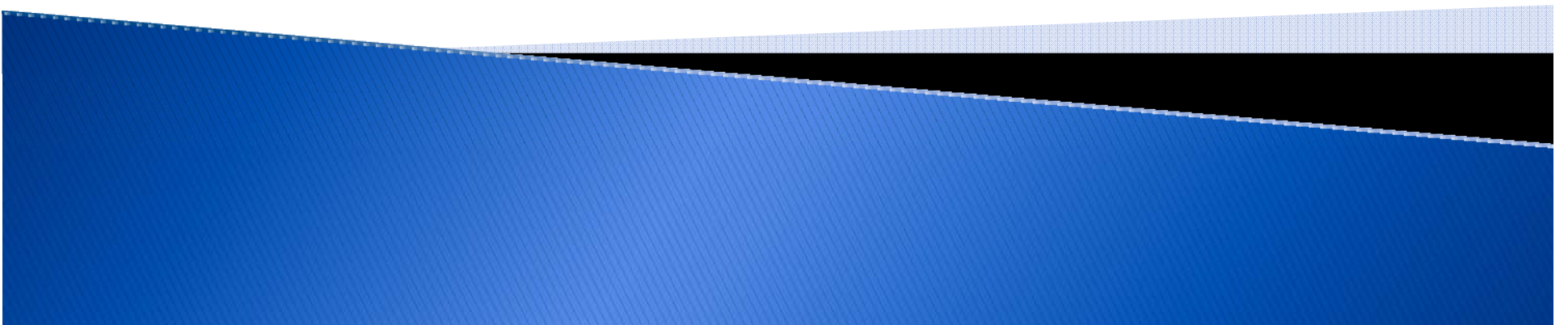
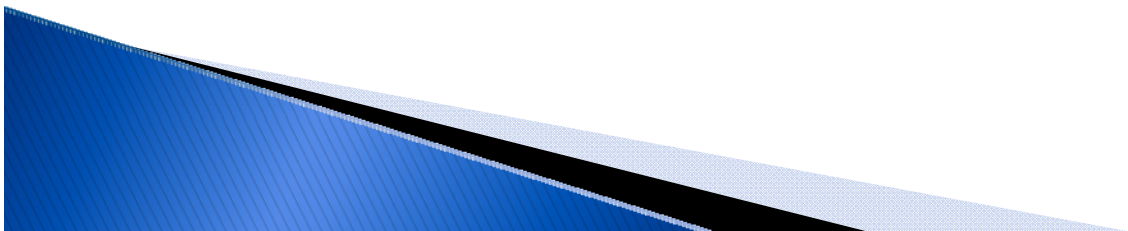


Bleeders

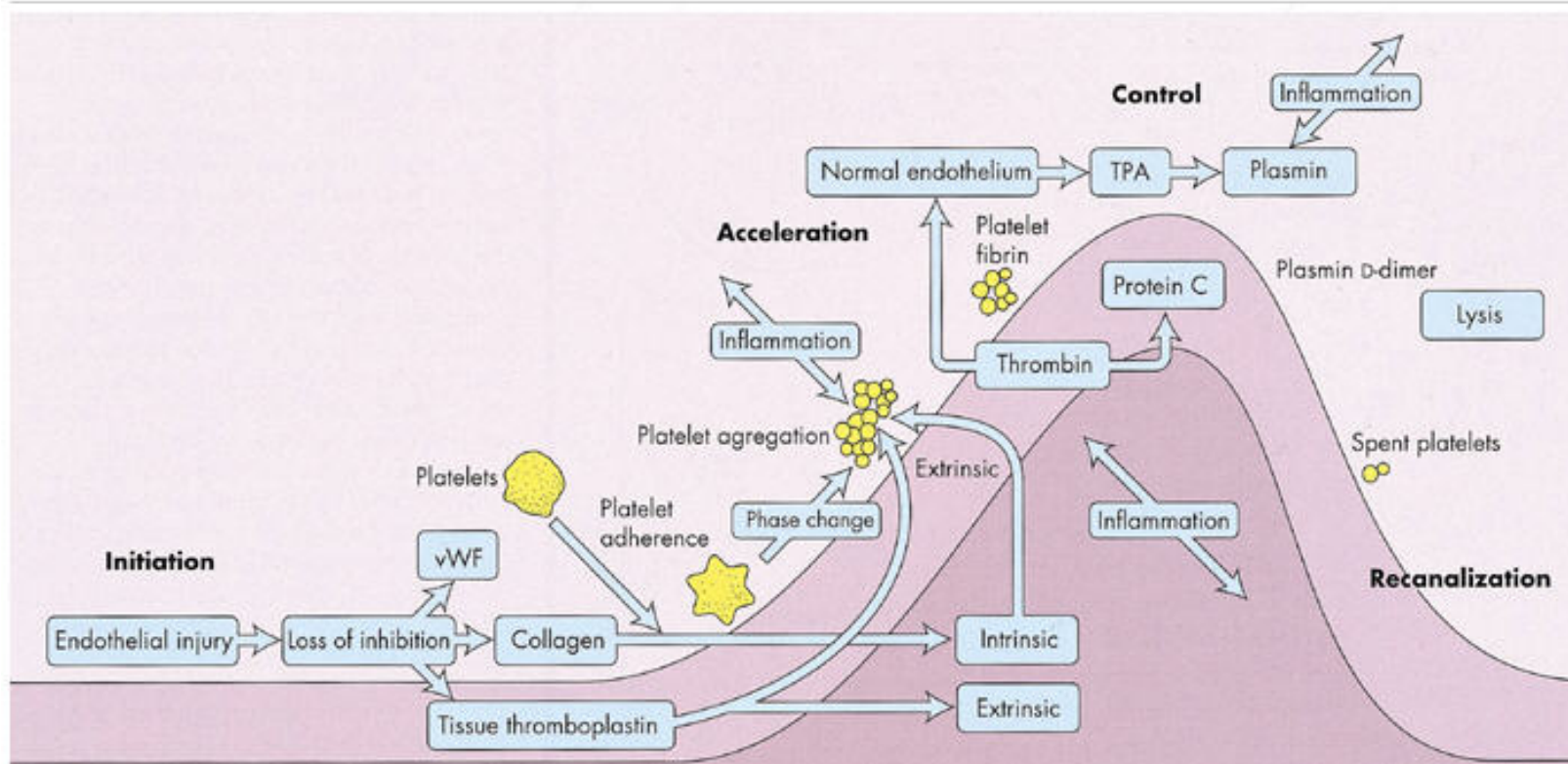


Objectives

- ▶ Identify the basic components of the coagulation process
- ▶ Describe findings associated with unusual bleeding
- ▶ List causes of abnormal bleeding
- ▶ Describe important aspects of pre-hospital care in the management of patients with abnormal bleeding.

