#### MONITORING OF RAISED INTRA CRANIAL PRESSURE AND MANAGEMENT OF RAISED INTRA CRANIAL PRESSURE

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#### Normal ICP

a. Adults and older children <10-15mmHg

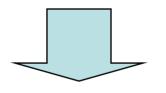
a. Young children 3-7

a. Term Infant

1.5-6

# Intra Cranial Cavity

- Volume is virtually constant
- Filled to capacity with fluids and solid material that are not compressible



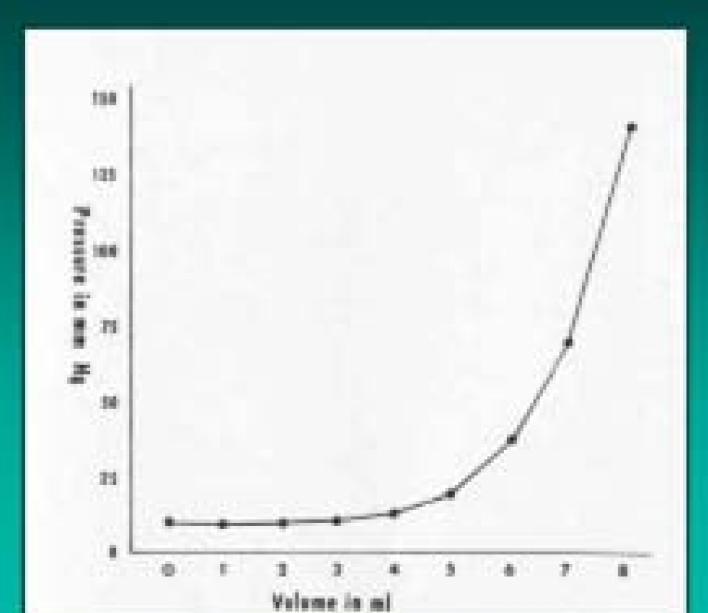
Increase in one constituent or an expanding mass with in the intra cranial space results in Raised ICP (Monro Kellie Doctrine)

# Intra Cranial Cavity Contents

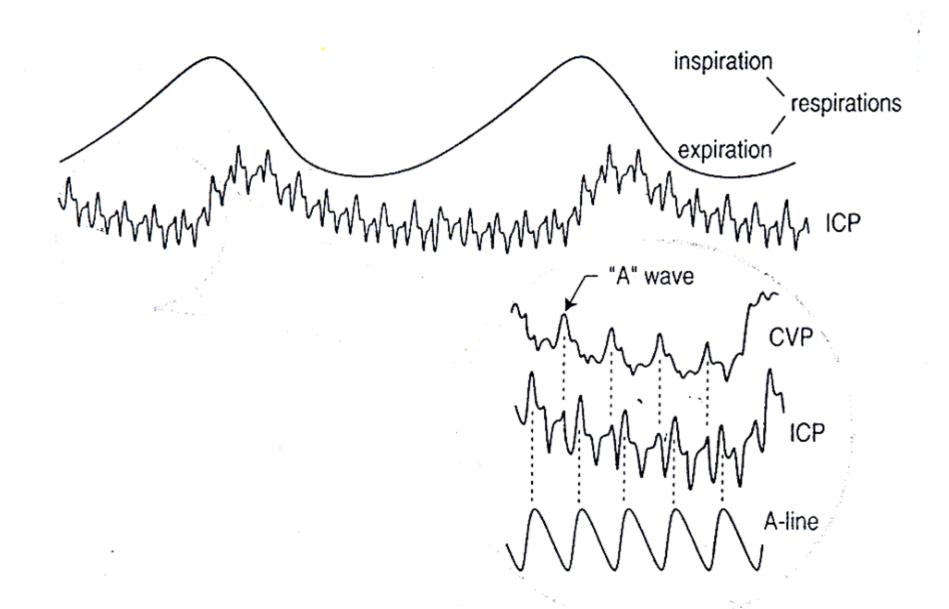
- Brain (1300-1750ml)
  - Neurons 500-700ml
  - Glia 700-900ml
  - Extra cellular fluid 100-150ml
- Blood 100-150ml
- CSF 100-150ml

