

Review on Haematologists on Hematological Disorders

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Review Article

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ABSTRACT

Study of blood and its components is called haematology. The branch of science that involves study and treatment of blood. The malfunction in production of blood and its components is called as Hematological disorders. The diseases include hemophilia, blood clots, other bleeding disorders and blood cancers such as lymphoma, leukemia and myeloma. The laboratory work that goes into the study of blood is frequently performed by medical technicians or medical lab scientist. Hematologists work as hematologist or oncologists providing medical treatment for all cancers. Pathologists who are specialised in blood diagnosis are called hematologists. Hematologists and hematopathologists do combined work to formulate a diagnosis resulting the most appropriate therapy to the patient. Hematology is very near to internal medicine and differs from medical oncology.

INTRODUCTION

Hematology is a branch of science that deals with the study of blood and its blood-forming organs, and blood diseases. Hematology is practised by clinicians who deal with the blood diagnosis its treatment and overall blood disorders ranging from anemia to blood cancer. The diseases which are mostly treated by haematologists include: Iron deficiency anaemia and types of anemia e.g. sickle cell anemia or trauma-related anemia, polycythemia, myelofibrosis, leukemia, platelet and bleeding disorders such as hemophilia, idiopathic thrombocytopenic purpura and Von Willebrand disease, The myelodysplastic syndromes, hemoglobinopathies such as thalassemia and sickle cell disease, multiple myeloma, malignant lymphomas, blood transfusion, bone marrow stem cell transplantation [1-10].

The main medical blood treatment methods are:

1. Using blood products
2. Intravenous administration
3. Bloodletting

TRANSFUSION OF BLOOD AND BLOOD PRODUCTS

Blood transfusion is the commonly used treatment of hemorrhage and to improve transport of oxygen to tissues. Transfusion of red blood cells is based on the patient's clinical and health condition. Indications for transfusion include symptomatic anemia, lack of sickle cells and acute loss of blood i.e more than 30% of blood volume. Anticoagulation effect can be reversed with the help of frozen plasma infusion. Platelet function defects can be stopped by Platelet transfusion which helps in the prevention of hemorrhage in thrombocytopenia patients. Cryoprecipitate is used in hypofibrinogenemia cases and is most often seen in setting the massive hemorrhage or consumptive coagulopathy [20-40]. Transfusion-related infections are less commonly seen with noninfectious complications. Noninfectious complications of transfusions are divided into noninfectious and serious hazards of transfusion. Acute complications are seen in minutes to 24 hours of transfusion and delayed complications may prolong to days, months or even years together. Blood transfusion is a lifesaving procedure and has its own merits and demerits like infectious and noninfectious complications. Clinical trials investigation confirmed that transfer of blood at lower hemoglobin levels is beneficial. Packed red blood cells (RBCs) are prepared from whole blood by removing approximately 250 ml of plasma. One packed unit RBCs should increase levels of hemoglobin by 1 g/dl (10 g/L) and hematocrit by 3%. Packed RBC units are filtered to reduce leukocytes which limits febrile nonhemolytic transfusion reactions (FNHTRs) and are considered safe [41-60].

INTRAVENOUS THERAPY

Intravenous mean infusion of liquid substances directly into the vein. Therapies administered intravenously are called as intravenous therapy and are often done with the use of drugs. Intravenous infusions are commonly referred as drips because many administration systems employ a drip chamber that prevents air from disturbing the blood stream allowing the smooth blood flow. This therapy is used to correct the electrolyte imbalances, blood transfusion or fluid replacement e.g. dehydration. It can also be used in chemotherapy. If compared with other routes, the intravenous route is the fastest way to inject fluids and medications throughout the body [61-70]. The bioavailability is 100% by using IV treatment. Some commonly found adverse effects of Intravenous therapy are pain, infection, phlebitis, infiltration/extravasation, fluid overload, hypothermia, electrolyte imbalance, embolism, glucose [71-85].

