

**SNS COLLEGE OF ALLIED HEALTH SCIENCE**  
Affiliated to The Tamil Nadu Dr. M.G.R Medical University, Chennai



**DEPARTMENT OF CARDIAC TECHNOLOGY**

**COURSE NAME : CF & BLS**

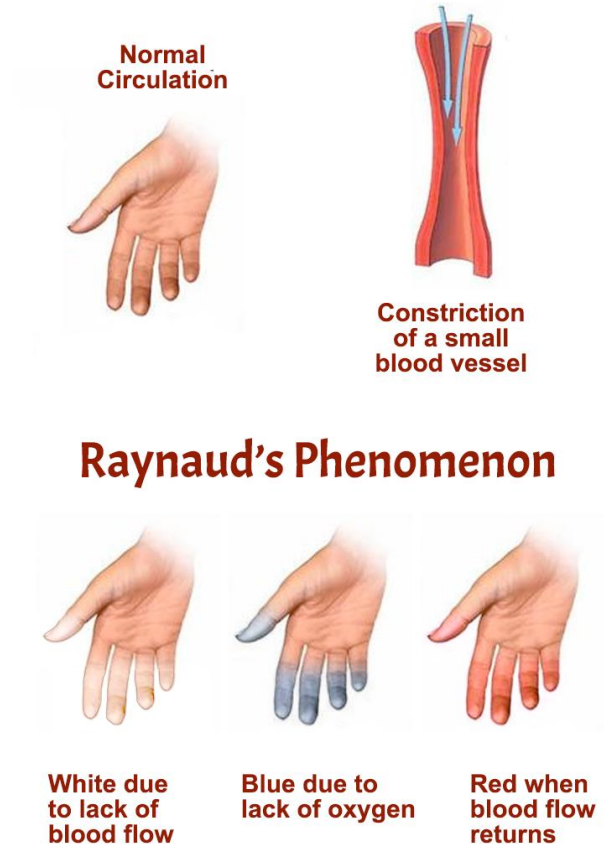
**UNIT : 1**

**TOPIC : RAYNAUD'S PHENOMENON**

**FACULTY NAME : Ms. HARSHITHA S**

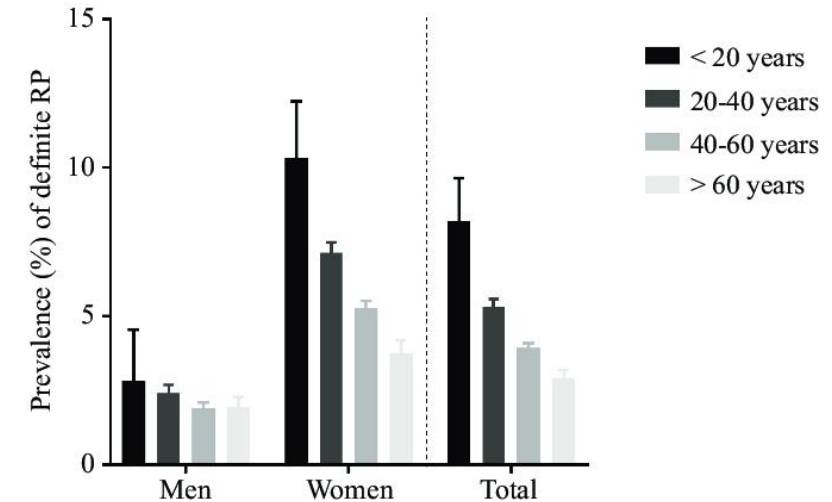
# What is Raynaud's Phenomenon?

