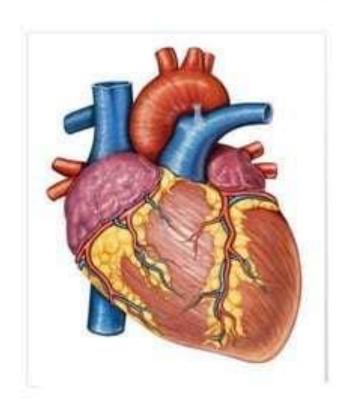
## CARDIOMYOPATHY



SUDESHNA BANERJEE DUTTA LECTURER S.R.S.V.M B.SC NURSING COLLEGE

## **CARDIOMYOPATHY**

- Disease of the heart muscle in which the heart loses its ability to pump blood effectively
- The heart muscle becomes enlarged or abnormally thick or rigid.
- In rare cases, the muscle tissue in the heart is replaced with scar tissue.

#### CLASSIFICATION

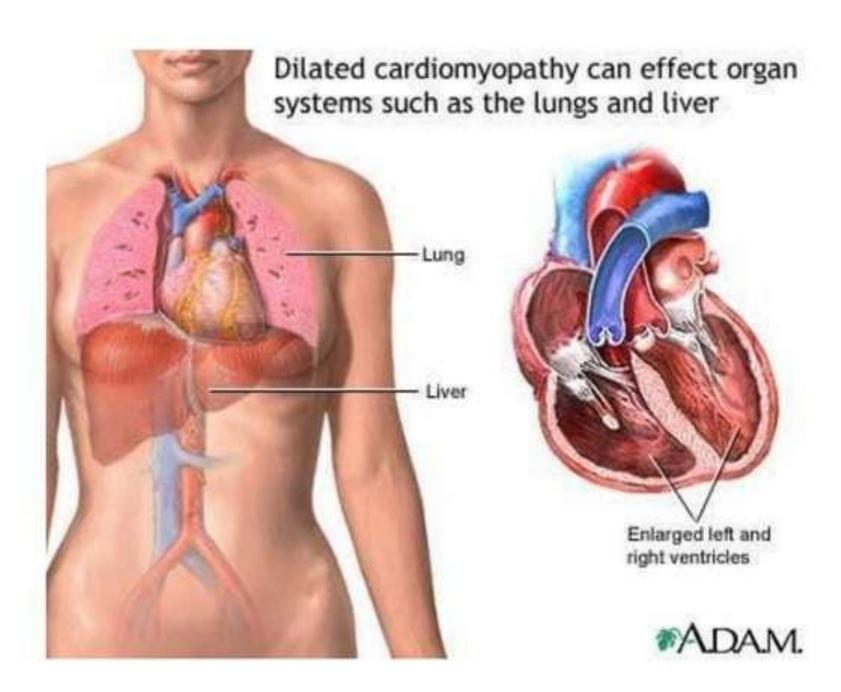
#### The 3 main types of cardiomyopathy are:

- Dilated cardiomyopathy
- ✓ Hypertrophic cardiomyopathy
- ✓ Restrictive cardiomyopathy

## Dilated Cardiomyopathy

A condition in which the hearts ability to pump blood is decreased because the hearts main pumping chamber; the left ventricle, is enlarged or weakened.

- ✓ Most common form of cardiomyopathy
- ✓ Generally occurs in adults aged 20 to 60 years
- ✓ More common in men



# Pathophysiology of Dilated Cardiomyopathy

Diffuse inflammation and rapid degeneration of myocardium

Ventricular dilation

Impairment of systolic function (impaired emptying of LV)

Left atrial enlargement and stasis of blood in the left ventricle

Cardiomegaly

#### Causes:

- ➤ Cardiotoxic agents like alcohol or coccaine
- ▶ Genetic
- >Hypertension
- ➤Ischemia(CAD)
- Muscular dystrophy (Weakening and wasting of muscles)
- ➤ Myocarditis
- ➤Valvular disease