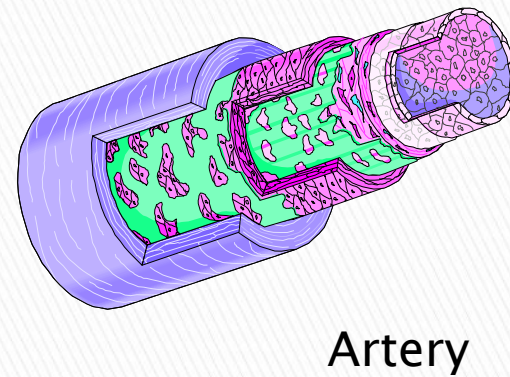
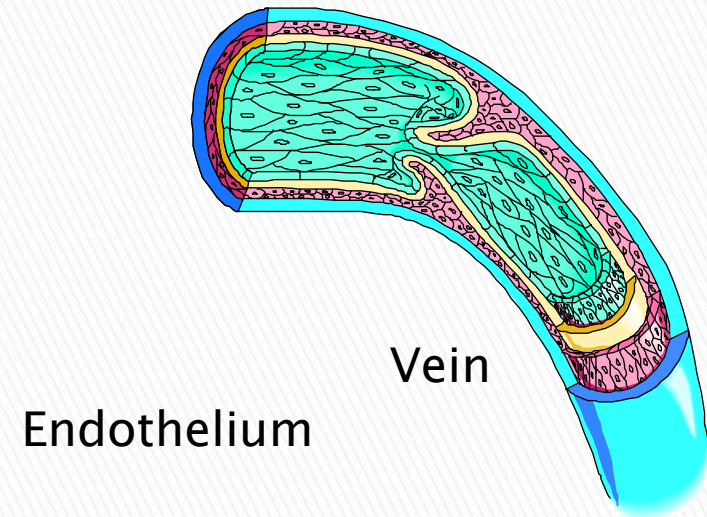


Clots: making them and breaking them

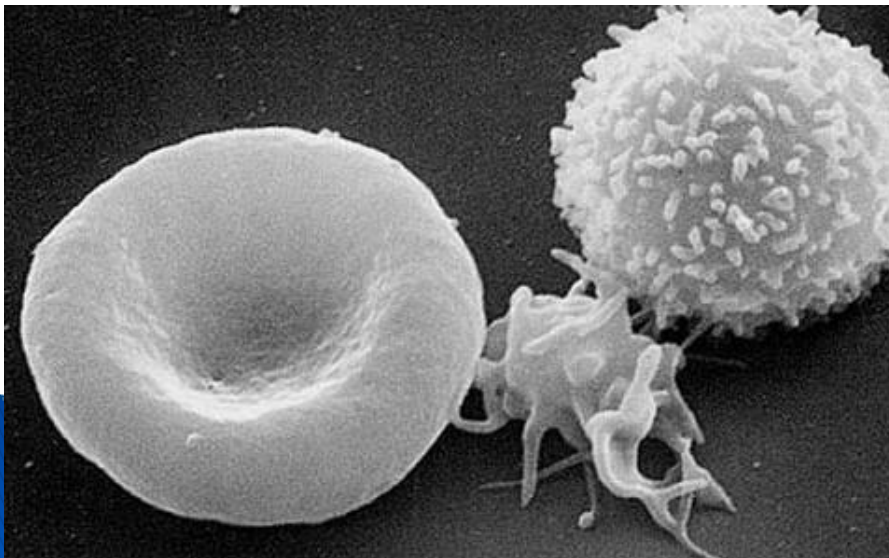
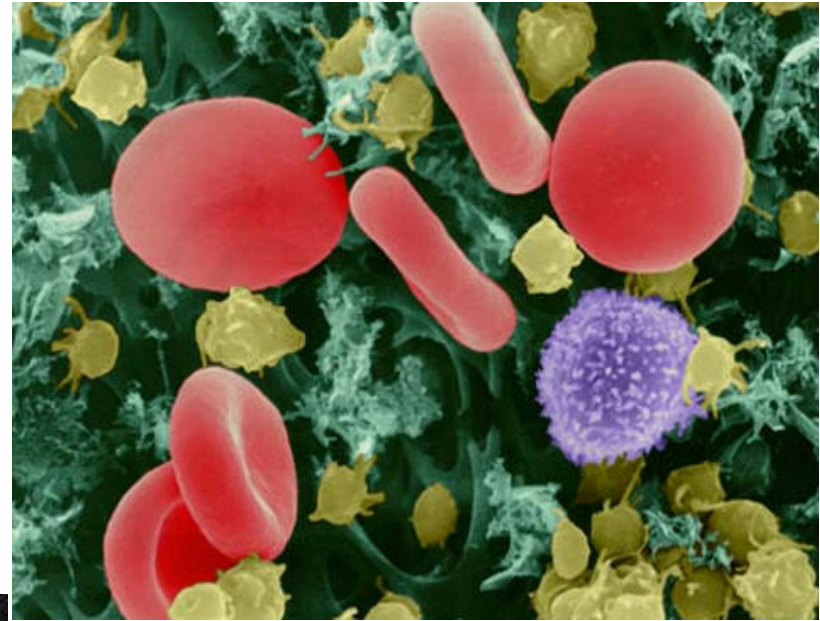