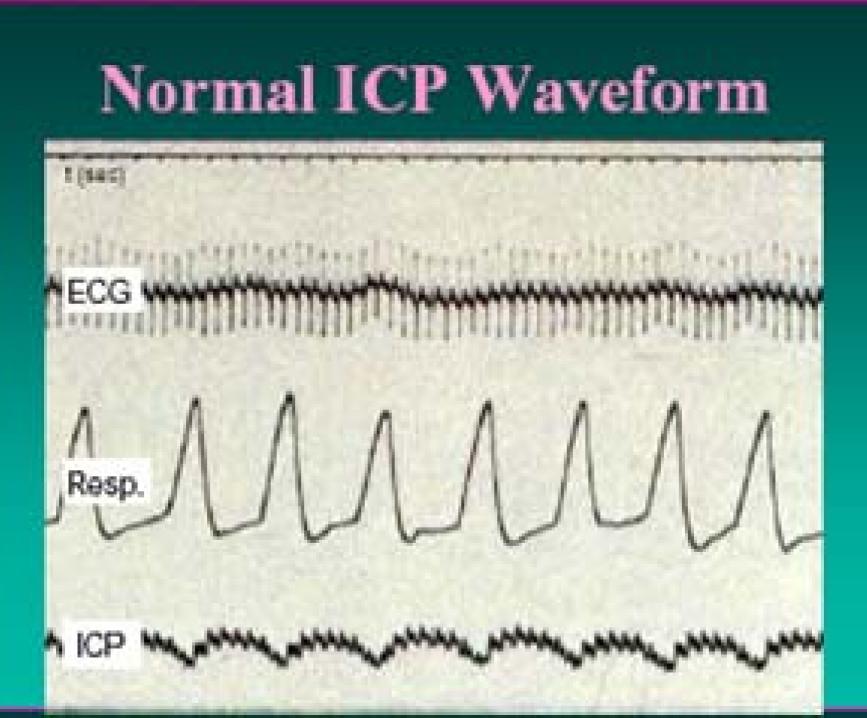
Constituents are non compressible but partially displaceable

# **ICP/Volume Curve**



#### ICP Wave





## Clinical Symptoms and Signs of Raised ICP

- Headache
- Nausea and vomiting
- Impairment of consciousness
- Papilloedema
- 6<sup>th</sup> C N Palsy False localizing sign
- Herniation signs (late)

## Causes of Increased ICP

- Increased volume of normal intra cranial constituents
  - Brain Cerebral oedema
  - CSF Hydrocephalous
  - Blood volume Vasodilatation secondary to increased Co<sub>2</sub>
- Space occupying lesion Tumour, Hematoma, Abscess,
- Idiopathic Pseudotumour Cerebri

Cerebral Perfusion Pressure and Auto Regulation

#### CPP = MAP-ICP

Cerebral auto regulation is a mechanism were by over a wide large changes in systemic BP produces only small changes in CBF

# ICP

- Cerebral oedema
- Hyperemia
- Intra cranial Mass
- HCP
- Hypoventilation
- Systemic Hypertension
- Venous sinus thrombosis
- Increased Muscle tone and valsalva maneuver
- Sustained Post traumatic seizure

Secondary Increase in ICP (3-10 Days Following trauma)

- Delayed hematoma formation
- Cerebral vasospasm
- ARDS with Hypoventilation
- Delayed Oedema formation
- Hyponatremia

## **Cushing Triad**

• Hypertension

• Bradycardia

Respiratory Irregularity

# **ICP** Monitoring

- Indication
  - Severe Head injury (GCS<8 after CPR) and either
    - Abnormal admitting CT

Or

- Normal CT with <u>></u> 2 of risk factors (Age >40yr, SBP <90mm Hg, Decerebrate or decorticate posturing</li>
- Multiple System Injured with altered level of consciousness
- Traumatic Intra cranial mass
  - A physician may choose to monitor ICP in some the patients
  - Post op subsequent to removal

#### **Contra indications**

Awake Patient

Coagulopathy

#### Duration of monitoring

 Till ICP is normal for 48-72 Hours after withdrawal of therapy

# Types of Monitor

• Intra ventricular catheter (Standard) –

#### – Advantages –

- Low cost
- Therapeutic CSF Drainage
- May be recalibrated to minimize measurement drift

