981, 21, 15-23. | 1.2 | 17 |
| 97 | Population screening for hemochromatosis: Has the time finally come?. Current Gastroenterology Reports, 2000, 2, 18-26. | 2.5 | 17 |
| 98 | Hemochromatosis: population genetics. , 2000, , 42-50. | | 17 |
| 99 | Variation of hemochromatosis prevalence and genotype in national groups. , 2000, , 51-62. | | 17 |
| 100 | Allele frequencies of hemojuvelin gene (HJV) I222N and G320V missense mutations in white and African American subjects from the general Alabama population. BMC Medical Genetics, 2004, 5, 29. | 2.1 | 17 |
| 101 | Total blood lymphocyte counts in hemochromatosis probands with HFEC282Y homozygosity: relationship to severity of iron overload and HLA-A and -B alleles and haplotypes. BMC Hematology, 2005, 5, 5. | 2.6 | 17 |
| 102 | <i>SLC40A1</i> c.1402Gâ†'A Results in Aberrant Splicing, Ferroportin Truncation after Glycine 330, and an Autosomal Dominant Hemochromatosis Phenotype. Acta Haematologica, 2007, 118, 237-241. | 1.4 | 17 |
| 103 | Common variable immunodeficiency and IgG subclass deficiency in central Alabama hemochromatosis probands homozygous for HFE C282Y. Blood Cells, Molecules, and Diseases, 2003, 31, 102-111. | 1.4 | 16 |
| 104 | Accuracy of Family History of Hemochromatosis or Iron Overload: The Hemochromatosis and Iron Overload Screening Study. Clinical Gastroenterology and Hepatology, 2008, 6, 934-938. | 4.4 | 16 |
| 105 | Thyroid-Stimulating Hormone and Free Thyroxine Levels in Persons with HFE C282Y Homozygosity, a Common Hemochromatosis Genotype: The HEIRS Study. Thyroid, 2008, 18, 831-838. | 4.5 | 16 |
| 106 | Hemochromatosis Detection in a Health Screening Program at an Alabama Forest Products Mill. Journal of Occupational and Environmental Medicine, 2002, 44, 745-751. | 1.7 | 15 |
| 107 | Phlebotomy-Mobilized Iron as a Surrogate for Liver Iron Content in Hemochromatosis Patients. Hematology, 2003, 8, 429-432. | 1.5 | 15 |
| 108 | Psychosocial Impact of Genetic Testing for Hemochromatosis in The HEIRS Study: A Comparison of Participants Recruited in Canada And in The United States. Genetic Testing and Molecular Biomarkers, 2007, 11, 55-64. | 1.7 | 15 |

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|---|---|---|---|
| 109 | Effects of Highly Conserved Major Histocompatibility Complex (MHC) Extended Haplotypes on Iron and Low CD8+ T Lymphocyte Phenotypes in HFE C282Y Homozygous Hemochromatosis Patients from Three Geographically Distant Areas. PLoS ONE, 2013, 8, e79990. | 2.5 | 15 |
| 110 | Neutrophil Function in Chronic Neutrophilic Leukemia: Defective Respiratory Burst in Response to Phorbol Esters. Acta Haematologica, 1992, 87, 16-21. | 1.4 | 14 |
| 111 | Transferrin Saturation Phenotype andHFEGenotype Screening for Hemochromatosis and Primary Iron Overload: Predictions from a Model Based on National, Racial, and Ethnic Group Composition in Central Alabama. Genetic Testing and Molecular Biomarkers, 2000, 4, 199-206. | 1.7 | 14 |
| 112 | Disparate phenotypic expression of ALAS2 R452H (nt 1407 GÂâ†'ÂA) in two brothers, one with severe sideroblastic anemia and iron overload, hepatic cirrhosis, and hepatocellular carcinoma. Blood Cells, Molecules, and Diseases, 2006, 36, 342-346. | 1.4 | 14 |
| 113 | Heritability of serum iron measures in the hemochromatosis and iron overload screening (HEIRS) family study. American Journal of Hematology, 2010, 85, 101-105. | 4.1 | 14 |
| 114 | Selective Subnormal IgG1 in 54 Adult Index Patients with Frequent or Severe Bacterial Respiratory Tract Infections. Journal of Immunology Research, 2016, 2016, 1-10. | 2.2 | 14 |
| 115 | Retention of radiolead by human erythrocytes in vitro. Toxicology and Applied Pharmacology, 1989, 99, 314-322. | 2.8 | 13 |