- Episodic vasospasm of small arteries & arterioles of fingers/toes
- Triggered by cold or emotional stress
- Triphasic color change (not always present):
  1. Pallor (white) → ischemia
  2. Cyanosis (blue) → deoxygenation
  3. Rubor (red) → reactive hyperemia
- First described by Maurice Raynaud in 1862



# Epidemiology

- Prevalence: 3–20% in general population
- Primary Raynaud's: 80–90% of cases
- Female : Male = 9:1 (primary)
- Onset usually 15–30 years (primary)
- Secondary: older onset, more severe



WHITE  
POOR BLOOD CIRCULATION



BLUE  
LOW OXYGENATION



RED  
BLOOD RETURNS AND MAY HURT

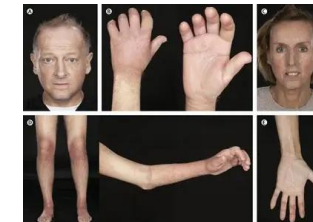
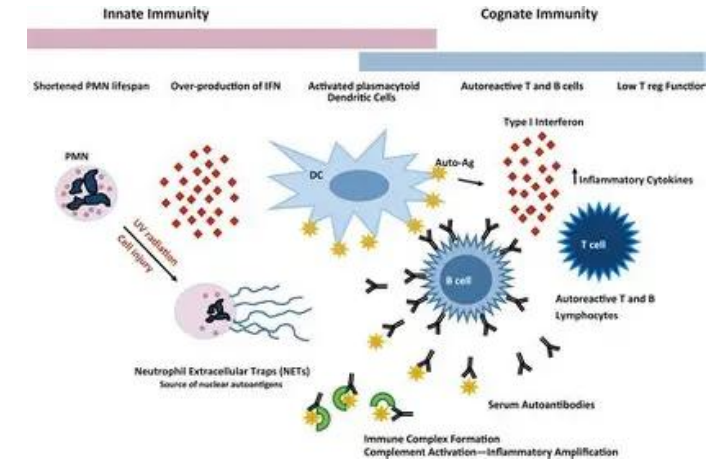


# Classification

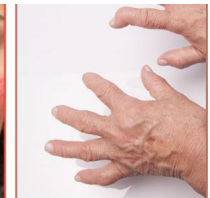
Feature	Primary Raynaud's	Secondary Raynaud's
Age of onset	<30–40 years	>30–40 years
Symmetry	Symmetric	Often asymmetric
Severity	Mild	Severe, digital ulcers, gangrene
Autoantibodies	Negative	Positive (ANA, ENA, etc.)
Capillaroscopy	Normal	Abnormal (SD pattern, scleroderma)
Associated disease	None	Connective tissue diseases, etc.

# Common Causes of Secondary Raynaud's

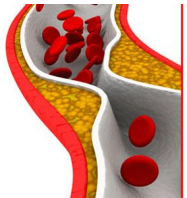
- Connective tissue diseases (90% of secondary cases)
  1. Systemic sclerosis (SSc) – strongest association
  2. SLE, Sjögren's, mixed CTD, dermatomyositis
- Vibration injury (vibration white finger)
- Drugs:  $\beta$ -blockers, ergotamine, amphetamines, vinblastine
- Hypothyroidism, cryoglobulinemia, malignancy



Lupus



Rheumatoid Arthritis



Atherosclerosis

# Clinical Features – Classic Triad

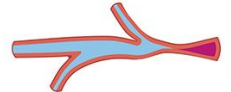
1. Pallor (white) → fingers/toes turn dead white
  2. Cyanosis (blue) → dusky bluish hue
  3. Hyperemia (red) → bright red flushing on rewarming
- Attacks last 5–30 minutes
  - Numbness, tingling, burning pain on rewarming
  - Spares the thumb in early disease



