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**DEPARTMENT OF CARDIO PULMONARY PERFUSION CARE**  
**TECHNOLOGY**

**COURSE NAME : PRINCIPLES OF PERFUSION TECHNOLOGY I**

**II YEAR**

**TOPIC : HYPOTHERMIA**



# HYPOTHERMIA



- Hypothermia is defined as a decrease in a core body temperature below ***37 degree Celsius.***
- The principle reason for hypothermic CPB is to protect the heart and other organs by ***reducing metabolic rate and thus oxygen requirements.***
- In the myocardium, hypothermia sustains intracellular reserves of high – energy phosphates and reserves high intracellular pH and electrochemical neutrality.



# HISTORY OF HYPOTHERMIA



- In **1950, Bigelow** (canada) first demonstrated the linear relationship between falling temperature and falling metabolic rate when anaesthesia was used to control shivering and the increased muscle tone generated in response to cold.
- In **1952, Lewis & Taufic** used surface cooling to 28°C with 5.5 minutes of inflow occlusion to facilitate successful closure of an atrial septal defect in a 5 year old child.



# HISTORY OF HYPOTHERMIA



- Hypothermia equipment, used at the University of *Colorado Medical Center* in 1953, is in the Smithsonian Institution in Washington, D.C., as a medical “landmark”
- **1953 Swan** (US) experimented with hypothermia further, and used this knowledge to the success of his first open heart surgery.
- **1955 Cooley** (US) first use of hypothermia for cerebral protection during first aortic arch aneurysm repair with a homograft.



# HISTORY OF HYPOTHERMIA



- **1955 Lillehei and Kirklin** (US) noticed and published that better outcomes occurred when body temperature cooled spontaneously during oxygenation.
- After development of the pump oxygenator by **Gibbon**, cpb and hypothermia were combined by **Sealy in 1958**.
- **1959 Sealy** (US) continued Lillehei and Kirklin's development and added a heat exchanger to a Dewall oxygenator to use hypothermia alongside it.



# HISTORY OF HYPOTHERMIA



- **1963, Barnard and Schrire** (south africa) First used DHCA and CPB at the same time on an ascending and arch aortic aneurysm.
- **1975 Griep** (US) used surface cooling with CPB to resect aortic arch aneurysms in four patients.



# USES OF HYPOTHERMIA



- Major vascular diseases (aortic aneurysm)
- Removal of hepatic and renal tumour
- Intracranial surgery
- Multiple cerebral diseases
- Better myocardial protection
- Better organ preservation
- Lower pump flows – less blood trauma
- Safety margin during CPB
- Bloodless and motionless surgical field.



# MYOCARDIAL COOLING



Myocardial cooling can be achieved with,

- ***Cold cardioplegia***
- ***Pouring cold tropical solution on the heart***
- ***Cooling jackets as well as by systemic hypothermia***





## Q10 EFFECT



- ***For every one degree reduction in temperature there is reduction of 7% of metabolic needs.***
- Systemic O<sub>2</sub> consumption is reduced by approximately 50% for every 7 degree Celsius reduction in core temperature below hypothermia.
- ***For 30° Celsius = 50%***
- ***For 23° Celsius = 25%***
- ***For 16° Celsius = 12.5%***



# CLASSIFICATION OF HYPOTHERMIA



- ***Mild***            ***36 - 34 ° c***
- ***Moderate***    ***33 - 28 ° c***
- ***Deep***            ***27 - 18 ° c***
- ***Profound***      ***< 18 ° c***



# METABOLIC LEVEL IN HYPOTHERMIA



## HYPOTHERMIA

## METABOLIC RATE

- *Mild* 80%
- *Moderate* 61%
- *Deep* 41%
- *Profound* 18 - 7 %



# TEMPERATURE MONITORING SITES IN PATIENTS



Temperature should be monitored in multiple sites they are,

- *Nasopharynx*
- *Tympanic membrane*
- *Pulmonary artery*
- *Bladder*
- *Rectum*
- *Distal esophagus*



# TEMPERATURE MONITORING SITES IN HEART LUNG MACHINE



In CPB circuit the temperature probes are placed at,

- *The venous inlet*
- *The arterial outlet*
- *The CPDS*



# TEMPERATURE MONITORING SITES AND ITS ACCURACY



- ***Nasopharyngeal temperature probes*** underestimate but ***approximate to brain temperature***, with the mixed venous temperature on the CPB circuit being an approximation of average body temperature.
- ***Nasopharyngeal and tympanic membrane*** probes shows ***surface temperature***.



# TEMPERATURE MONITORING SITES AND ITS ACCURACY



- ***Bladder and Rectal temperature*** given an ***indication of core body temperature***, but these can be erroneous due to interference from varying urine production and faecal matter, respectively. These low blood flow sites tend to underestimate temperature so are particularly valuable following deeper levels of hypothermia.



