

## **CLASSIFICATION OF PULMONARY HYPERTENSION**

### **Group 1. Pulmonary Arterial Hypertension**

- a) Idiopathic (formerly primary PH)
- a) Congenital systemic-to-pulmonic shunts
  - i. Atrial septal defects
  - ii. Ventricular septal defects
  - iii. Patent ductus arteriosus
- b) Associated with drugs, toxins, inflammatory conditions

### **Group 2. Pulmonary Hypertension associated with left sided heart disease**

- b) Left ventricular or arterial disease
- c) Left-sided valvular disease

### **Group 3. Pulmonary Hypertension Associated with Respiratory Disease and/or Hypoxemia**

- a) Interstitial pulmonary fibrosis
- b) Chronic obstructive pulmonary disease
- c) High-altitude disease
- d) Neoplasia

### **Group 4. Pulmonary Hypertension Due to Thromboembolic Disease**

- a) Primary pulmonary arterial lesion (e.g. *Dirofilaria immitis*, *Angiostrongylus vasorum*)
- b) Pulmonary Thromboembolism
  - i. Protein-losing disease
  - ii. Neoplasia
  - iii. Immune-mediated hemolytic anemia
  - iv. Hyperadrenocorticism
  - v. Disseminated intravascular coagulation
  - vi. Sepsis
  - vii. Recent surgery
  - viii. Cardiac Disease

### **Group 5. Miscellaneous**

- a) Compressive masses (neoplasia, granuloma)