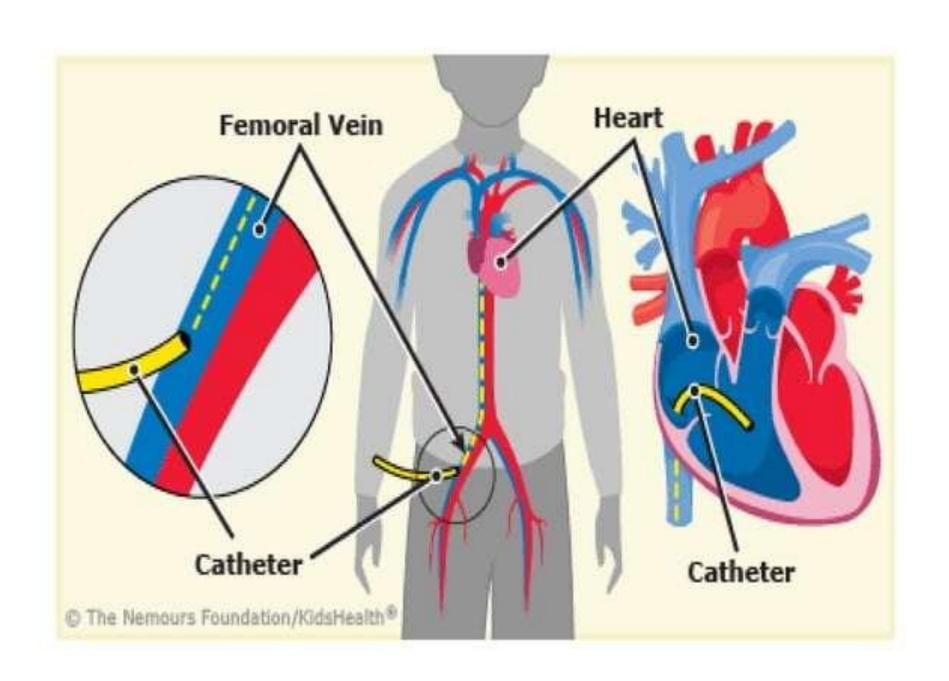
#### **Features**

- Decreased exercise capacity
- Fatigue
- Dyspnea
- Paroxysmal nocturnal dyspnea
- Orthopnea as the disease progresses
- Dry cough, palpitations
- Abdominal bloating
- Nausea, vomiting
- Anorexia

- Abnormal S3 and S4 sound
- Tachycardia or bradycardia
- ➤ Edema
- Pulmonary crackles
- Weak peripheral pulses
- Jugular venous distention

## Diagnostic tests

- ➤ History
- Echocardiography
- Chest x-ray: shows the signs of cardiomegaly
- ECG: reveals tachycardia, bradycardia and dysrythmias.
- Cardiac catheterization: it is performed to confirm CAD



## Medical management

- Nitrates: eg- isosorbitrate
- Loop diuretics: eg-furosemide
- ACE inhibitors: eg- captopril
- Beta adrenergic blockers: eg- atenolol
- Aldosterone agonists: eg- spironolactone
- Cardiac glycoside : eg-digoxin
- Anticoagulation therapy

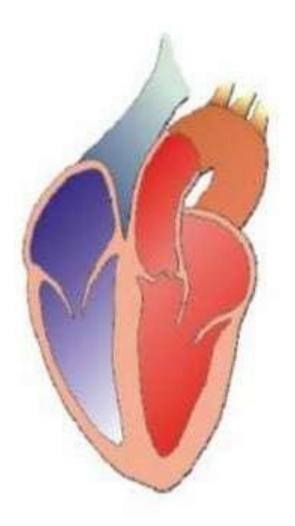
## Surgical management

Cardiac transplantation

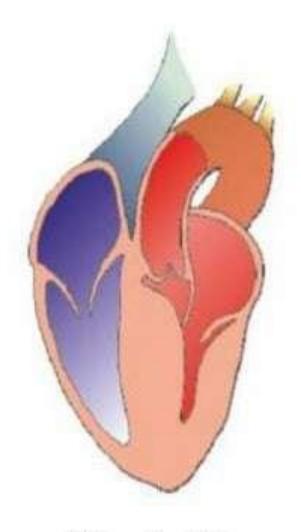
## Hypertrophic Cardiomyopathy

Assymetric left ventricular hypertrophy without ventricular dilation.