BLOODLETTING

It is an extraction of blood from a patient to cure illness. Bloodletting was based on an ancient system of science in which blood and other fluids that had to remain in proper balance to maintain health. The most commonly used medical practice done by surgeons until the late 19th century. The practice has been stopped by modern medicine because of some specific conditions. Bloodletting has a beneficial effect in temporarily decreasing the blood pressure in the absence of other hypertension treatments. Hypertension is an asymptomatic and un-diagnosable condition and is unintentional. The use of bloodletting was harmful to patients in most of the cases. Phlebotomy means extraction of blood for laboratory analysis or blood transfusion. The drawing of blood to analyse specific cases like hemochromatosis, polycythemia vera etc. comes under Therapeutic phlebotomy. Presently the medical practice of bloodletting is considered to be a pseudoscience [86-100].

CONCLUSION

Scientists have attained tremendous progress in treating blood disorders by using different sources like myelofibrosis, Multiple myeloma, myelodysplastic syndromes, and blood transfusion etc. They have invented many blood treatment methods and are successfully applied in saving human life. All of these are successful blood treatment methods. If one or more parts of the blood is affected and prevent to perform its function leads to Blood disorders. Blood disorders can be acute, chronic or inherited. Other causes include medicinal side effects and lack of certain diet nutrients.

REFERENCES

1. Reuter DA, et al. Stroke volume variation for assessment of cardiac responsiveness to volume loading in mechanically ventilated patients after cardiac surgery. *Intensive Care Med.* 2002;28:392-398.
2. Rajput RS, et al. Comparison of Cardiac output measurement by noninvasive method with electrical cardiometry and invasive method with thermodilution technique in patients undergoing coronary artery bypass grafting. *World Journal of Cardiovascular Surgery.* 2014;4:123-130.
3. Zoremba N, et al. LTE Comparison of electrical velocimetry and thermodilution techniques for the measurement of cardiac output. *Acta Anaesthesiol Scand.* 2007;51:1314-1319.
4. Schmidt C, et al. A Comparison of electrical velocimetry and transoesophageal Doppler echocardiography for measuring stroke volume and cardiac output. *British Journal of Anaesthesia.* 2005;95:603-610.
5. Narula J, et al. Electrical Cardiometry in Patients undergoing Cardiac Catheterisation. *International Journal of Perioperative Ultrasound and Applied Technologies.* 2013;2:102-107.
6. Zhang L, et al. Early goal-directed therapy in the management of severe sepsis or septic shock in adults: a meta-analysis of randomized controlled trials. *BMC Medicine.* 2015;13:71.
7. Bernstein DP and Lemmens HJ Stroke volume equation for impedance cardiography. *Med Biol Eng Comput.* 2005;43:443-450.
8. Bernstein DP Bernstein-Osypka stroke volume equation for impedance cardiography: citation correction. *Intensive Care Med.* 33: 923.
9. Eng J. Sample Size Estimation: How Many Individuals Should Be Studied? *Radiology.* 2003;227:309-313.