#### – Disadvantages –

- Difficult to insert
- Obstruction of fluid column
- Some effort is required to check any monitoring function
- Transducer must be consistently maintain at a fixed reference point

- Intra parenchymal monitor Similar to IVC usually not subjected to measurement drift
- Less Accurate-
  - Subarachnoid screw- At high ICP brain may block Lumen
  - Subdural
  - Epidural
  - Infants-
    - Aplanation Principle
    - Fontometry

#### Intra Ventricular Catheter

• Right side frontal horn

• EAC is external land mark for Zeroing



# Type of Wave Form

- Normal wave form
  - Small pulsation transmitted from the systolic BP to intra cranial cavity
    - Large 1-2mm Hg Peak corresponding to altered systolic Pressure wave with small dicrotic notch.
    - The Peak is followed by smaller and less distinct peak
    - Followed by peaks corresponding to central venous A wave from right atrium.
  - Blood pressure pulsation are superimposed on slower respiratory variation
  - During expiration SVC Pressure Increases so ICP increase

## Pathological Wave Form

- As ICP rises cerebral compliance decreased the venous component disappears and arterial pulsation become more pronounced
- a. Lundberg A- (Plateau wave)
- b. Lundberg B- amplitude 10-20mmHg, last
  30 second to 2 minutes. Respiratory
  variation present.
- c. Lundberg C- frequency 4-8 per Minutes. Also called as Traube Hering Waves

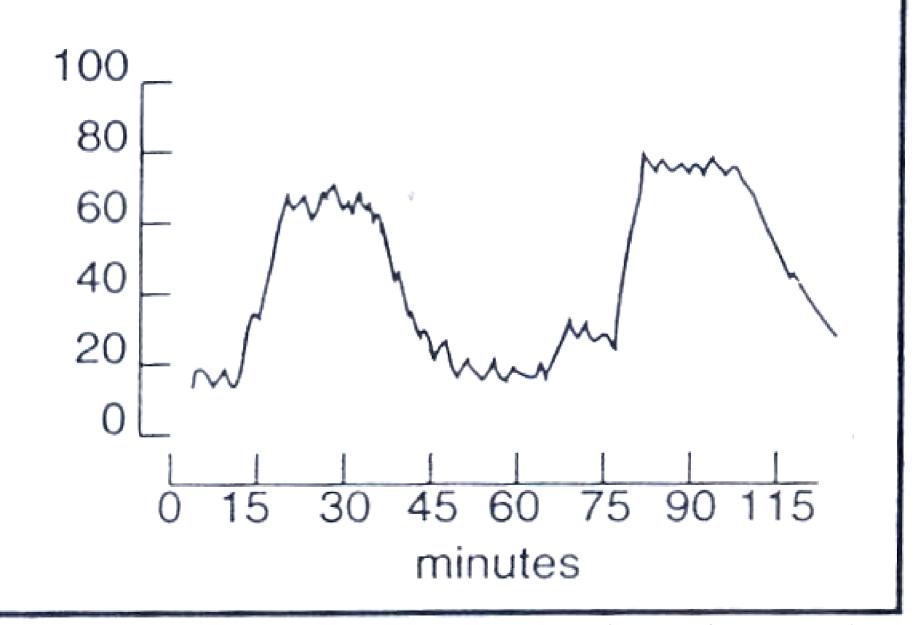


Figure 24-4 Plateau waves (Lundberg A waves)

# Normal Function of IVC System

- Wave form
- Pulsation
- Output
- Lowering of Head of Bed increases ICP.
   Bilateral JV pressure increases ICP

## **IVC Problems**

- Block
- Improper connection
- Change of position of head of bed
- Drip chamber fall

### **IVC Trouble Shooting**

- IVC no longer works
- ICP wave form dampened

# Adjuncts to ICP Monitoring

- Evoked potential
- SJO2
- TCD
- PET & SPECT

#### **ICP** Treatment Measures

- Guidelines
  - ICP treatment should be initiated for ICP >20-25mm Hg
- Options
  - Interpretation and treatment of ICP based on any threshold should be corroborated by frequent clinical examination and CPP data
  - CPP should be maintain ≥70mm Hg

