

An overview of anaemia

Keywords: anaemia, haemoglobin, sickle, microcytic, normocytic, macrocytic

Abbreviations: g/dl, gram per deciliter; fL, femtolitre; G6PD, glucose-6-phosphate dehydrogenase deficiency

Introduction

Anaemia is a commonly used term in clinical settings. Etymologically, it is from Greek (An-no or lacking; haemia-blood). It is important that anaemia does not lose its meaning, so that patients will continue to receive proper care.

Definition

Anaemia can be defined as a decrease in the number of red blood cells or their haemoglobin content, for an individual's age, sex or altitude. Anaemia is not a disease or diagnosis but a clinico-pathologic condition. It requires thorough investigation. It is estimated that about 30% of the world population is anaemic and half of these, some 600million people, have iron deficiency.¹ Although normal haemoglobin values can vary between laboratories, typical values would be less than 13.5g/dl in adult males and less than 11.0g/dl in adult females. From the age of 2 years to puberty, less than 11.0g/dl indicates anaemia. As newborn infants have a high haemoglobin level, 14.0g/dl is taken as the lower limit at birth.²

Classification

The morphological classification is given here.²

Microcytic anaemia

The red blood cells appear small microscopically. Mean corpuscular volume is <80fL. Causes of this type include iron deficiency anaemia, thalassaemia, anaemia of chronic disease (some cases), lead poisoning and sideroblastic anaemia.

Normocytic anaemia

The red blood cells are normal in size microscopically. Mean corpuscular volume is 80-95fL. Causes of this type include many haemolytic anaemias (e.g. sickle cell anaemia), anaemia of chronic disease (some cases), acute blood loss, renal disease, mixed deficiencies and bone marrow failure.

Macrocytic anaemia

The red blood cells appear larger than normal microscopically. Mean corpuscular volume is >95fL. Causes of this type may be megaloblastic or non-megaloblastic. Megaloblastic causes include vitamin B₁₂ deficiency or folate deficiency. Non-megaloblastic causes include alcohol, liver disease, myelodysplasia, aplastic anaemia, etc.

Clinical features

Symptoms³

- i. Fatigue
- ii. Breathlessness

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- iii. Malaise
- iv. Angina
- v. Dizziness
- vi. Paresthesiae
- vii. Visual disturbances
- viii. Symptoms of the underlying cause:
- ix. Bone Pain and jaundice in sickle cell disease
- x. Passage of dark urine in G6PD deficiency
- xi. Symptoms associated with severe malnutrition
- xii. Symptoms of complications e.g. severe dyspnoea and edema in heart failure

Signs.²

General signs: include pallor of mucous membranes, tachycardia, bounding pulse, systolic flow murmur and signs of heart failure. The tongue is said to be the most accurate site for the detection of pallor caused by anemia.⁴

Specific: associated with a particular cause such as koilonychia (spoon nails) in iron deficiency, jaundice in haemolytic anaemia and leg ulcers in sickle cell anaemia.

Assessment

Basically, an assessment requires:

- i. A comprehensive history.
- ii. A thorough physical examination.
- iii. Full blood count with differentials.
- iv. Peripheral blood film.

Further investigations are directed at the likely cause (e.g. iron studies, thick and thin films, serum bilirubin, haemoglobin electrophoresis and bone marrow examination) or complication.

Management

Management may involve dietary improvement, nutritional

supplementation, administration of haematinics, administration of antimicrobials, avoidance of implicated drugs or foods, blood transfusion, splenectomy and bone marrow transplant. Exact action is directed at the cause of anaemia along with other instituted treatment. Blood transfusion may not be necessary in a well-compensated patient with chronic anaemia.

Nursing

- i. Health history and examination.
- ii. Ensuring optimal in-patient overall care
- iii. Monitoring of vital signs especially during blood transfusion.
- iv. Ensuring adequate nutrition.
- v. Ensuring compliance with treatment.
- vi. Identification of patient risk factors with a view to preventing recurrence.
- vii. Educating the patient about his condition.

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Conflict of interest

The author declares no conflict of interest.

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