| 116 | Abnormalities of flavin monooxygenase as an etiology for sideroblastic anemia. American Journal of Hematology, 2000, 65, 149-153. | 4.1 | 13 |
| 117 | Mechanisms of iron toxicity. , 2000, , 229-238. | | 13 |
| 118 | Cardiac abnormalities in hemochromatosis. , 2000, , 297-311. | | 13 |
| 119 | Vibrio vulnificus Infection in a Hemodialysis Patient Receiving Intravenous Iron Therapy. Clinical Infectious Diseases, 2003, 37, e63-e67. | 5.8 | 13 |
| 120 | GNPAT p.D519G is independently associated with markedly increased iron stores in HFE p.C282Y homozygotes. Blood Cells, Molecules, and Diseases, 2017, 63, 15-20. | 1.4 | 13 |
| 121 | Countries of ancestry reported by hemochromatosis probands and control subjects in central Alabama. Ethnicity and Disease, 2004, 14, 73-81. | 2.3 | 13 |
| 122 | Isolated Chylopericardium Associated With Lymphoma. Southern Medical Journal, 1980, 73, 1551-1552. | 0.7 | 12 |
| 123 | Neutrophil lactoferrin content: Variation among mammals. The Anatomical Record, 1988, 221, 567-575. | 1.8 | 12 |
| 124 | Non-diabetic endocrinopathy in hemochromatosis. , 2000, , 278-289. | | 12 |
| 125 | The mitochondrial nt 16189 polymorphism and hereditary hemochromatosis. Blood Cells, Molecules, and Diseases, 2004, 33, 344-345. | 1.4 | 12 |
| 126 | Three kinships with ALAS2 P520L (c. 1559 CÂâ†'ÂT) mutation, two in association with severe iron overload, and one with sideroblastic anemia and severe iron overload. Blood Cells, Molecules, and Diseases, 2006, 36, 292-297. | 1.4 | 12 |

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|---|---|---|---|
| 127 | Characteristics of participants with selfâ€reported hemochromatosis or iron overload at HEIRS study initial screening. American Journal of Hematology, 2008, 83, 126-132. | 4.1 | 12 |
| 128 | Common TMPRSS6 mutations and iron, erythrocyte, and pica phenotypes in 48 women with iron deficiency or depletion. Blood Cells, Molecules, and Diseases, 2012, 48, 124-127. | 1.4 | 12 |
| 129 | Serum Ferritin, Insulin Resistance, and Metabolic Syndrome: Clinical and Laboratory Associations in 769 Non-Hispanic Whites Without Diabetes Mellitus in the HEIRS Study. Metabolic Syndrome and Related Disorders, 2015, 13, 57-63. | 1.3 | 12 |
| 130 | Assessment of total immunoreactive lactoferrin in hematopoietic cells using flow cytometry. Journal of Immunological Methods, 1988, 108, 159-170. | 1.4 | 11 |
| 131 | Normal Transferrin Saturation in Hemochromatosis. Hospital Practice (1995), 1991, 26, 45-48. | 1.0 | 11 |
| 132 | Hemochromatosis, iron overload, and porphyria cutanea tarda. , 2000, , 453-467. | | 11 |
| 133 | Survey of physician knowledge about hemochromatosis. Genetics in Medicine, 2002, 4, 136-141. | 2.4 | 11 |
| 134 | Stainable hepatic iron in 341 African American adults at coroner/medical examiner autopsy. BMC Clinical Pathology, 2005, 5, 2. | 1.8 | 11 |
| 135 | Hemochromatosis,<I> HFE</I> C282Y Homozygosity, and Bariatric Surgery: Report of Three Cases. Obesity Surgery, 2004, 14, 1409-1414. | 2.1 | 10 |
| 136 | HFE Hemochromatosis and Hepatic Sarcoid. American Journal of the Medical Sciences, 2009, 337, 386-390. | 1.1 | 10 |
| 137 | Mild Iron Overload in an African American Man with SLC40A1 D270V. Acta Haematologica, 2012, 128, 28-32. | 1.4 | 10 |
| 138 | Autoimmune Conditions in 235 Hemochromatosis Probands withHFEC282Y Homozygosity and Their First-Degree Relatives. Journal of Immunology Research, 2015, 2015, 1-11. | 2.2 | 10 |
| 139 | Duration of frequent or severe respiratory tract infection in adults before diagnosis of IgG subclass deficiency. PLoS ONE, 2019, 14, e0216940. | 2.5 | 10 |