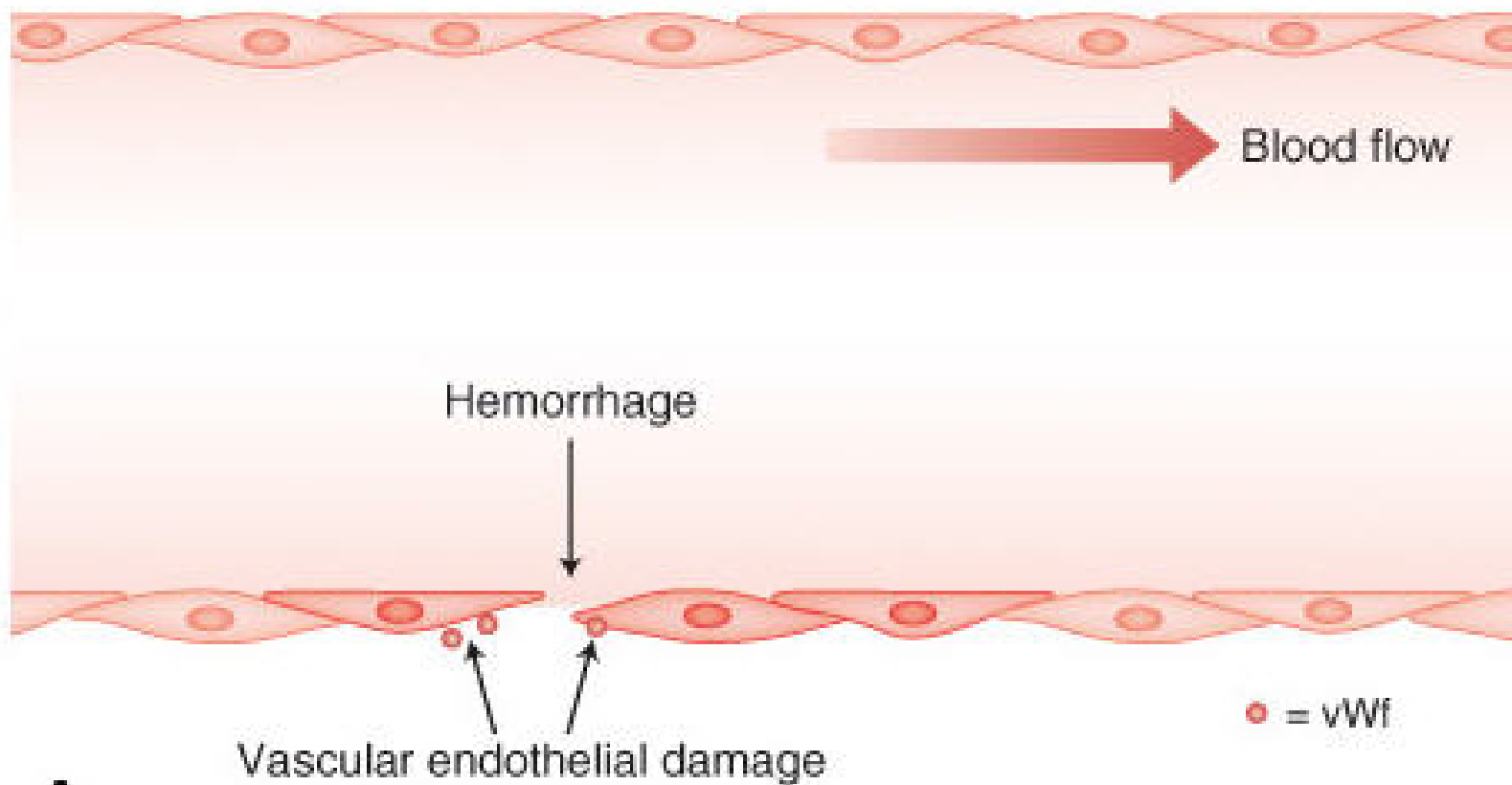
Endothelium

- ▶ Lines vessels
- ▶ Prevents coagulation
- ▶ Produces active substances

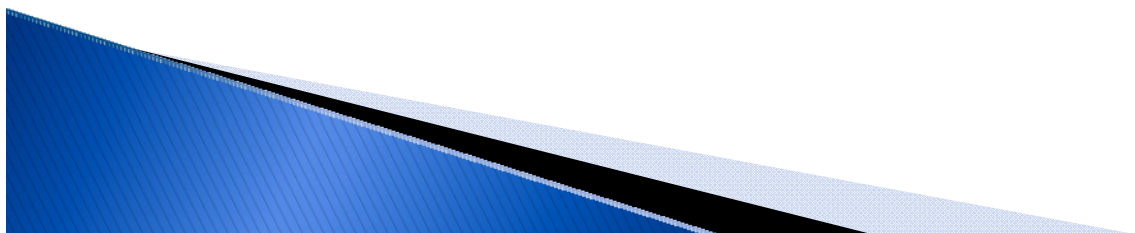


