

Cyanosis

Dr Jon Bramall

Consultant in Anaesthesia and Intensive Care Medicine

CAP10 Cyanosis

The trainee will be able to assess a patient presenting with cyanosis to produce a valid differential diagnosis, investigate appropriately, formulate and implement a management plan		
Knowledge	Assessment Methods	GMP Domains
Know the causes of cyanosis, cardiac & respiratory	E, C, Mi, ACAT	1
Know how to formulate a differential diagnosis and be able to differentiate from methaemoglobinaemia	E, C, Mi, ACAT	1
Skills		
Perform a full clinical examination differentiating between the various causes of cyanosis	E, C, D	1
Be able to perform and interpret the appropriate tests, e.g. x-rays and ECG	E, C, D	1
Understand the safe prescribing of oxygen therapy	E, C	1
Behaviour		
Involve senior promptly in event of significant airway compromise	ACAT, C	2
Involve specialist team as appropriate	ACAT, C	2

What colour
do Smurfs go
when you
choke them?

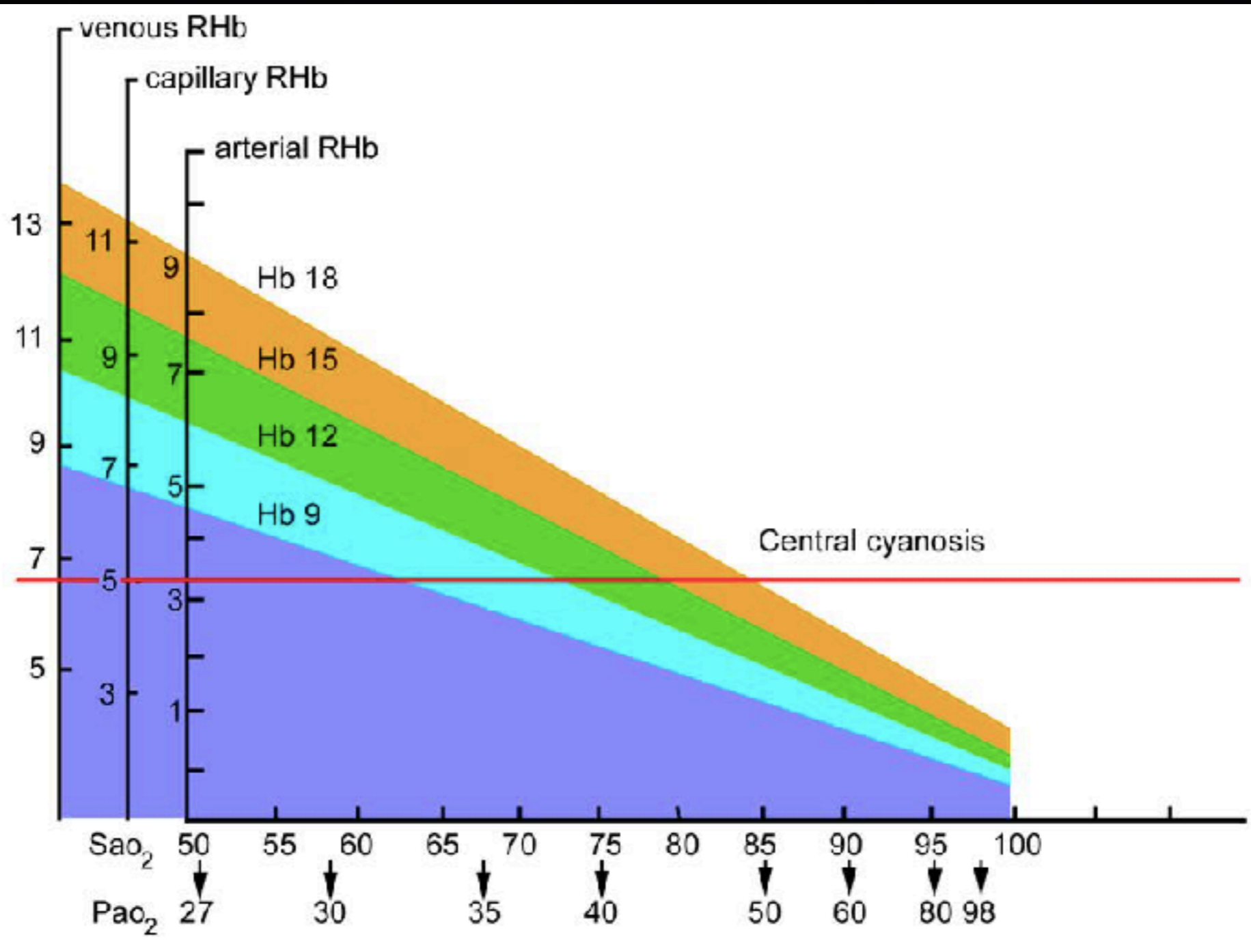


“Cyanosis is defined as the bluish or purplish discolouration of the skin or mucous membranes due to the tissues near the skin surface having low oxygen saturation”