WHITE  
POOR BLOOD CIRCULATION



BLUE  
LOW OXYGENATION

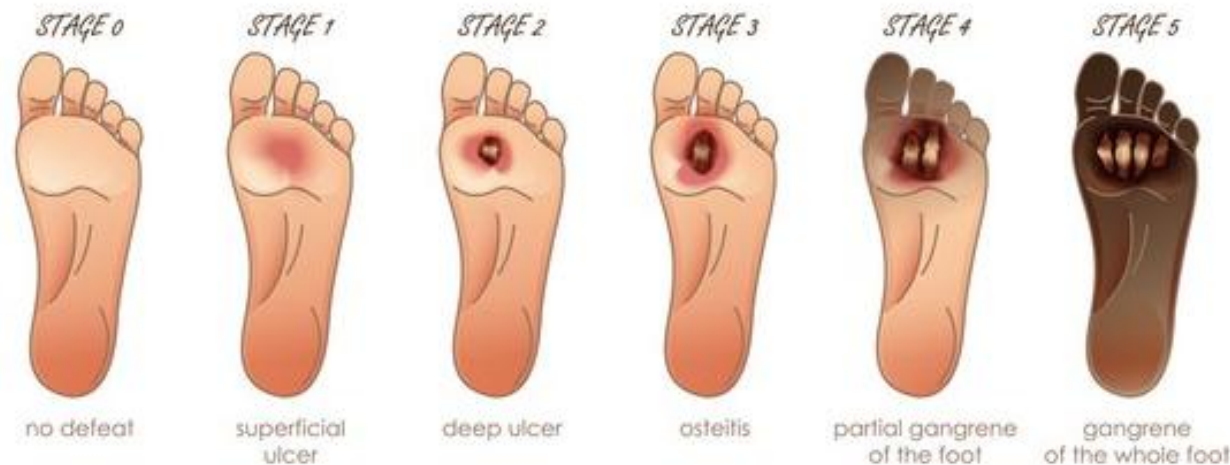


RED  
BLOOD RETURNS AND MAY HURT



# Severe Manifestations (Secondary)

- Critical ischemia
- Digital pitting scars
- Ulceration
- Gangrene & auto-amputation (rare)

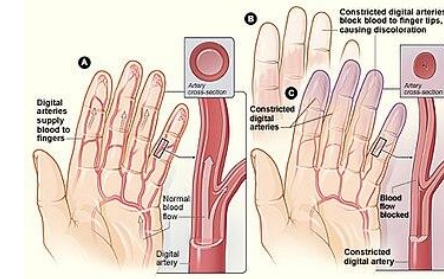




# Diagnosis

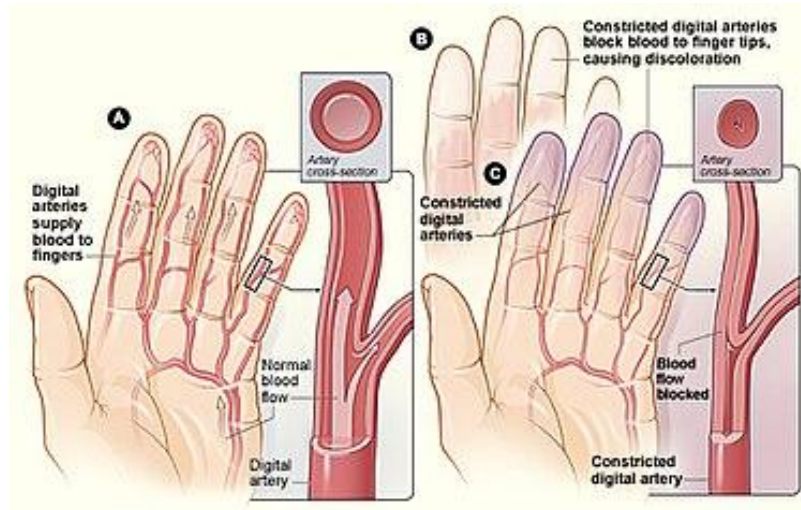
## Clinical Criteria (Allen & Brown 1932, updated)

- Primary Raynaud's diagnosis requires ALL:
- Episodic biphasic or triphasic color changes triggered by cold/stress
- Symmetric involvement
- No evidence of tissue necrosis/ulceration
- No secondary cause on history/exam
- Normal nailfold capillaroscopy
- Negative or low-titer autoantibodies





# Diagnosis



## Three-step approach to diagnosis of Raynaud's Phenomenon

### Step 1: Ask screening Question

Are your fingers unusually sensitive to cold?