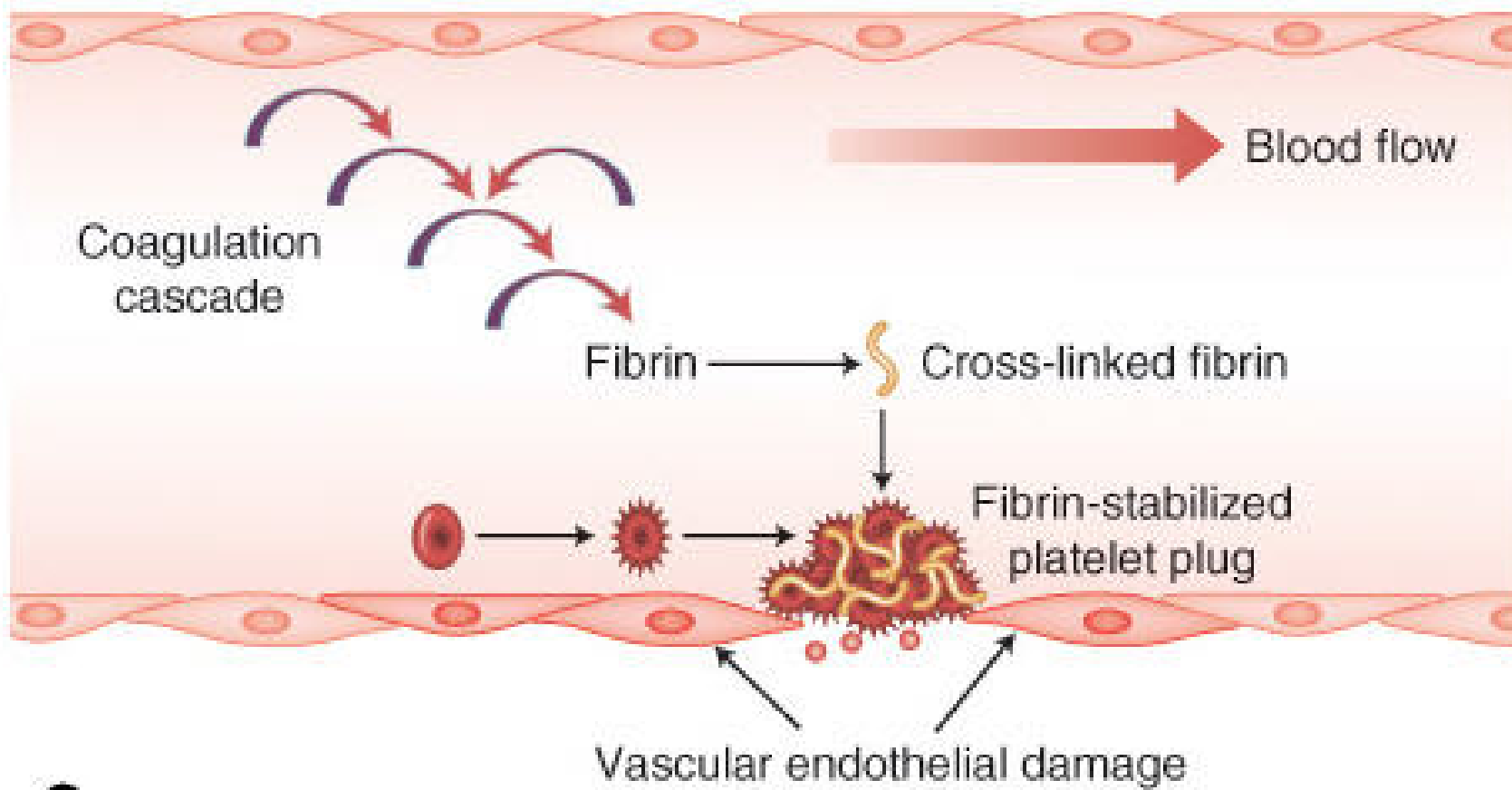
Blood Cells



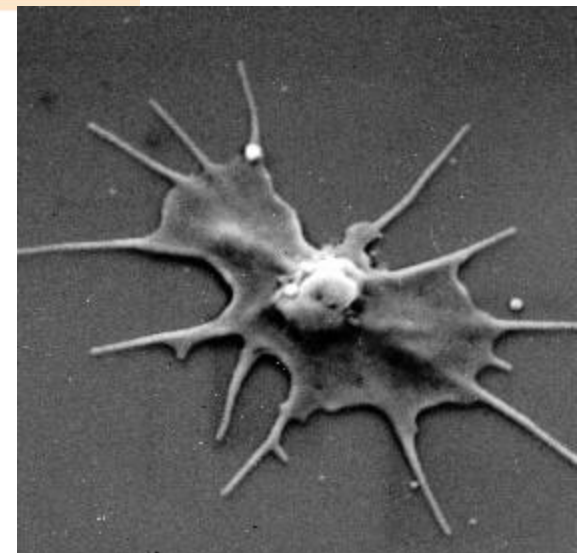
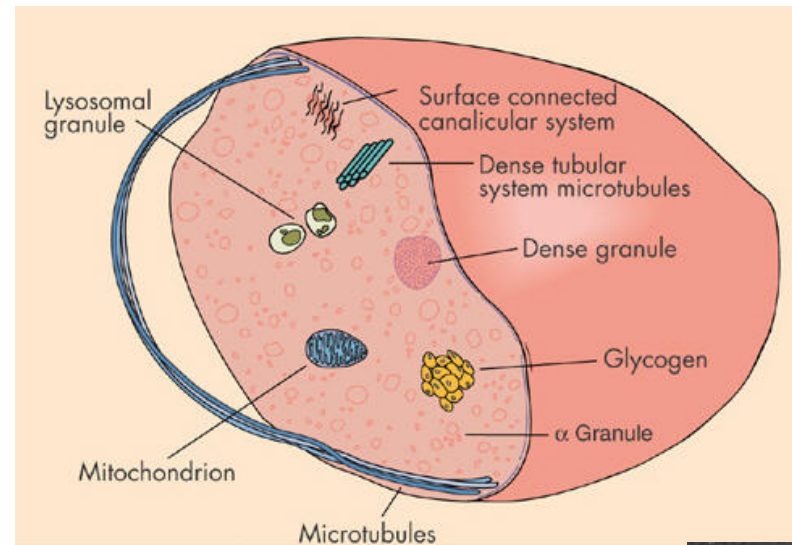
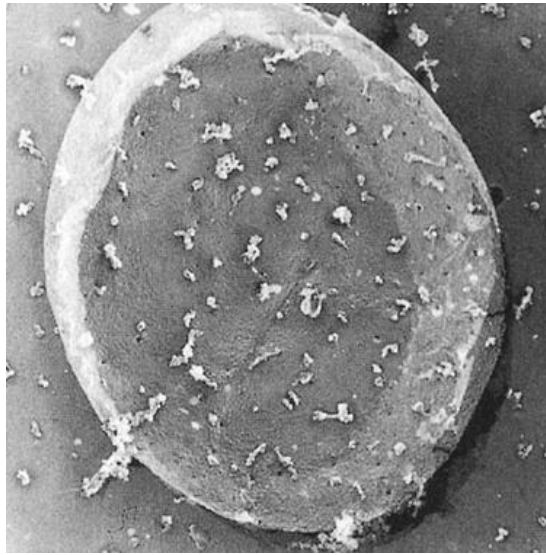