–Wikipedia

Cyanosis

**Bluish hue that occurs in the presence of
~60 g/L deoxyhaemoglobin or
dyshaemoglobinaemias**

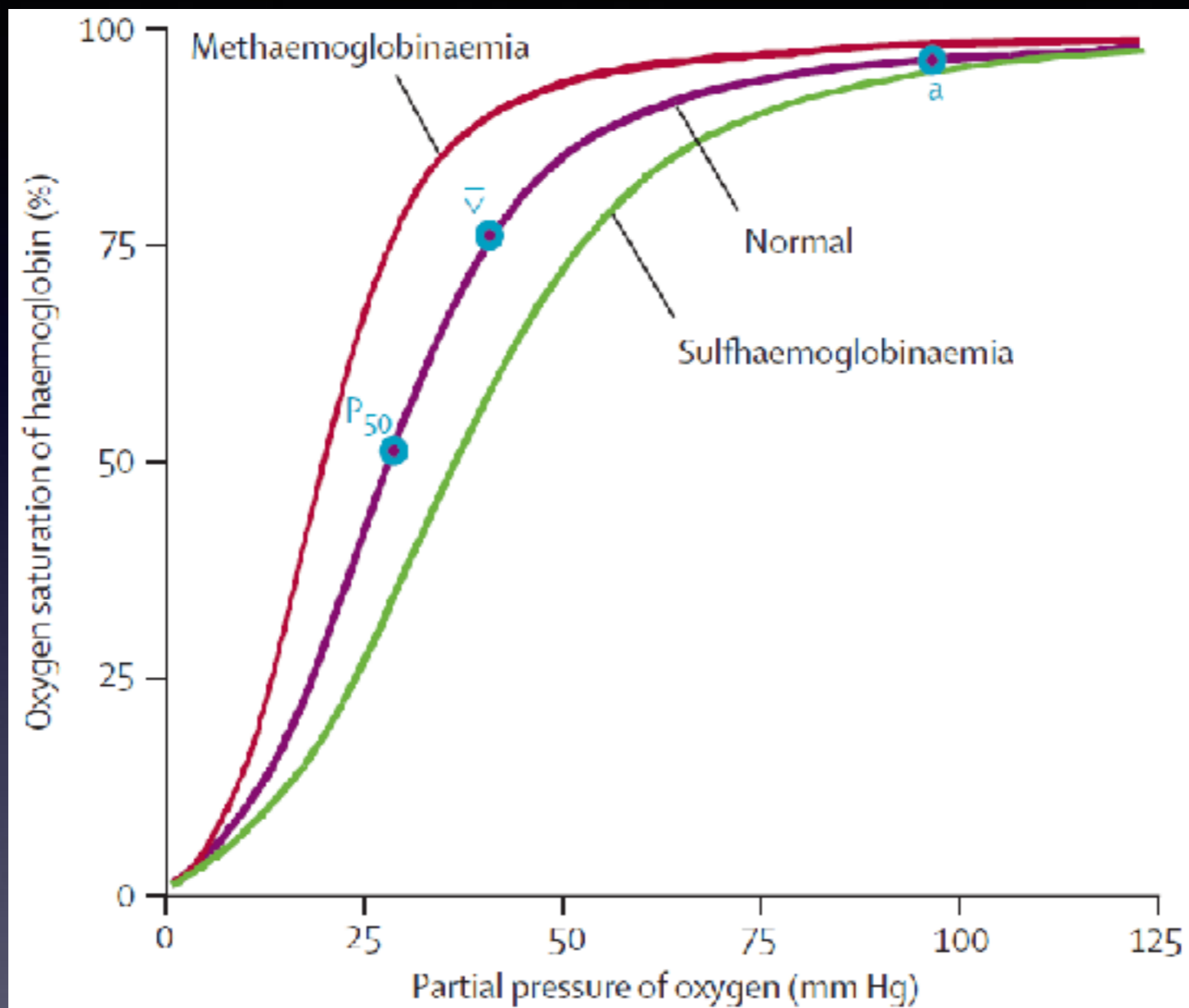


Cyanosis Causes

- Diseases causing hypoxaemia
- Shunt
- Peripheral
- Abnormal Haemoglobin

Peripheral Cyanosis

- Normal PaO₂ and Normal SpO₂?
- Causes:
 - Central causes
 - Reduced Cardiac output
 - Cold
 - Redistribution
 - Arterial or venous obstruction



Oxygen Content

$$\text{CaO}_2 = \begin{array}{l} \text{Oxygen carried by blood} \\ 1.34 \times \text{Hb} \times \text{SaO}_2 \end{array} + \begin{array}{l} \text{Oxygen in solution} \\ 0.01 \times \text{PaO}_2 \end{array}$$

Methaemoglobinaemia

- Fe^{2+} oxidised to Fe^{3+}
- Unable to bind to O_2
- Normal level $>1.5\%$

Methaemoglobinaemia

Causes

Congenital

- cytochrome b5 reductase deficiency
- haemoglobin M disease

Acquired (toxin/drugs)

- aniline dyes
- benzene derivatives
- chloroquine
- dapsone
- prilocaine
- metoclopramide
- nitrites (nitroglycerin, NO, sodium nitroprusside)
- sulphonamides

