# **Anesthesia** Sanda-Maria Copotoiu

Anestezia V MG

**Preanaesthetic consult I.** Objectives **III.** Types I. GA II. Loco-regional an **Epidural** Ι. **II.** Spinal **III.** Perriferal nerve blocks **IV.** Local - topical

#### V. Basic principles

# **Preanesthetic consult**

- History
- Physical exam
- Anterior lab+ supplemental
- Patient information certainty as to his ability to understant and take decisions
- Infromed consent
- Primary evidence- drug interactions, preop medication
- ASA risk assessment

#### **ASA risk assessment**

- ASA 1 = no organic, functional, biocehmical, psychiatric disease
- ASA 2 = Mild to moderate systemic disease, resoc or not for surgery eg:
  - cardiopathy moderately impinging on physical activity
  - Essential HBP,
  - Diabetis
  - 🌻 anemia,
  - Age extremes,
  - Morbid obesity,
  - Chronic bronchitis

#### **ASA risk assessment cont**

#### 🌻 ASA 3

#### Severe systemic disturbances, cause or not for surgery eg:

- Cardiac diseases limiting physical activity
- Essential HBP difficult to cope with,
- Diabetes + vascular complications,
- Chronic pulmonary disease
- 🌻 angina ,
- History of AMI

#### **ASA risk assessment cont**

ASA 4 Severe life threatening systemic diseases  $\pm$  surgery e.g:

- Congestive cardiac failure,
- Persistent angina,
- Pulmonary failure,
- Cardiac failure,

\*

Liver advanvced failure

#### **ASA risk assessment cont**

ASA 5 Moribund patient, limitted survival chances, surgery is the last remedy (resuscitation effort)

ASA 6 Brain dead, donor

**E** Any patient operated in emergency

#### **Anesthesia choice**

Coexisting diseases ± connection with surgery

- Site of surgery
- Position during surgery
- Elective character, or emergency
- One-day surgery
- Full stomach risk (occlusion, postingestion)
- Age

Patient' wish, verbaly or expressed in writing, fully consciousness

# Anesthetists' responsibilities

Physical status evaluation – functional assessment
Anesthesia risk assessment
Tailoring according to

comorbidities
Pts' wishes
Surgery needs submitted to the best interest of the patient

Choice of the adequate anesthetic technique

#### Cardiac Risk Factors in Patients Undergoing Elective Major Noncardiac Surgery

Surgery	
High-risk surgery	Abdominal aortic aneurysm
1 point	Peripheral vascular operation
	Thoracotomy
	Major abdominal operation
Ischemic heart disease	History of myocardial infarction
1point	History of a positive exercise test
	Current complaints of angina pectoris
	Use of nitrate therapy
	Q waves on electrocardiogram
Congestive heart failure	History of congestive heart failure
1 point	History of pulmonary edema
	History of paroxysmal nocturnal dyspnea
	Physical examination showing rales or S₃ gallop
	Chest radiograph showing pulmonary vascular redistribution
Cerebrovascular disease 1point	History of stroke
	History of transient ischemic attack
Insulin-dependent diabetes mellitus 1point	
Preoperative serum creatinine	

concentration > 2 mg/dL 1 point

- 1. Points 0: Class I Very Low (0.4% complications)
- 2. Points 1: Class II Low (0.9% complications)
- 3. Points 2: Class III Moderate (6.6% complications)
- 4. Points 3: Class IV High (>11% complications)