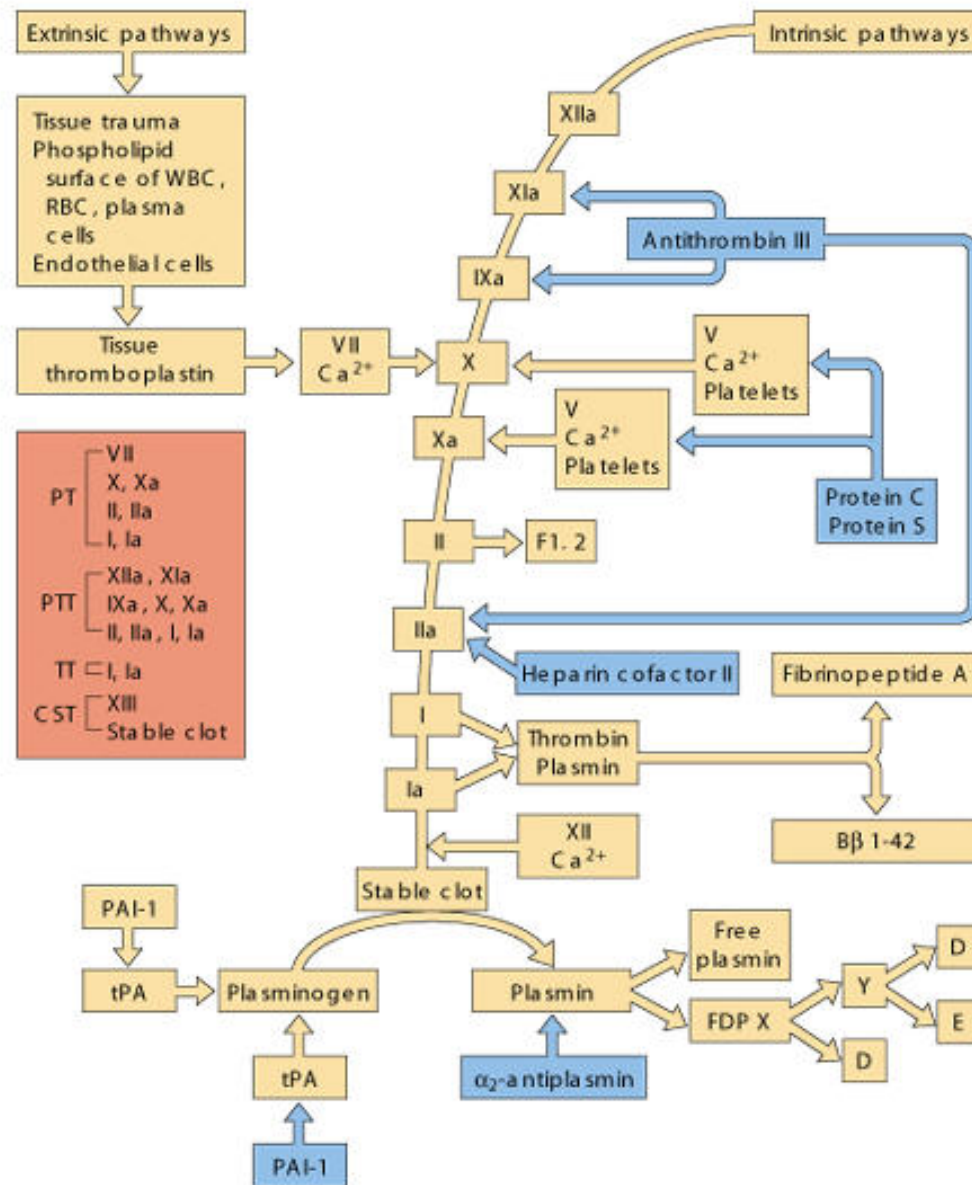
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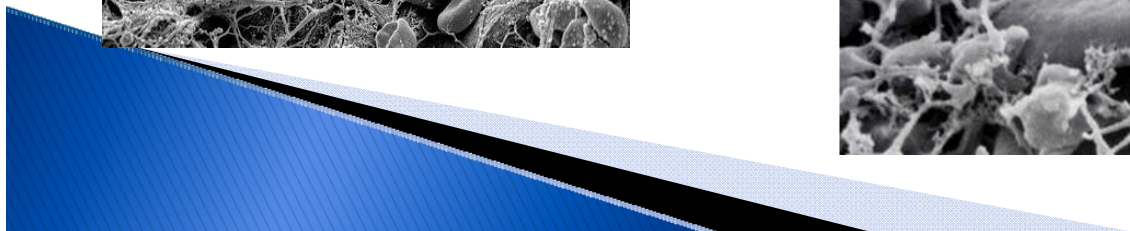
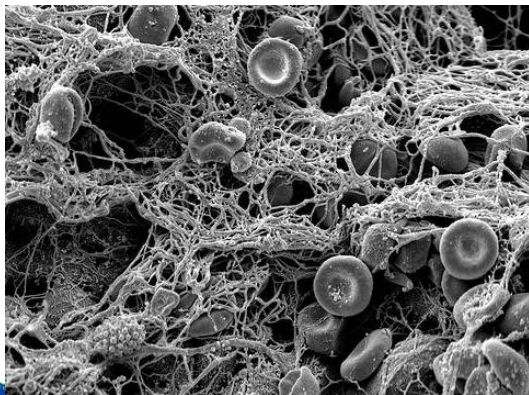
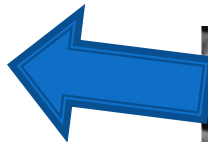
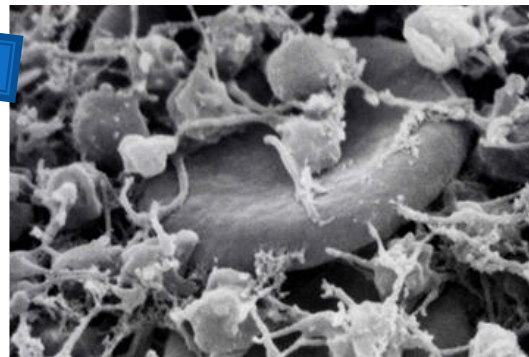
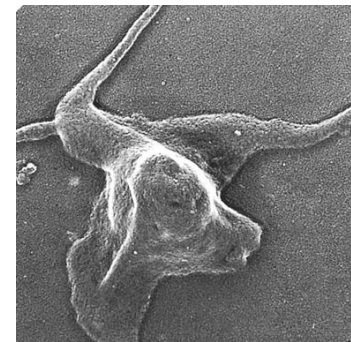
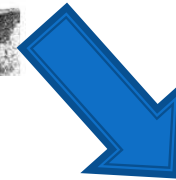
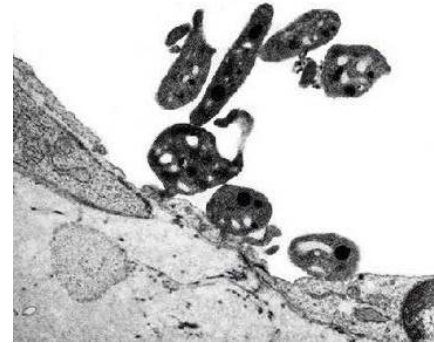
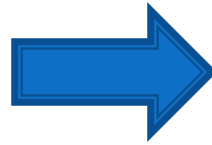
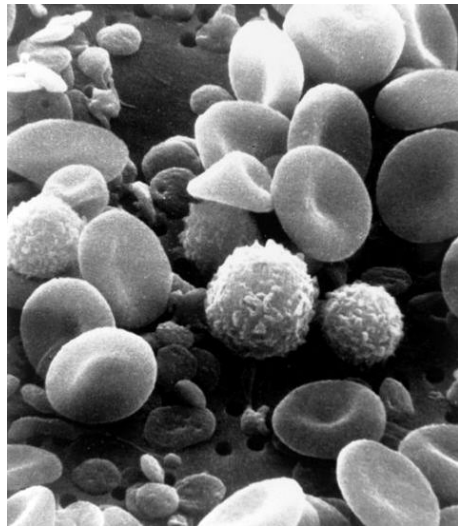




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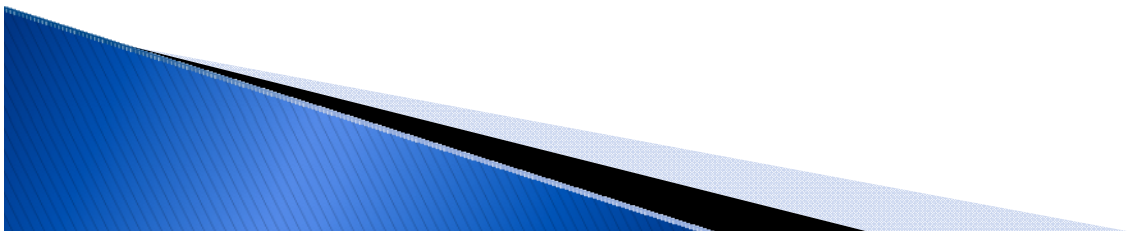






Signs of bleeding

- ▶ Blood!?
- ▶ Bruising or ecchymosis
- ▶ Petechiae
 - Pinpoint red dots– usually related to low platelets