## Temperature monitoring

Measuring site	Advantages	Limitations
Nasopharyngeal	easy to introduce	measurement errors due to leakage of air; nosebleeds; the core temperature is not measured
Oesophageal	reliable	dislocation; gastric catheters
Rectal	easy to introduce	not always accurate; the faeces act as insulation; the core temperature is not measured
In the bladder	can be used during both general and locoregional anaesthesia	the urine flow affects the temperature
Tympanic	can be used during both general and locoregional anaesthesia	potentially traumatic; unreliable if not inserted by an expert
Axillary	easy to introduce	only moderately reliable; the core temperature is not measured
Via the blood, tip of a Swan-Ganz catheter or CVC, the pulmonary artery	the core temperature	only during invasive pressure measurement







# BLOOD GAS FLOW RATIO



TEMPERATURE	GAS / BLOOD FLOW RATIO	FI02
37°C	1:1	80
34°C	8:1	70
30°C	7:1	65
28°C	6:1	60
22°C	5:1	50



# SAFE ARREST PERIOD FOR TEMPERATURE



## *TEMPERATURE*

32<sup>o</sup>c

28<sup>o</sup>c

18<sup>o</sup>c

< 18<sup>o</sup>c

## *ARREST PERIOD*

under 10 minute

10 to 15 minute

16 to 45 minute

46 to 60 minute



# GRADIENT OF TEMPERATURE MANAGEMENT



## *Temperature gradient between Hemotherm & pt. blood*

Adult            10 - 12° c

Paediatric    8 - 10° c

## *Gradient for temperature management*

Cooling 1° c for 1 minute

Rewarming 1° c for 3 minute



# CHANGES IN OXYGEN – HEMOGLOBIN DISSOCIATION CURVE



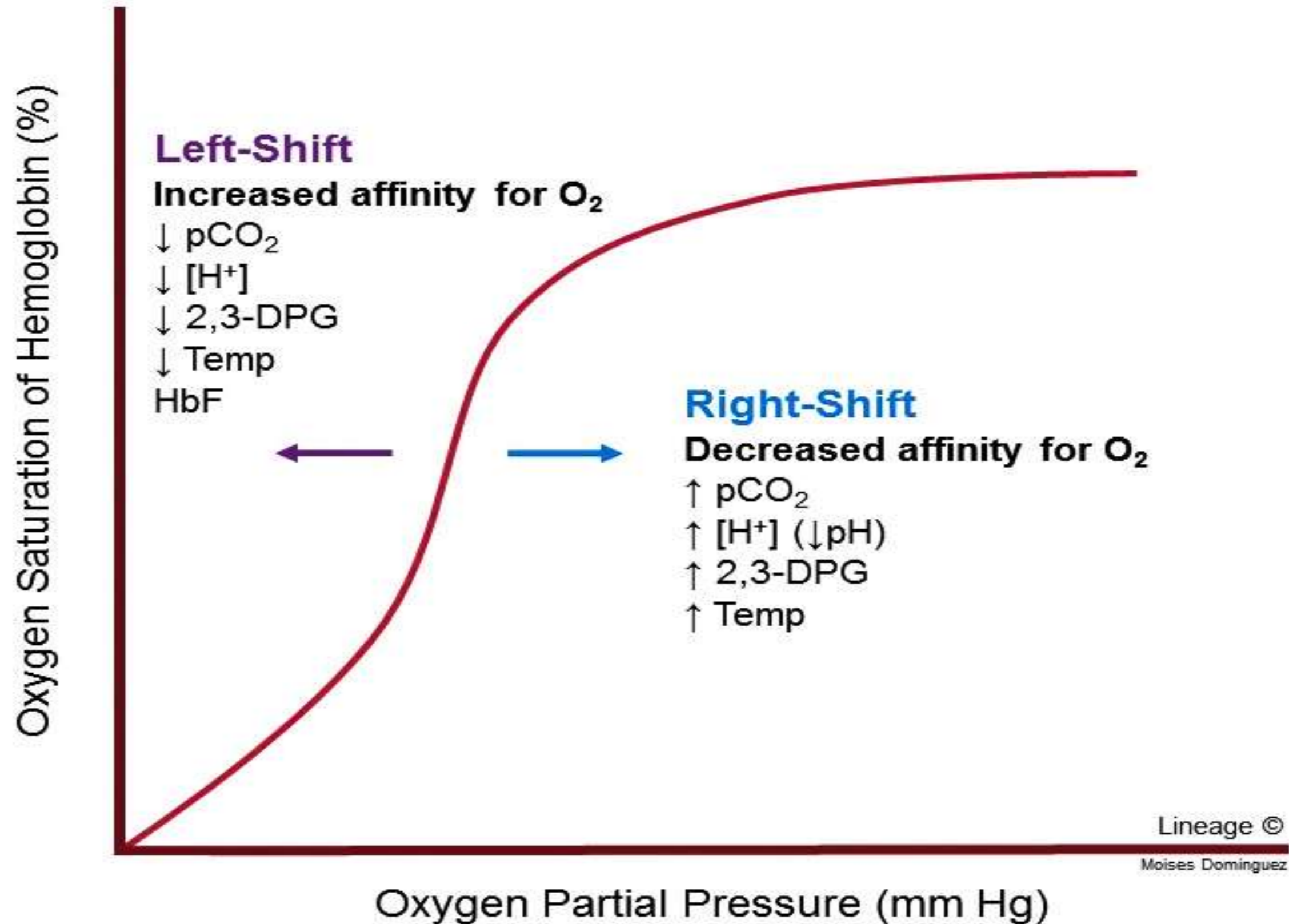
- *As the temperature decreases, the affinity or strength of binding between O<sub>2</sub> and Hemoglobin increases.  
( a lower partial pressure of O<sub>2</sub> is required to force a given amount of O<sub>2</sub> into hemoglobin molecule)*
- The oxygen – hemoglobin dissociation curve is shifted to left. Release of O<sub>2</sub> from hemoglobin at the tissue level is less efficient.