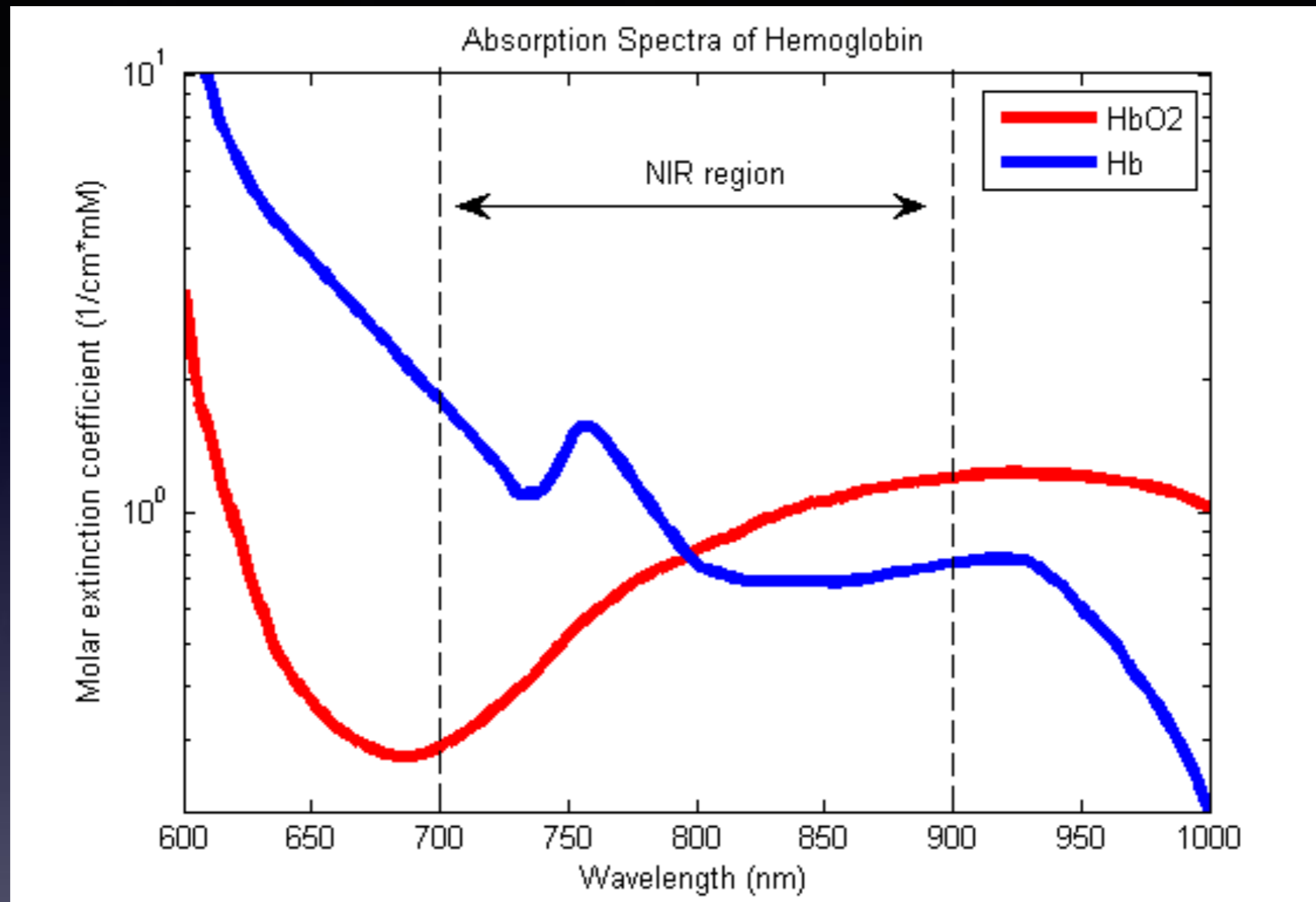
Methaemoglobinemia

Clinical Features

- cyanosis!
- symptoms and signs of decreased oxygen delivery
 - e.g. chest pain, dyspnea, altered mental state, end organ damage
- SpO₂ reading 85-90%
- blood samples typically have a chocolate brown hue
- Normal PaO₂

Methaemoglobinemia Management

- High flow O₂
- Stop precipitants (or avoid if congenital)
- methylene blue (1-2mg/kg over 5 minutes)
 - provides an artificial electron acceptor to facilitate the reduction of MetHb via the NADPH-dependent pathway
- Alternatives:
 - ascorbic acid (if methylene blue contra-indicated, e.g. G6PD deficiency)
 - exchange transfusion
 - hyperbaric oxygen



How does a sats probe work?