Yes, proceed to step 2

### Step 2: Assess color changes

Occurrence of biphasic color changes during the vasospastic episodes (white and blue)

Yes, proceed to step 3

### Step 3: Calculate disease score

- a) Episodes are triggered by things other than cold (i.e. emotional stressors)
- b) Episodes involve both hands, even if the involvement is asynchronous and/or asymmetric
- c) Episodes are accompanied by numbness and/or paresthesias
- d) Observed color changes are often characterized by a well-demarcated border between affected and unaffected skin.
- e) Patient provided photograph(s) strongly support a diagnosis of RP.
- f) Episodes sometimes occur at other body sites (e.g. nose, ears, feet, and areolas).
- g) Occurrence of triphasic color changes during the vasospastic episodes (white, blue, red)

If 3 or more criteria met from Step 3 (a - g), then the patient has RP.

# Key Investigations

Test	Purpose	Typical Finding in Secondary
Full blood count	Anaemia, thrombocytopenia	May be abnormal
ESR/CRP	Inflammation	Raised
ANA (IF)	Screening for CTD	Positive $\geq 1:160$ (SSc pattern)
ENA (anti-centromere, Scl-70, etc.)	Specific CTDs	Positive
Nailfold capillaroscopy	Gold standard for distinguishing primary vs secondary	Giant capillaries, avascular areas in SSc
Thermography	Objective documentation of attacks	Prolonged rewarming time

# Management – Non-Pharmacological (First Line)

## Patient education & lifestyle modification (cornerstone)

- Keep core body & extremities warm (gloves, hats, heated clothing)
- Avoid sudden cold exposure
- Stop smoking (vasoconstrictor)
- Minimize stress (biofeedback, relaxation techniques)
- Avoid trigger drugs ( $\beta$ -blockers, clonidine)

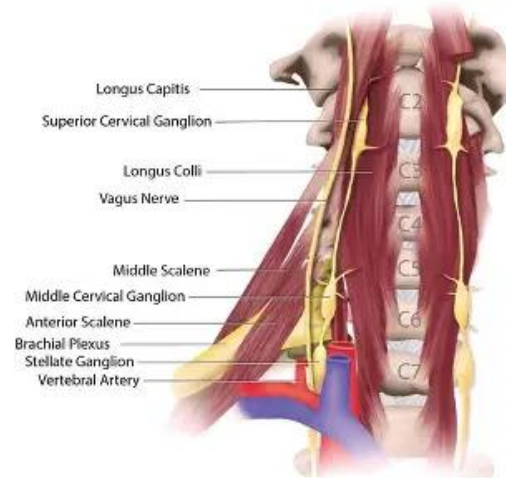
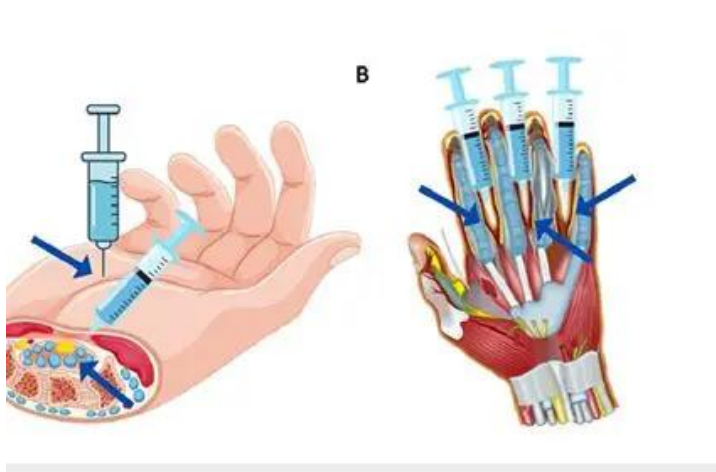
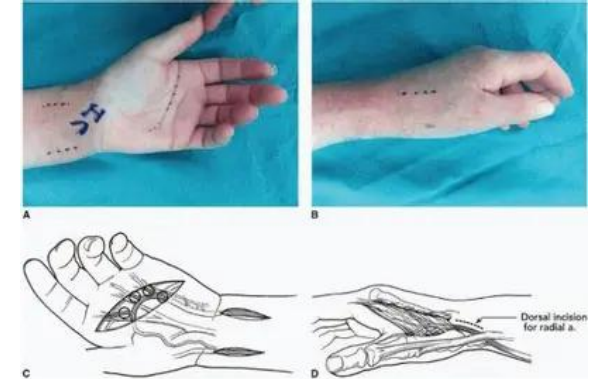