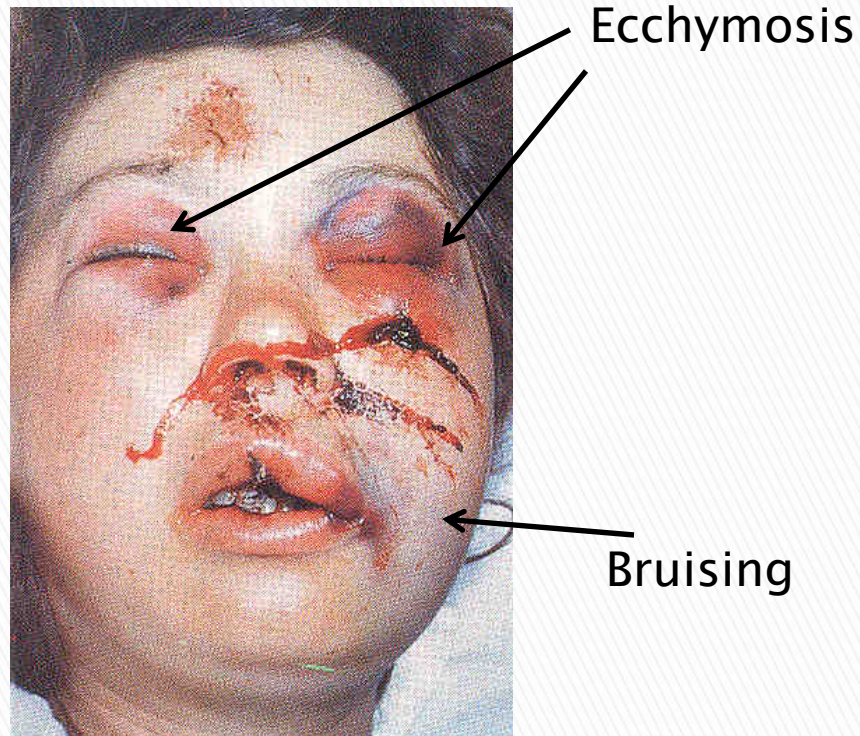
Bruising and ecchymosis

- ▶ Color varies generally with age
 - You can only tell relative not precise age
 - Sequence: Red–Blue–Brown–Green/yellow



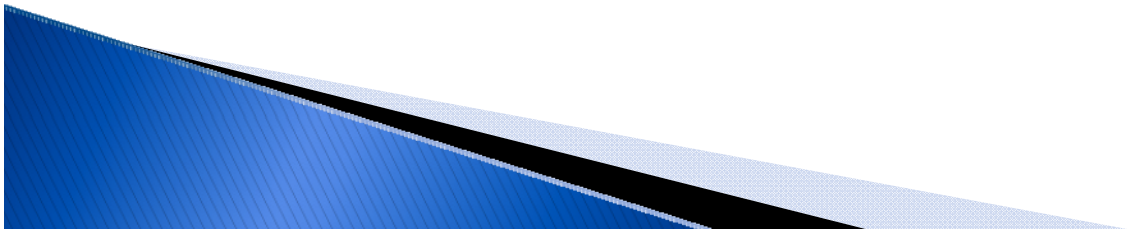
Bruising and ecchymosis

- ▶ Bruising: discoloration resulting from traumatic impact
- ▶ Ecchymosis: discoloration from migration of blood through tissues



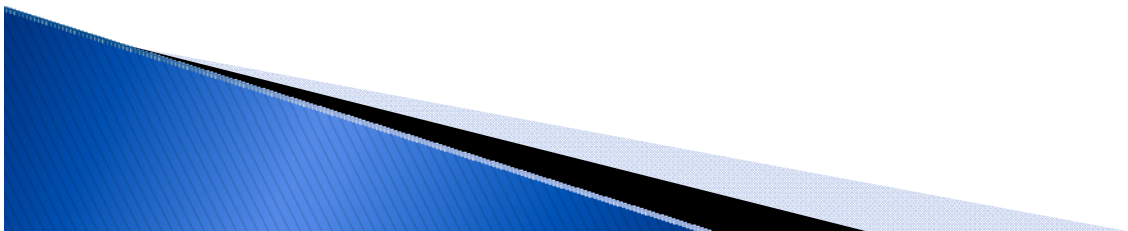
Other bleeding problems

- ▶ Nosebleed (epistaxis)
- ▶ Bleeding into joints (hemarthrosis)
- ▶ Bleeding into muscle
- ▶ Intracerebral bleed
- ▶ Excessive menses



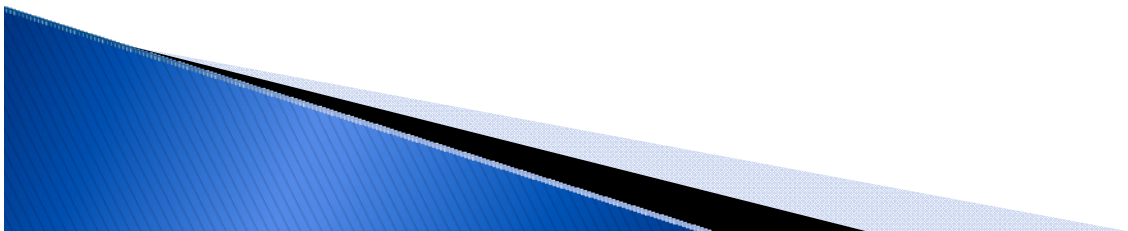
History

- ▶ Medications
 - Coumadin
 - Heparin
 - Aspirin
 - Plavix
- ▶ Known bleeding condition
- ▶ History of heavy bleeding
 - Dental work
 - Menstrual period
 - Minor surgery



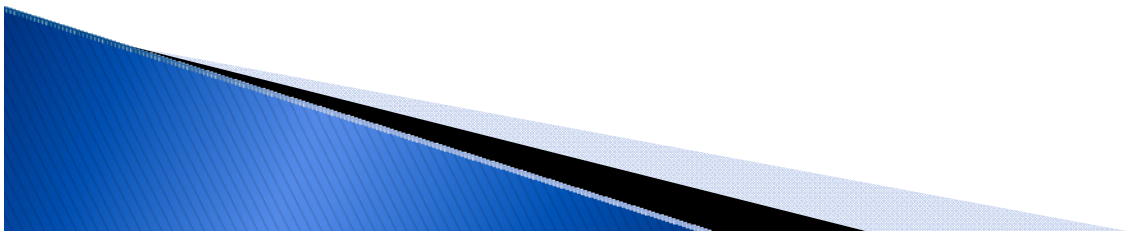
Who Bleeds?