10. Guinot PG, et al. Mini-fluid challenge can predict arterial pressure response to volume expansion in spontaneously breathing patients under spinal anaesthesia. *Anaesth Crit Care Pain Med.* 2015;32:645-649.
11. Cheng Li, et al. Stroke Volume Variation for Prediction of Fluid Responsiveness in Patients Undergoing Gastrointestinal Surgery. *Int J Med Sci.* 2013;10:148-155.
12. Soliman R, et al. Stroke volume variation compared with pulse pressure variation and cardiac index changes for prediction of fluid responsiveness in mechanically ventilated patients. *EJCCM.* 2015;3:9-16.
13. Marx G, et al. Assessing fluid responsiveness by stroke volume variation in mechanically ventilated patients with severe sepsis. *Eur J Anaesthesiol.* 2004;02:132-138.
14. Peake SL, et al. Goal-directed resuscitation for patients with early septic shock. *N Engl J Med.* 2014;371:1496-1506.
15. Yealy DM, et al. A randomized trial of protocol-based care for early septic shock. *N Engl J Med.* 2014;370:1683-1693.
16. IBM Corp. IBM SPSS Statistics for Windows, Version 22.0. Armonk, NY: IBM Corp 2013.
17. Youden WJ. An index for rating diagnostic test. *Cancer.* 1950;3:32-35.
18. Lopes MR, et al. Goal-directed fluid management based on pulse pressure variation monitoring during high-risk surgery: a pilot randomized controlled trial. *Crit Care.* 2007;11:R100.
19. Angappan S, Parida S, Vasudevan A, Badhe AS (2015) The comparison of stroke volume variation with central venous pressure in predicting fluid responsiveness in septic patients with acute circulatory failure. *Indian J Crit Care Med.* 19: 394-400.
20. Berkenstadt H, et al. Stroke volume variation as a predictor of fluid responsiveness in patients undergoing brain surgery. *Anesth Analg.* 2001;92:984-989.
21. Kim KM, et al. Pulse pressure variation and stroke volume variation to predict fluid responsiveness in patients undergoing carotid endarterectomy. *Korean J Anesthesiol.* 2013;65:237-243.
22. Suehiro K, et al. Stroke volume variation as a predictor of fluid responsiveness in patients undergoing airway pressure release ventilation. *Anaesth Intensive Care.* 2012;40:767-772.
23. Peng K, et al. Goal-Directed Fluid Therapy Based on Stroke Volume Variations Improves Fluid Management and Gastrointestinal Perfusion in Patients Undergoing Major Orthopedic Surgery. *Med Princ Pract.* 2014;23:413-420.
24. Zhang J, et al. Goal-directed fluid optimization based on stroke volume variation and cardiac index during one-lung ventilation in patients undergoing thoracoscopy lobectomy operations: a pilot study. *Clinics (Sao Paulo).* 2013;68:1065-1070.
25. Gallagher JE, et al. Public health aspects of tobacco control revisited. *Int Dent J.* 2010;60:31-49.
26. Nisar N, et al. Pattern of tobacco consumption among adult women of low socioeconomic community Karachi, Pakistan. *J Pak Med Assoc.* 2005;55:111-114.
27. Schroeder SA. Tobacco control in the wake of the 1998 master settlement agreement. *New England Journal of Medicine.* 2004;350:293-301.
28. WHO. World Health Organization Framework Convention on Tobacco Control. Geneva 2003.
29. Petersen PE. World Health Organization global policy for improvement of oral health: World Health Assembly 2007. *Int Dent J.* 2008;58:115-121.
30. Davis JM, et al. Education of tobacco use prevention and cessation for dental professionals-a paradigm shift. *Int Dent J.* 2010;60:60-72.
31. Wiener RC and Pla RMW. Evaluation of educational material for tobacco prevention and cessation used in West Virginia University Dental Programs. *J Dent Hyg.* 2011;85:204-210.
32. Ehizele A, et al. Oral health knowledge, attitude and practices among Nigerian primary school teachers. *Int J Dent Hyg.* 2011;9:254-260.
33. Sood P, et al. Dental patient's knowledge and perceptions about the effects of smoking and role of dentists in smoking cessation activities. *Eur J Dent.* 2014;8:216-223.
34. Terrades M, et al. Patients' knowledge and views about the effects of smoking on their mouths and the involvement of their dentists in smoking cessation activities. *Br Dent J.* 2009;207:542-543.
35. Rikard-Bell G, et al. Preventive dentistry: what do Australian patients endorse and recall of smoking cessation advice by their dentists? *Br Dent J.* 2003;194:159-164.