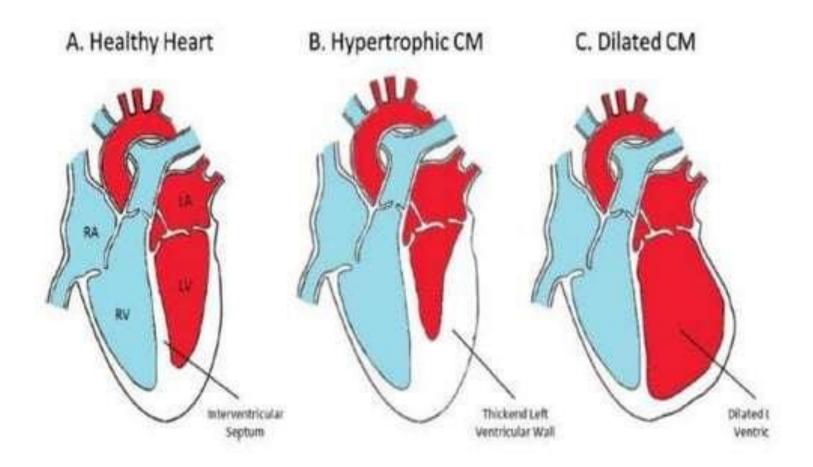
When the septum between two ventricles become enlarged and obstructs the blood flow from left ventricle, it is known as hypertrophic obstructive cardiomyopathy.



Normal heart



Heart with left ventricular hypertrophy



#### Causes

- Inherited because of a gene mutation
- Aortic stenosis
- Genetic
- Hypertension
- More common in men between ages 30 to 40

# Pathophysiology of Hypertrophic Cardiomyopathy

Thickened intra-ventricular septum and ventricular wall

Ventricular hypertrophy

Diastolic dysfunction

Impaired ventricular filling and obstruction to decreased outflow

Decreased cardiac output

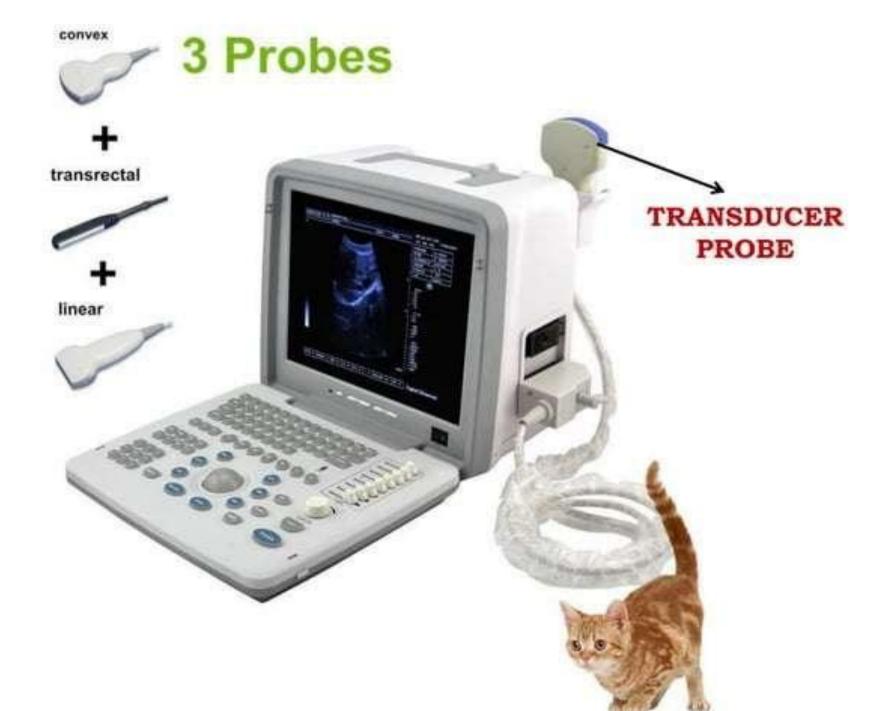
#### **Features**

- Exertional dyspnea (Shortness of breath during exercise)
- Decreased cardiac output
- Fatigue
- Angina
- Syncope
- Hypertension