## **ICP Management Protocol**

Goals of therapy

 Keep ICP <20mmHg (Prevents plateau waves from compromising CBF and causing cerebral ischemia and or brain death

– CPP >70mmHg (avoid hypotension)

#### Summary of measures to control IC-HTN\* Goals: keep ICP < 20 mm Hg, and CPP $\ge$ 70 mm Hg

| Step  | Rationale   |
|---|---|
| GENERAL MEASURES  | (should be utilized routinely)  |
| elevate HOB to 30-45°   | reduces ICP by enhancing venous outflow, but also re-<br>duces mean carotid pressure $\rightarrow$ no net change in CBF                                   |
| keep neck straight, avoid tight trach tape  | constriction of jugular venous outflow increases ICP  |
| avoid hypotension (SBP < 90 mm Hg)  | <ul> <li>normalize intravascular volume</li> <li>use pressors if needed</li> </ul>  |
| control hypertension if present   | <ul> <li>nitroprusside if not tachycardic</li> <li>beta-blocker if tachycardic (labetalol, esmolol)</li> <li>avoid overtreatment → hypotension</li> </ul> |
| avoid hypoxia ( $pO_2 < 60 \text{ mm Hg}$ )<br>(maintain airway and adequate oxygenation) | hypoxia may cause further ischemic brain injury   |
| ventilate to normocarbia ( $pCO_2 = 35-40 \text{ mm Hg}$ )                                | avoid prophylactic hyperventilation (see page 659)  |
| light sedation: codeine 30-60 mg IM q 4 hrs PRN   | (same as heavy sedation, see below)   |
| unenhanced head CT scan for ICP problems†   | rule out surgical condition   |

- Surgical treatment
  - Any subdural or epidural hematoma larger than 1cm maximal thickness should be removed
  - Patient with contused brain with progressive deterioration should be removed.
  - De-compressive craniectomy should be considered for ICP not controlled medically.
- General care
  - Avoid Hypoxia PaO<sub>2</sub><60mmHg
  - Avoid Hypo tension SBP ≤90mmHg

## Specific Treatment

- Prophylaxis against stress ulcer
- Control fever
- Arterial line
- CVP line
- IV fluid
  - NS +20Meqkcl/let.
  - Avoid hypotonic solution
  - Normalize intra vascular volume
  - Concept of running patient dry is obsolete
  - If injury to other system present they may dictate Fluid management
  - Pressors or preferable to IV fluid bolus in head injury

#### Measures to Lower ICP

- General Measure
- Specific measures

Second Tier Therapy for Persistent IC Hyper Tension

- High dose Barbiturate therapy if ICP>20-25mmHg
- Hyper ventilation
- Hypothermia
- De-compressive craniectomy
- Lumber drainage
- Hypertensive therapy

#### **Adjunctive Measures**

Lidocaine – 1.5 mg/kg IVP

• High frequency (Jet) ventilation

#### High Dose Barbiturate Therapy

- Guidelines
  - May be considered in hemodynamically stable salvageable severe TBI patient with Intra cranial Hyper tension refractory to maximal Medical and surgical ICP lowering therapy
  - Theoretical benefits of barbiturates in head injury is derived from

1. vasoconstriction in normal areas (shunting blood to ischemic brain tissue)

2. Decreased metabolic demand for oxygen (CMRO<sub>2</sub>) with accompanying reduction of CBF.

- 3. Free radical scavenging
- 4. Reduced intra cellular calcium
- 5. Lysosomal stabilization

- Pentobarbitol is choice
  - Load 10mg/kg/hr for 4 hrs.
    - 1hr. 2.5mg/kg every 15 min. doses
  - four doses Next 3hrs. 10mg/kg /hr infusion Maintain 1.5mg/kg/hr infusion
  - BSAER early. Repeat if Pentobarbitol level >6mg%
  - Goal ICP <24mmHg level 3-5mg%
  - If ICP <20 mmHg continue treatment for 48hrs. And then Taper dose

# THANK YOU