36. Lung ZH, et al. Poor patient awareness of the relationship between smoking and periodontal diseases. *Br Dent J.* 2005;199:731-737.
37. Al-Shammari KF, et al. Dental patient awareness of smoking effects on oral health: comparison of smokers and non-smokers. *J Dent.* 2006;34:173-178.
38. Campus G, Cagetti MG, Senna A, Blasi G, Mascolo A, et al. Does smoking increase risk for caries? A cross-sectional study in an Italian military academy. *Caries Res.* 2011;45:40-46.
39. Jette AM, Feldman HA, Tennstedt SL. Tobacco use: a modifiable risk factor for dental disease among the elderly. *Am J Public Health.* 1993;83:1271-1276.
40. Petersen PE. The World Oral Health Report 2003: continuous improvement of oral health in the 21st century—the approach of the WHO Global Oral Health Programme. *Community Dent Oral Epidemiol.* 2003;31:3-24.
41. Gallagher JE and Eaton KA. Health workforce governance and oral health: Diversity and challenges in Europe. *Health Policy.* 2015;119:1565-1575.
42. Butler CC, Pill R, Stott NCH. Qualitative study of patients' perceptions of doctors' advice to quit smoking: implications for opportunistic health promotion. *BMJ.* 1998;316:1878-1881.
43. Chaudry AI, et al. Carotid cavernous fistula: ophthalmological implications. *Middle East Afr J Ophthalmology.* 2009;16:57-63.
44. Oishi A, et al. (2009) Etiology of carotid cavernous fistula in Japanese. *Jpn J Ophthalmol.* 53: 40-43.
45. Uehara T, et al. Spontaneous dural carotid cavernous sinus fistula presenting isolated ophthalmoplegia: evaluation with MR angiography. *Neurology.* 1998;50:814-816.
46. Coskun O, et al. Carotid-cavernous fistula: diagnosis with spiral CT angiography. *AJNR Am J Neuroradiol.* 2000;21:712-716.
47. Aralasmak A, Karaali K, Cevikol C, Senol U, Sindel T, et al. (2014) Venous Drainage Patterns in Carotid Cavernous Fistulas. *ISRN Radiology.* 2014: 7.
48. Hirai T, et al. Three-dimensional FISP imaging in the evaluation of carotid cavernous fistula: comparison with contrast-enhanced CT and spin-echo MR. *AJNR Am J Neuroradiol.* 1998;19:253-259.
49. Harsha KJ, et al. Susceptibility-weighted imaging in carotido-cavernous fistulas. A case control study. *Interv Neuroradiol.* 2013;19:438-444.
50. Seeger A, et al. Feasibility of Noninvasive Diagnosis and Treatment Planning in a Case Series with Carotid-Cavernous Fistula using High-Resolution Time-Resolved MR-Angiography with Stochastic Trajectories (TWIST) and Extended Parallel Acquisition Technique (ePAT 6) at 3 T. *Clin Neuroradiol.* 2015;25:241-247.
51. Ching NG et al. A Man with Breast Cancer Following Hormonal Treatment for Prostate Cancer *J Med Diagn Meth.* 2013;2:112.
52. Iacono F et al. Treating Idiopathic Male Infertility with a Combination of Tamoxifen Citrate and a Natural Compost with Antioxidant and Androgen-Mimetic Action. *J Steroids Hormon Sci.* 2013;S5:002.
53. Shaaban MM, et al. Follicular Fluid Activin A and Leptin are not Correlated with IVF Outcome Measures. *J Steroids Horm Sci.* 2012;4:111.
54. Santos FGD, et al. Regulation of Glucose Transporter 1 (Slc2a1) in the Pituitary Gonadotrope of Mice after Puberty. *J Steroids Hormon Sci.* 2014;5:138.
55. Declercq J, et al. Replenishing a Balanced Mixture of Hormone Producing Cells: A Necessary Component in the Stem Cell Based Therapy for Diabetes? *J Steroids Horm Sci.* 2012;3:e111.
56. Wojcik M, et al. High Incidence of Abnormal Circadian Blood Pressure Profiles in Patients on Steroid Replacement Therapy due to Secondary Adrenal Insufficiency and Congenital Adrenal Hyperplasia without Overt Hypertension - Initial Results. *J Steroids Hormon Sci.* 2013;S12:005.
57. Neves EM, et al. Polycystic Ovary Syndrome: Correlation between Phenotypes and Metabolic Syndrome. *J Steroids Hormon Sci.* 2014;5:132.
58. McGrath KCY, et al. Androgens Rapidly Activate Nuclear Factor-Kappa B via Intracellular Ca²⁺ Signalling in Human Vascular Endothelial Cells. *J Steroids Hormon Sci.* 2012;S2:005.
59. Bandaru P, et al. The Impact of Obesity on Immune Response to Infection and Vaccine: An Insight into Plausible Mechanisms. *Endocrinol Metab Syndr.* 2013;2:113.
60. Genazzani AD, et al. Effects of a Combination of Alpha Lipoic Acid and Myo-Inositol on Insulin Dynamics in Overweight/Obese Patients with PCOS. *Endocrinol Metab Syndr.* 2014;3:140.