| 140 | Characterization of adult patients with IgG subclass deficiency and subnormal IgG2. PLoS ONE, 2020, 15, e0240522. | 2.5 | 10 |
| 141 | The aplastic anemia-paroxysmal nocturnal hemoglobinuria syndrome. American Journal of Hematology, 1979, 7, 61-67. | 4.1 | 9 |
| 142 | Human leukocyte antigen (HLA) association and typing in hemochromatosis. , 2000, , 63-74. | | 9 |
| 143 | The ancestral haplotype in hemochromatosis. , 2000, , 91-98. | | 9 |
| 144 | Vibrio vulnificus Bacteremia Associated with Chronic Lymphocytic Leukemia, Hypogammaglobulinemia, and Hepatic Cirrhosis: Relation to Host and Exposure Factors in 252 V. Vulnificus Infections Reported in Louisiana. American Journal of the Medical Sciences, 2006, 332, 216-220. | 1.1 | 9 |

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|---|---|---|---|
| 145 | Diabetes in First-Degree Family Members: A Predictor of Type 2 Diabetes in 159 Nonscreening Alabama Hemochromatosis Probands WithHFEC282Y Homozygosity. Diabetes Care, 2014, 37, 259-266. | 8.6 | 9 |
| 146 | Risk Factors for Insulin Resistance, Metabolic Syndrome, and Diabetes in 248HFEC282Y Homozygotes Identified by Population Screening in the HEIRS Study. Metabolic Syndrome and Related Disorders, 2016, 14, 94-101. | 1.3 | 9 |
| 147 | Case Report: Immune Thrombocytopenia: Effects of Maternal Gamma Globulin Infusion on Maternal and Fetal Serum, Platelet, and Monocyte IgG. American Journal of the Medical Sciences, 1987, 293, 112-118. | 1.1 | 8 |
| 148 | Immunoreactive lactoferrin in resting, activated, and neoplastic lymphocytes. Leukemia Research, 1990, 14, 441-447. | 0.8 | 8 |
| 149 | Iron as a carcinogen. , 2000, , 239-249. | | 8 |
| 150 | HFEGenotype Frequencies in Consecutive Reference Laboratory Specimens: Comparisons among Referral Sources and Association with Initial Diagnosis. Genetic Testing and Molecular Biomarkers, 2001, 5, 299-306. | 1.7 | 8 |
| 151 | HFE C282Y Homozygotes Aged 25â€“29 Years at HEIRS Study Initial Screening. Genetic Testing and Molecular Biomarkers, 2007, 11, 269-275. | 1.7 | 8 |
| 152 | African Americans at Risk for Increased Iron Stores or Liver Disease. American Journal of Medicine, 2007, 120, 734.e1-734.e9. | 1.5 | 8 |
| 153 | Serum immunoglobulins in 28 adults with autoimmune sensorineural hearing loss: increased prevalence of subnormal immunoglobulin G1 and immunoglobulin G3. BMC Immunology, 2014, 15, 43. | 2.2 | 8 |
| 154 | Pneumococcal vaccination responses in adults with subnormal IgG subclass concentrations. BMC Immunology, 2019, 20, 29. | 2.2 | 8 |
| 155 | Differential staining of neutrophils and monocytes: Surface and cytoplasmic iron-binding proteins. The Histochemical Journal, 1988, 20, 147-155. | 0.6 | 7 |
| 156 | T-lymphocyte expression and function in hemochromatosis. , 2000, , 396-408. | | 7 |
| 157 | Clinical spectrum of hepatic disease in hemochromatosis. , 2000, , 250-257. | | 7 |
| 158 | Juvenile hemochromatosis. , 2000, , 318-326. | | 7 |
| 159 | The iron phenotype of hemochromatosis heterozygotes. , 2000, , 411-418. | | 7 |
| 160 | Characteristics of HFE C282Y Homozygotes Younger than Age 30 Years. Acta Haematologica, 2004, 112, 219-221. | 1.4 | 7 |
| 161 | Longer survival associated with HLAâ€A*03, B*14 among 212 hemochromatosis probands with HFE C282Y homozygosity and HLAâ€A and â€B typing and haplotyping<sup>1</sup>. European Journal of Haematology, 2010, 85, 439-447. | 2.2 | 7 |
| 162 | Tumor necrosis factor-alpha promoter variants and iron phenotypes in 785 Hemochromatosis and Iron Overload Screening (HEIRS) Study participants. Blood Cells, Molecules, and Diseases, 2010, 44, 252-256. | 1.4 | 7 |

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