- ▶ Congenital
 - Hemophilias
 - VonWillebrands Disease
- ▶ Acquired
 - Infection
 - ITP
 - Liver or Kidney Disease
 - Toxic exposure
 - Iatrogenic
 - Medication
 - Hypothermia
 - Dilution



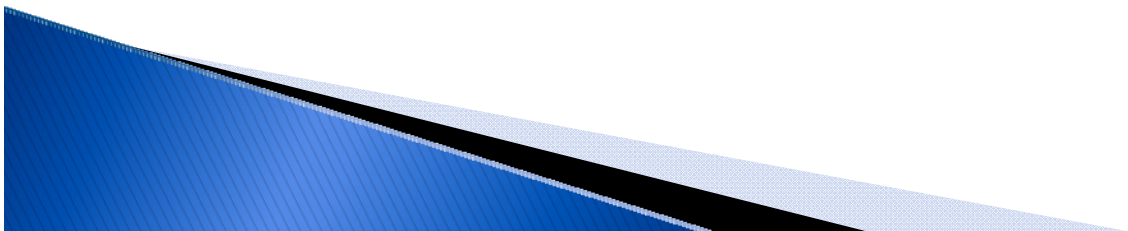
Von Willebrand's Disease

- ▶ Most common inherited defect
- ▶ Decreased or abnormal Von Willebrands Factor (required to attach platelets to endothelium and to activate)
- ▶ Usually (90–98% of time mild to moderate)
- ▶ Often appears as increased blood loss during menstruation or at time of miscarriage
- ▶ Autosomal gene– woman can have condition
- ▶ Long term effects rare



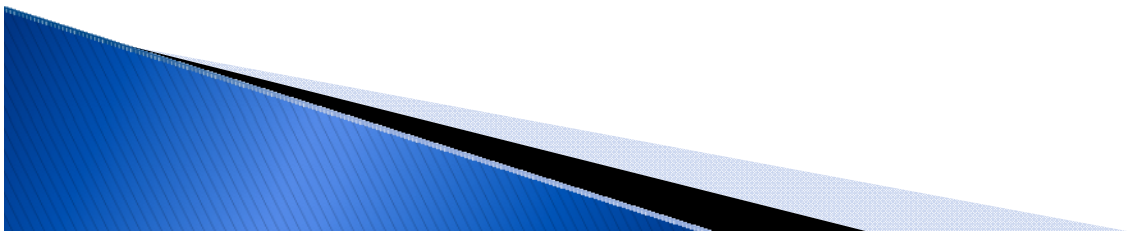
Von Willebrand's Disease– Treatment

- ▶ Usually none required unless dental procedure or surgical procedure
- ▶ Birth control pills for young women
- ▶ Vasopressin for high risk periods
- ▶ On rare occasions factor VIII or IX required
- ▶ In field usual protocols



Hemophilias

- ▶ Two major types
 - A (Factor VII)
 - B (Factor IX) – “Christmas disease”
- ▶ Sex linked recessive
 - Woman may be carriers
 - Almost all patients are males
- ▶ Level of severity depends upon amount of factor produced by patient



Hemophilias

- ▶ Significant long term affects



Retroperitoneal Hematoma



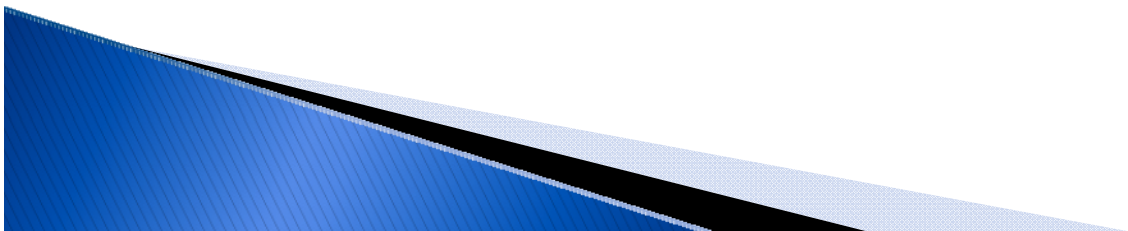
Intracranial Hemmorage



Ankle Hemarthrosis

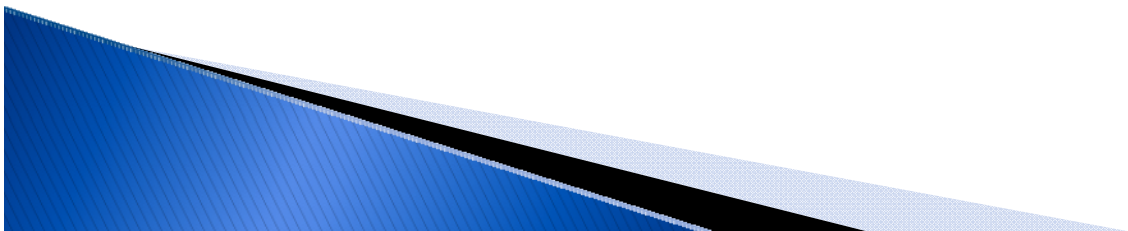
Treatment

- ▶ Prevention of contact trauma
- ▶ IV infusion of factor VIII or IX
 - In hospital
 - At home by family
 - By Patient
- ▶ Case management by HTC
- ▶ Topical agents
- ▶ Physical therapy and conditioning