61. Mohamed WS, et al. Role of Ghrelin, Leptin and Insulin Resistance in Development of Metabolic Syndrome in Obese Patients. *Endocrinol Metab Synd.* 2014;3: 122.
62. Pramanik KC and Pandey AK. Natural Compounds: Prospective of Chemoprevention. *Endocrinol Metab Synd.* 2013;2:e115.
63. Nakaoka K, et al. A Case of Primary Adrenal Tuberculosis - A Diagnostic Quandary. *Endocrinol Metabol Syndrome.* 2012;1:103.
64. Chakraborty PP and Chowdhury S. A Look Inside the Pancreas: The "Endocrine-Exocrine Cross-talk". *Endocrinol Metab Synd.* 2015;4:160.
65. Shpakov AO. GPCR-Peptides: Prospective Use in Biology and Medicine. *Endocrinol Metab Synd.* 2013;2:e116.
66. Harinarayan CV, et al. Efficacy and Safety of Cholecalciferol Supplementation in Vitamin D Deficient Subjects Based on Endocrine Society Clinical Practice Guidelines. *Endocrinol Metab Synd.* 2012;S4:004.
67. Stoll H, et al. Mechanical Control of Mesenchymal Stem Cell Adipogenesis. *Endocrinol Metab Synd.* 2015;4:152.
68. Horita S, et al. Metabolic syndrome and insulin signaling in kidney. *Endocrinol Metab Synd.* 2011;S1:005.
69. Pirasath S. Glycemic Index of Traditional Foods in Northern Sri Lanka. *Endocrinol Metab Synd.* 2015;4:154.
70. Bohra A and Bhateja S. Carcinogenesis and Sex Hormones: A Review. *Endocrinol Metab Synd.* 2015;4:156.
71. Peppia M, et al. Body Composition as an Important Determinant of Metabolic Syndrome in Postmenopausal Women. *Endocrinol Metabol Syndrome.* 2012;S1:009.
72. de Piano A, et al. Nonalcoholic Fatty Liver Disease (NAFLD), a Manifestation of the Metabolic Syndrome: New Perspectives on the Nutritional Therapy. *Endocrinol Metab Synd.* 2014;3:135.
73. Guénard F, et al. Common Sequence Variants in CD163 Gene are Associated with Plasma Triglyceride and Total Cholesterol Levels in Severely Obese Individuals. *Endocrinol Metab Synd.* 2014;3:146.
74. Granados H and Phulwani P. Absent Visualization of a Hypoplastic Uterus in a 16 Year Old with Complete 46 XY Gonadal Dysgenesis (Swyer Syndrome). *Endocrinol Metab Synd.* 2013;2:114.
75. Aina O. Adrenal Psychosis, A Diagnostic Challenge. *Endocrinol Metab Synd.* 2013;2:115.
76. Mohammed SA. Differentiation between the Anterior Pituitary Cells of the Egyptian Insectivorous Bats *Rhinopoma hardwickei* using Transmission Electron Microscope. *Endocrinol Metab Synd.* 2015;4:151.
77. Burini RC, et al. Dietary Intake Association with IFG and Responses of a Lifestyle Changing Protocol in a Community-B based Adult Cohort. *Endocrinol Metab Synd.* 2014;3:125.
78. Derar DR, et al. Postpartum Ovarian Resumption in Native Dairy Cows in Upper Egypt and their Relation to Oxidant Antioxidant Status. *Endocrinol Metab Synd.* 2011;S4:002.
79. Ekmekci A, et al. Presence of Metabolic Syndrome is not an Independent Predictor of In-hospital Adverse Events in Patients with ST Elevation Myocardial Infarction that Underwent Primary Percutaneous Coronary Intervention. *Endocrinol Metab Synd.* 2013;2:112.
80. Hossein-nezhad A, et al. Circulating Omentin-1 in Obesity and Metabolic Syndrome Status Compared to Control Subjects. *Endocrinol Metab Synd.* 2012;S1:008.
81. Levine J, et al. Proton-Pump Inhibitor Treatment in Eosinophilic Esophagitis is Associated with Decreased Eosinophil Degranulation. *J Gastrointest Dig Syst.* 2015;5:259.
82. Abegunde A, et al. Change in Bowel Habit and Heme Positive Stool. *J Gastrointest Dig Syst* 2015;5:i104.
83. Hopp RN, et al. Digit Ratio is Associated with Colorectal Cancer. *J Gastrointest Dig Syst.* 5:253.
84. Taboada S and Whitney-Miller CL (2013) Updates in HER2 Testing in Gastric Cancer. *J Gastrointest Dig Syst.* 2015;3:131.
85. Tasar PT, et al. Hemosiderosis due to Chronic Alcoholism. *J Gastrointest Dig Syst.* 2014;4:182.
86. Bergholt MS, et al. Raman Endoscopy for Objective Diagnosis of Early Cancer in the Gastrointestinal System. *J Gastrointest Dig Syst.* 2013;S1:008.
87. Protic MB, et al. An Unusual Cause of Gastrointestinal Bleeding: Gastric Fundic Gland Polyps. *J Gastrointest Dig Syst.* 2014;4:203
88. Patil R, et al. Characteristics and Risk Stratification of Colon Polyps among Asymptomatic Hispanic Patients Undergoing First Time Screening Colonoscopy: A Retrospective Study. *J Gastrointest Dig Syst.* 2013;3:153.
89. Trabulo D, et al. Sweet Syndrome and Pulmonary Tuberculosis in a Crohn's Disease Patient Treated with Anti-TNF α . *J Gastrointest Dig Syst.* 2015;5:262.

