Medical Management of Adult Congenital Heart Disease

1. Erythrocytosis
2. Pulmonary Hemorrhage
3. The Thrombosis Dilemma
4. Non-Cardiac Surgery:
   a) The anesthesiologist
   b) Bilirubin kinetics
   c) Post operative acute gouty arthritis
5. Gynecologic Endocrinology
6. Abnormalities of great arterial walls
A Multi-System Systemic Disorder
Cyanotic Congenital Heart Disease

Erythrocytosis is a physiologically appropriate response to the decrease in tissue oxygenation caused by arterial hypoxemia. Decreased tissue oxygenation stimulates renal release of erythropoietin.

Phlebotomy reduces red blood cell mass, reduces delivery of oxygen to metabolizing tissues, and stimulates further release of erythropoietin.

Deformable biconcave discs become nondeformable iron deficient microspherocytes, thus increasing whole blood viscosity.
The Virtues of Erythrocytosis

The viscous erythrocytotic perfusate increases endothelial shear stress

The increase in shear stress causes elaboration of endothelial vasodilator substances

Vasodilatation augments flow to metabolizing tissues

NO in red blood cells enhances transfer of $O_2$ from hemoglobin to tissues

The transfer is enhanced by the increased number of red blood cells
Erythrocytosis in an iron replete state is not a risk factor for stroke due to cerebral arterial thrombotic occlusion. Iron deficient erythrocytosis is a risk factor for *venous* thrombotic stroke in infants but not in older children and adults.
Recommendations for Phlebotomy in an Iron Replete State

1. Not based on hematocrit (automated electronic particle counter) irrespective of level.
2. Recommended for temporary relief of intrusive hyperviscosity symptoms.
3. Minimum phlebotomy that achieves symptomatic relief, generally one unit with isovolumetric saline replacement.
4. Hydroxyurea blunts the erythropoietin-induced rebound.
Preoperative Phlebotomy

Whole blood is removed isovolumetrically in daily amounts of 500ml to reduce the hematocrit to just below 65%. Within hours after phlebotomy, platelet counts increase, and platelet aggregation and hemostasis improve.
“The temptation to use the anticoagulant drugs may be great. On the basis of the present studies, their use would appear to be fraught with danger.”

Robert C. Hartmann
Bull. Johns Hopkins Hospital 1952

Abnormal Hemostasis in CCHD

Intrinsic hemostatic defect(s)

Increased tissue vascularity
Pulmonary Hemorrhage in Eisenmenger Syndrome

External Hemorrhage: Hemoptysis
Internal Hemorrhage: Intrapulmonary

A common cause of sudden death
Hemoptysis in Eisenmenger Syndrome

1. *Do not bronchoscope.*
2. Determine if there is a history of antiplatelet or anti-inflammatory agents
4. CT scan if infiltrates are present.
5. Hospitalize for all but mild or moderate intrapulmonary hemorrhage.
Treatment of Pulmonary Hemorrhage in Eisenmenger Syndrome

1. Thrombocytopenia – platelet transfusion vs phlebotomy

2. Platelet Counts in Normal Range
   a) Fresh frozen plasma
   b) Cryoprecipitate
   c) Human factor VIII – safe but efficacy unproven
   d) Desmopressin (DDAVP), synthetic analog of vasopressin – neither safety nor efficacy has been proven

3. Excessively low HCT -- transfuse
Thrombosis in Dilated Hypertensive Proximal Pulmonary Arteries. A Therapeutic Dilemma

1. Anticoagulants – Efficacy is nil. The risk of reinforcing intrinsic hemostatic defect(s) and provoking hemorrhage is high.
2. Thrombolytic Agents – The efficacy of even intrapulmonary administration is nil.
The pivotal role of the cardiac anesthesiologist
Bilirubin Kinetics

- Bilirubin is formed from the breakdown of heme, a process that is excessive in the presence of the erythrocytosis of cyanotic congenital heart disease and that coincides with a substantial increase in the amount of unconjugated bilirubin.
Acute Gouty Arthritis

Prophylaxis After Resolution

Low dose oral colchicine is recommended because non-steroidal anti-inflammatory agents, even low doses, reinforce the hemostatic defects in CCHD.
CCHD
Gynecologic Endocrinology
Dysfunctional bleeding may reflect an anovulatory state. Chronic unopposed estrogen production due to anovulation leads to continuous uterine stimulation and increases the risk of endometrial hyperplasia and adenocarcinoma.
Abnormalities of Great Arterial Walls in Congenital Heart Disease
“The presence of a bicuspid aortic valve appears to indicate, at least in a portion of the cases in which it occurs, a tendency for spontaneous rupture.”

Maude Abbott 1928
Bicuspid Aortic Valve
Coarctation of the Aorta
Fallot’s Tetralogy

AAo
Fallot’s Tetralogy
Aortic Root & AR