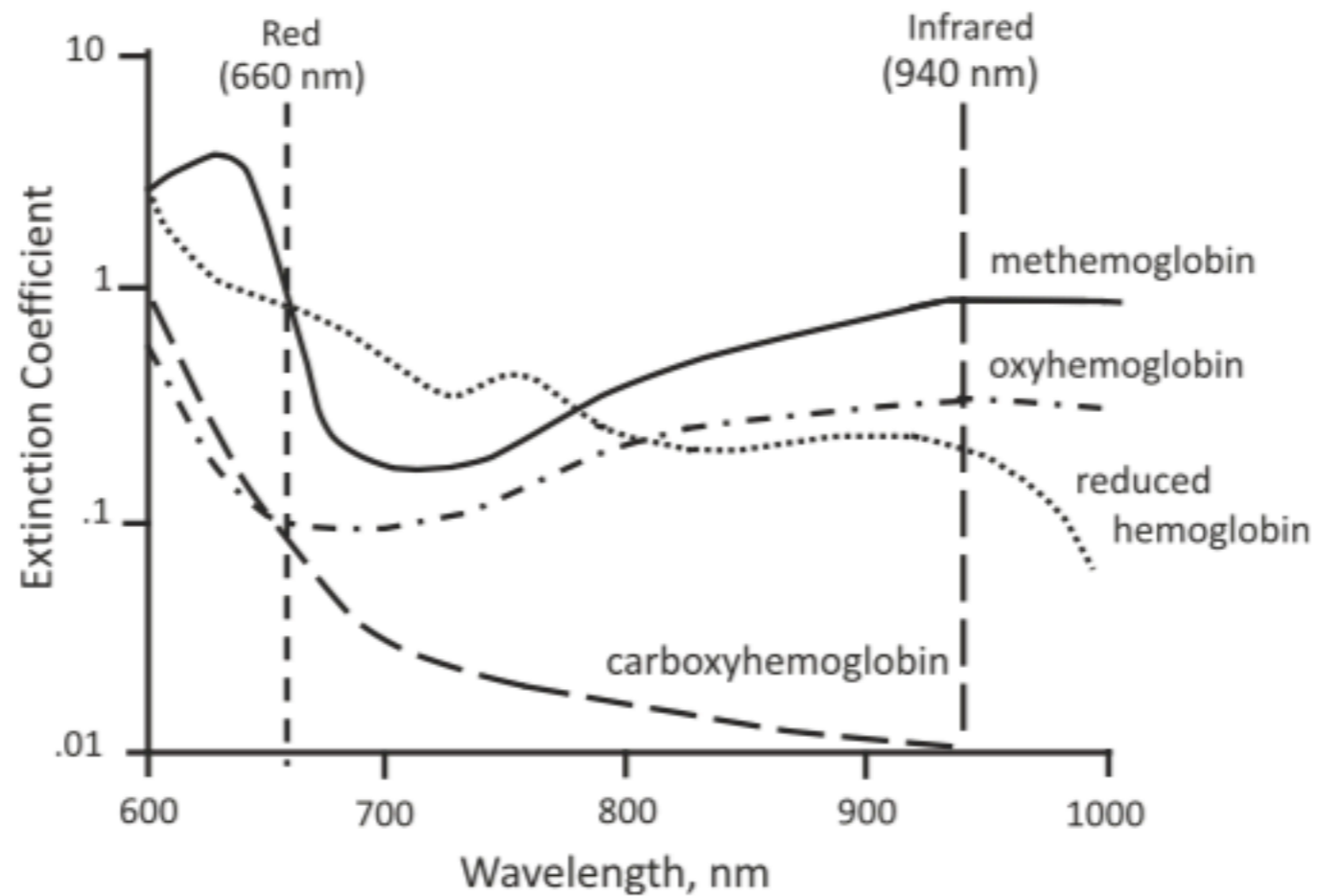


Fig. 1 Transmitted light absorbance spectra of four hemoglobin species: oxyhemoglobin, reduced hemoglobin, carboxyhemoglobin, and methemoglobin

How does a sats probe work?

Pseudocyanosis



Summary

- Cyanosis occurs when there is 60g/L deoxyhaemoglobin
- Hb 8 cyanosis at SpO₂ 60%; Hb18 SpO₂ 87%
- Causes include resp and cardiac causes of hyoxaemia
- TREAT THE CAUSE
- Don't forget blue dye, abnormal haemoglobins and pseudo cyanosis
- Methaemoglobinaemia causes "cyanosis"
 - Treat with Methylene blue

What Questions
do you have?