90. Bertino G, et al. Management of Hepatocellular Carcinoma: An Update at the Start of 2014. *J Gastroint Dig Syst.* 2014;4:178.
91. Acar S. Plantar Erythema Nodosum Associated with Crohn's Disease. *J Gastrointest Dig Syst.* 2015;5: i102.
92. Yildirim AE, et al. An Unexpected Cause of Hyperactive Delirium in Patients with Decompensated Nonalcoholic Cirrhosis. *J Gastrointest Dig Syst.* 2015;5:261.
93. Rino Y and Yukawa N. Vitamin A, D, and E after Gastrectomy for Gastric Cancer. *J Gastroint Dig Syst.* 2013;S12:009.
94. Ebert EC. Gastrointestinal Manifestations of Churg-Strauss Syndrome. *J Gastrointest Dig Syst.* 2011;1: 101.
95. Shi D, et al. Current Status of Metal Stents for Malignant Gastro-Duodenal Obstruction. *J Gastroint Dig Syst.* 2013;3:140.
96. Salem A and Roland BC. Small Intestinal Bacterial Overgrowth (SIBO). *J Gastroint Dig Syst.* 2014;4:225.
97. Oldfield EC, et al. Nonalcoholic Fatty Liver Disease and the Gut Microbiota: Exploring the Connection. *J Gastrointest Dig Syst.* 2014;4:245.
98. Nerome K, et al. The Usefulness of an Influenza Virus-Like Particle (VLP) Vaccine Produced in Silkworm Pupae and Virosomes and Liposomes Prepared by Chemical Means: From Virosome to VLP and the Future of Vaccines. *J Gastrointest Dig Syst.* 2015;5:256.
99. Liang J, et al. Standards for Local Recurrence Rates in Both Open and Laparoscopic Rectal Cancer Surgery. How do you Measure Up?. *J Gastrointest Dig Syst.* 2015;5:260.
100. Chen GD, et al. Gender Differences in Coping Strategies for Troublesome Lower Urinary Tract Symptoms Prior to Seeking Treatment. *J Gen Practice.* 2014;2:181.