## Diagnostic tests

History and physical examination

Transthoracic echocardiogram. In this test, a device (transducer) is pressed firmly against your skin. The transducer aims an ultrasound beam through your chest to your heart, producing moving images of the working of the heart.





- Electrocardiogram (ECG): Wires (electrodes) attached to adhesive pads on your skin measure electrical impulses from your heart. An ECG can detect enlarged chambers of your heart and abnormal heart rhythms.
- Cardiac MRI: A cardiac MRI uses magnetic fields and radio waves to create images of your heart.

### Medical management

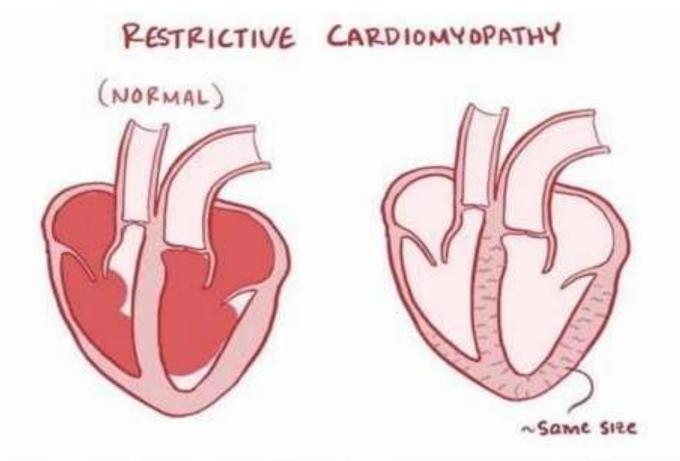
- Beta adrenergic blockers: eg- atenolol
- Calcium channel blocker: eg- verapamil
- Antidysrhythmic drugs : eg- amiodarone

## Surgical management

- Septal myectomy: It is an open heart surgical procedure in which the surgeon removes the part of thickened, over grown septum between the ventricles.
- Septal ablation: In this procedure a small portion of the thickened heart muscle is destroyed by injecting alcohol through a long, thin tube into the artery supplying blood to that area.

## Restrictive cardiomyopathy

Disease of the heart muscle that impairs diastolic filling and stretch and the systolic function remains unaffected.



## Etiology

Unknown etiology

Myocardial fibrosis, endocardial fibrosis, sarcoidosis and radiation to the thorax

## Pathophysiology

**Etiologic factors** 

Stiffness of the ventricular wall with loss of ventricular compliance

Ventricles become resistant to filling

Decrease cardiac output

#### **Features**

- Fatigue
- Exercise intolerance
- Dyspnea
- Orthopnea(shortness of breath (dyspnea) which occurs when lying flat)
- Syncope
- Palpitations
- Peripheral edema
- Jugular venous distention

### Diagnostic tests

Chest x-ray: shows cardiomegaly

ECG: shows tachycardia

Echocardiography : for the visualization of left ventricle

CT-Scan and MRI Scan

## Medical management

- Beta adrenergic blockers: eg- atenolol
- Calcium channel blocker: eg- verapamil
- Steriods: hydrocortisone
- Antidysrhythmic drugs : eg- amiodarone

## Surgical management

A heart transplantation may be considered if the heart function is very poor and the symptoms are severe

## Nursing management

- Instruct the patient to take all medicines on prescribed time.
- Encourage to use low sodium diet
- Instruct to drink more water
- Instruct the patient to maintain proper body weight
- Teach the patient to balance activity and rest
- Instruct the patient to avoid vigorous activities and exercises

- Encourage to perform stress reduction activities.
- Teach about breathing and coughing exercise
- Suggest the family members to learn about CPR.