# CHANGES IN OXYGEN – HEMOGLOBIN DISSOCIATION CURVE



Oxygen-Hemoglobin Dissociation Curve





# CHANGES IN SOLUBILITY OF O<sub>2</sub> AND CO<sub>2</sub>



- As temperature decreases, gases become more soluble in liquid. For a given amount of O<sub>2</sub> and CO<sub>2</sub> more gas will be dissolved in the plasma and the partial pressure of the gas will decrease.
- This is much more significant for CO<sub>2</sub> because it is more soluble in plasma at any given temperature.



## MECHANISM OF ACTION



- Reduction in cerebral metabolism (CMRO<sub>2</sub>) by approximately 7% per 1° C. This leads to less oxygen and glucose consumption.
- Promotion of cerebral vasoconstriction, which can directly decrease ICP. Also, vascular permeability and therefore edema formation is decreased.
- Prevention of neuronal injury leading leading to programmed cell death (apoptosis) mainly by inhibition of caspase activation.



# MECHANISM OF ACTION



- Decrease the free radical formation
- Suppression of inflammatory cascade and decreased nitric oxide, cytokine and leukotriene production. Leukocyte migration from the damaged endothelium is diminished.





# ADVERSE EFFECT



- Mostly thrombocytopenia by reversible sequestration of platelets in portal circulation.
- RBC aggregate and leads to reduced perfusion to capillaries.
- Tissue H<sub>2</sub>O content increases due to hemodilution which can cause swelling and edema.
- O<sub>2</sub> availability decrease because of left shift of O<sub>2</sub> Hb curve.



# ADVERSE EFFECT



- It can cause coagulopathy disorder (eg) cold agglutination.
- Capillary resistance increases due to increase HCT have low / cerebral circulation. So, hypothermia can be performed slowly with low HCT.
- Importantly rewarming should not result in hyperthermia even 1<sup>o</sup>c to 2<sup>o</sup>c increase in brain temperature can exacerbate ischemic cerebral damage.



# SYMPTOMS FOR DECREASED BODY TEMPERATURE



BODY TEMPERATURE	SYMPTOMS
36° C	Normal core temperature
35° C	Vasoconstriction (peripheral), maximal shivering, speech disorders and hyperreflexia
34° C	Still conscious, but movements are difficult
33 - 31° C	Retrograde amnesia, no shivers, hypotension and dilation of pupils
30 - 28° C	Loss of consciousness, muscular rigidity, bradycardia and bradypnea
27 - 25° C	Loss of reflexes, ventricular fibrillation and suspended animation
17° C	Iso – electric ECG



# *Rewarming*



***Passive external rewarming***



***Active external rewarming***



***Active Internal rewarming***





# RATE OF REWARMING



- Maintain a gradient of nasal to water bath of at least 6<sup>o</sup>c in adult and 4<sup>o</sup>c in paediatrics.
- Warm at a rate of 1<sup>o</sup>c for 3 – 5 mins
- Plan accordingly with surgeon, consulting with anaesthetist.
- Consider – Bicarb, Mannitol, ZBUF ,CUF
- Proper gas flow



# TOO HIGH FLOW AT REDUCED TEMPERATURE



Too high flow at a reduced temperature may,

- Cause blood damage
- Impede the surgery by flooding the field
- Cause an excessive rise in venous pressure
- Cause rewarming of the heart when cardioplegia has been used.
- Cause underperfusion.



THANK YOU