Hemophilias

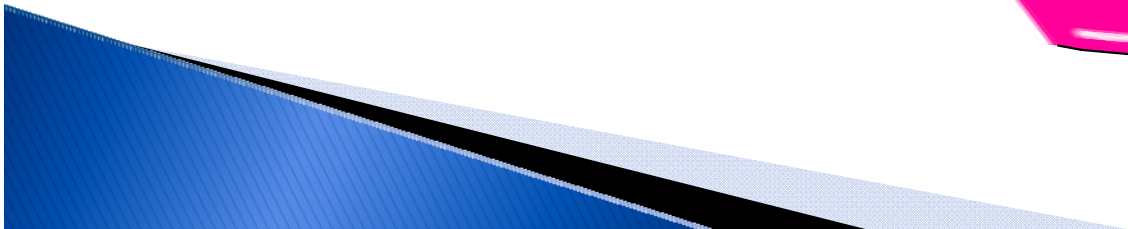
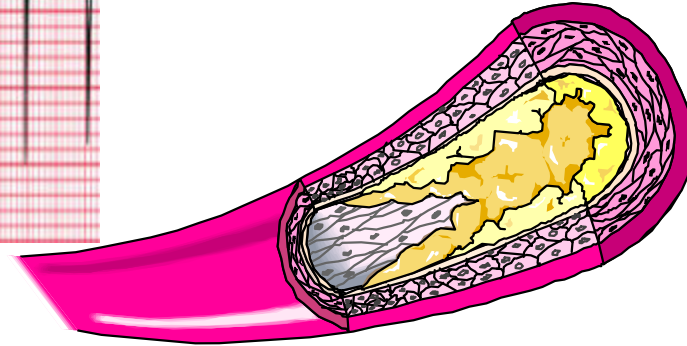
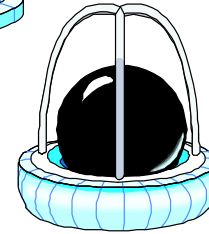
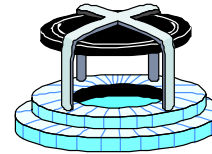
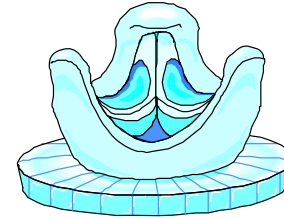
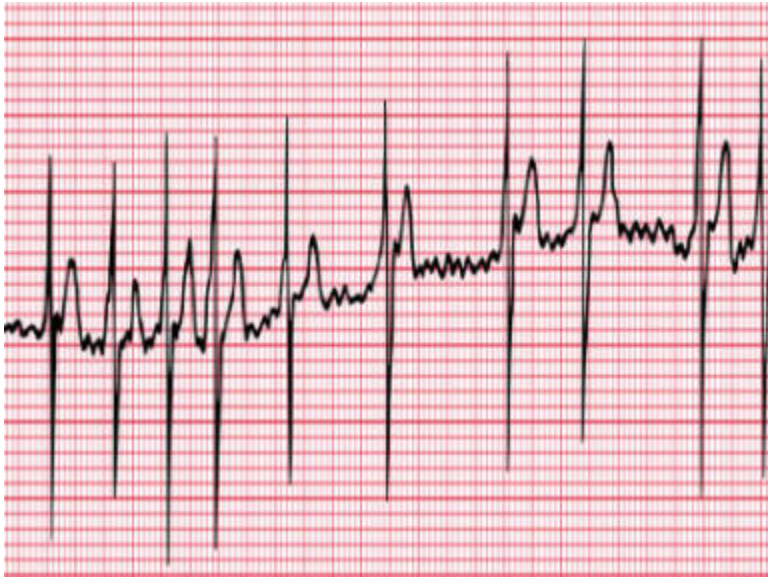
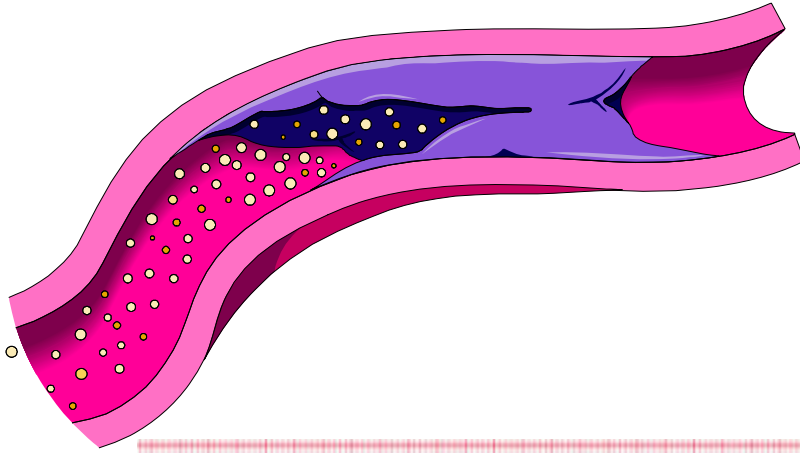
- ▶ Name and phone number of HTC
- ▶ Infusion?
 - Who?
 - Last infused?
 - When?
- ▶ Where is factor
 - Take to facility with you



Medication related

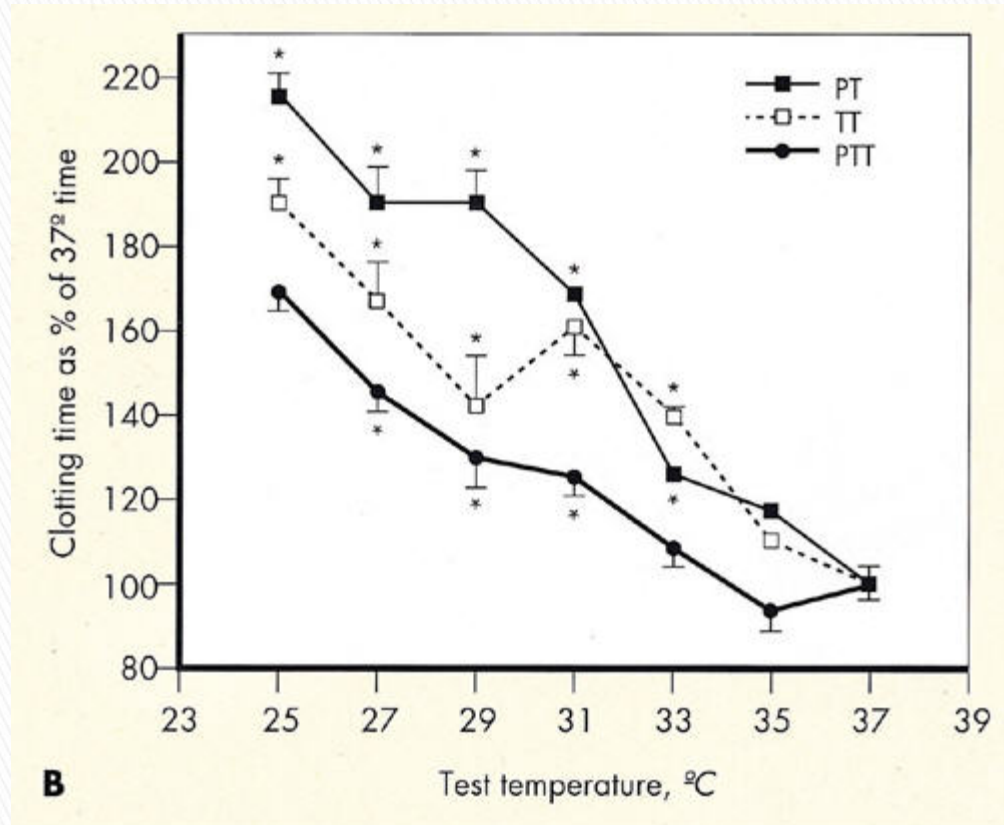
- ▶ Coumadin
 - Correct dose varies with medications and diet
 - Reverse with Vitamin K
 - Rapid reversal with coumadin
- ▶ Platelet inhibitors
- ▶ Heparin





Cold

- ▶ Accidental hypothermia
- ▶ Therapeutic



Decreased Availability

- ▶ Decreased manufacture of clotting factors
 - Liver disease–vitamin K dependent
- ▶ Decreased manufacture of platelets
 - Hypersplenism – often occurs in alcoholic cirrhosis
- ▶ Destruction of platelets
 - ITP
- ▶ Poorly functioning platelets
 - Increased BUN in kidney disease