# Pharmacological Treatment

Indication	Drug of Choice	Dose	Notes
Mild-moderate (primary)	Calcium channel blockers	Nifedipine XL 30-90 mg/day	First-line, 70% response
Alternative/adjunct	Losartan (ARB)	50-100 mg/day	Better tolerated than CCBs
Refractory	PDE-5 inhibitors	Sildenafil 25-50 mg TDS	Especially in SSc with ulcers
Severe/refractory	IV Iloprost (prostacyclin)	0.5-2 ng/kg/min × 3-5 days	Gold standard for critical ischemia/ulcers
Refractory (rare)	Bosentan (endothelin antagonist)	62.5-125 mg BD	Prevents new digital ulcers in SSc

# Surgical & Advanced Options

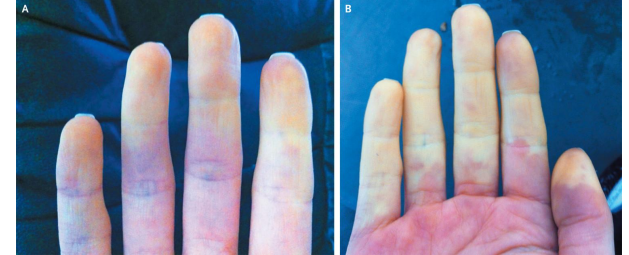
- Digital sympathectomy (last resort)
- Botulinum toxin injection into affected digits
- Cervical/thoracic sympathectomy (temporary benefit)
- Stem cell therapy (experimental)





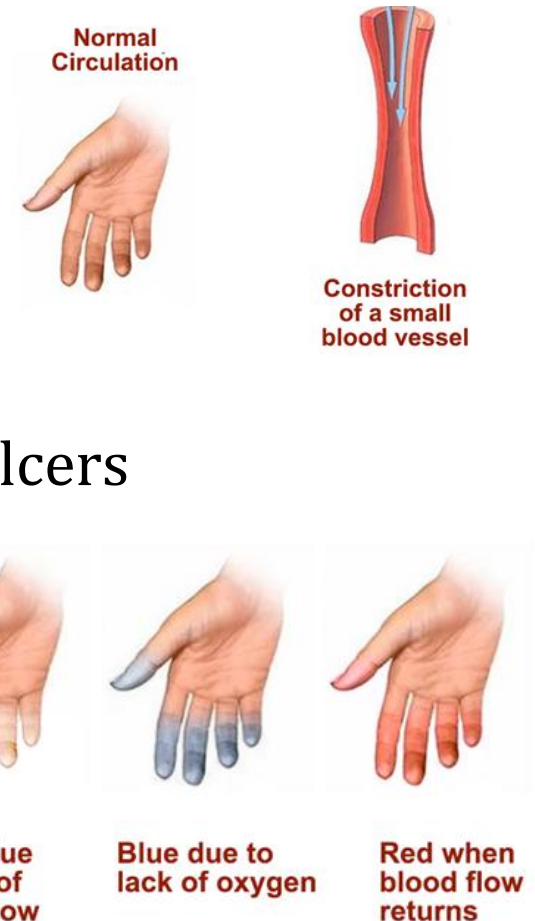
# Prognosis

- Primary Raynaud's: benign, rarely progresses
- Secondary Raynaud's:
  1. SSc patients: 50% develop ulcers within 10 yrs
  2. 10-year risk of gangrene ~5–10% in severe SSc
- Early diagnosis + aggressive treatment of underlying CTD improves outcomes



# Summary

- Raynaud's is common; most cases are primary & benign
- Asymmetric, severe, late-onset → investigate for secondary cause
- Nailfold capillaroscopy + ANA are the two most useful tests
- Lifestyle measures are first-line; CCBs are first drug
- Iloprost is the most effective therapy for critical ischemia/digital ulcers
- Treat the underlying disease in secondary Raynaud's







**THANK YOU**