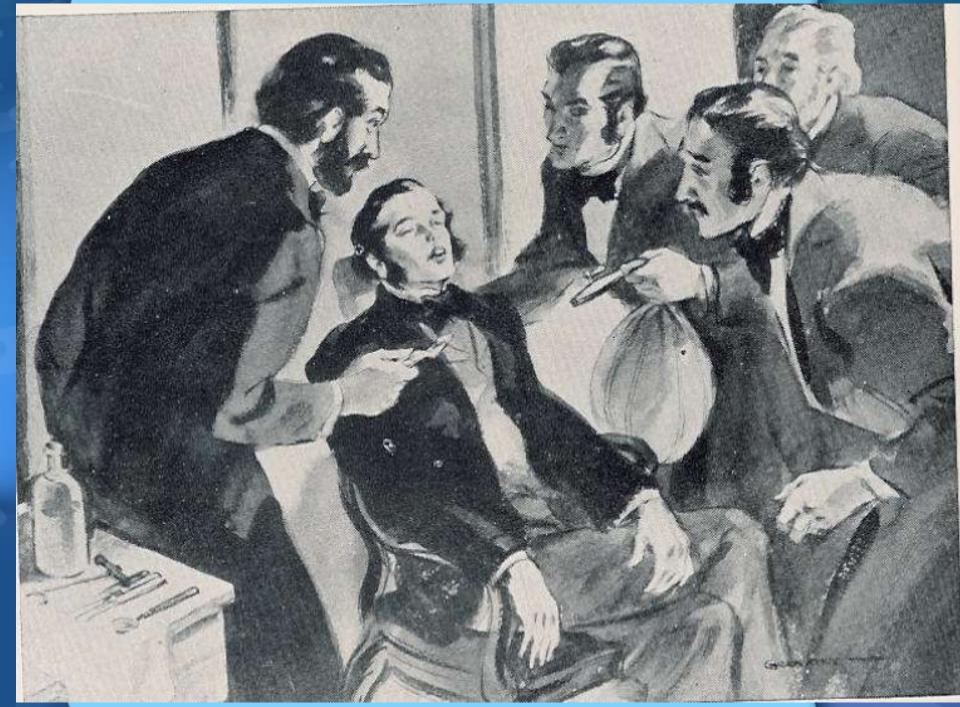
#### **Types of anesthesia**

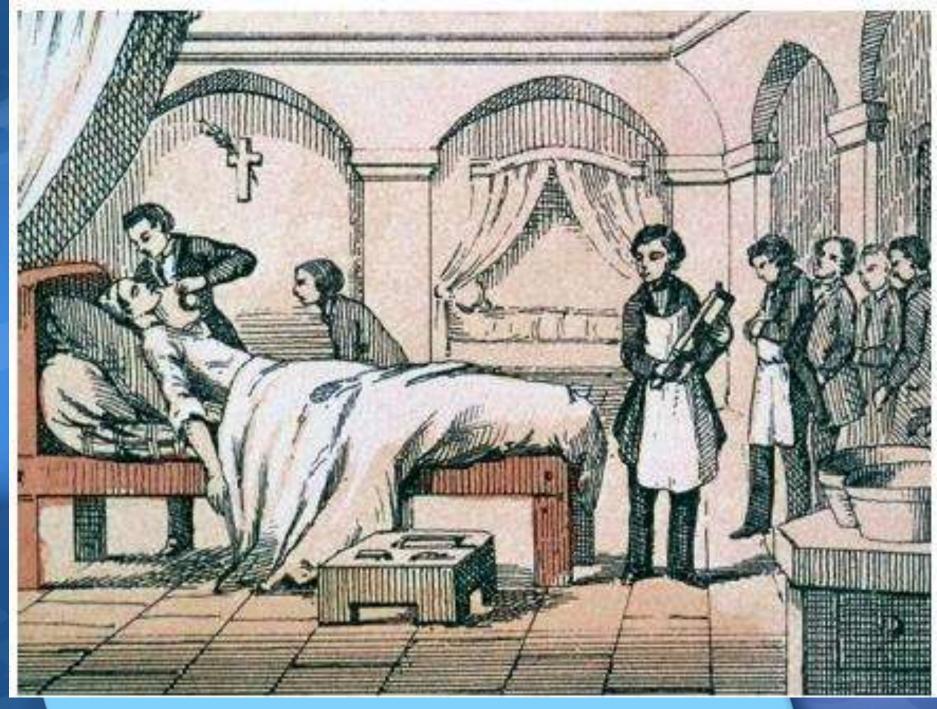
- **GA**
- Regional
- Perifferal nerve blocks
- Local an



#### •Shen Nung 2<sup>nd</sup> Chinese emperor cannabis

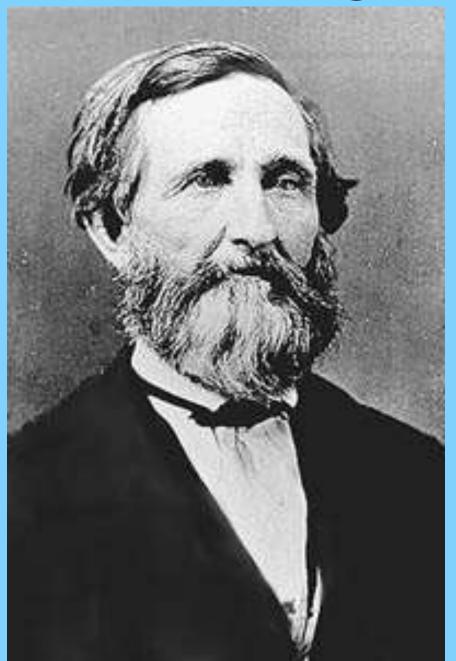








# **Crawford Long**



#### **Horace Wells**





#### **Morton Dagherotip**



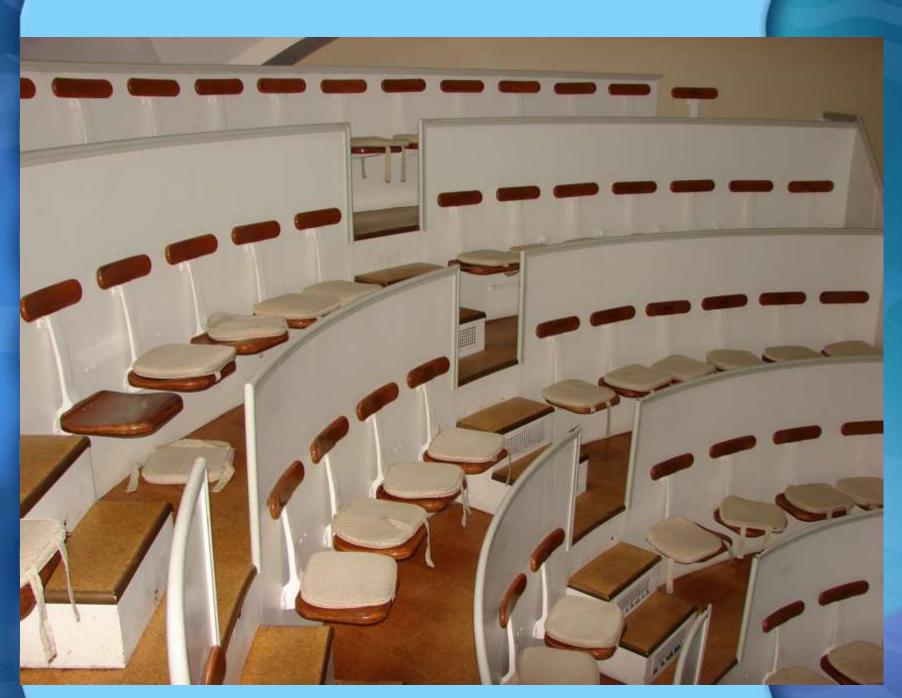












# Diabetes

 $\diamond$ 

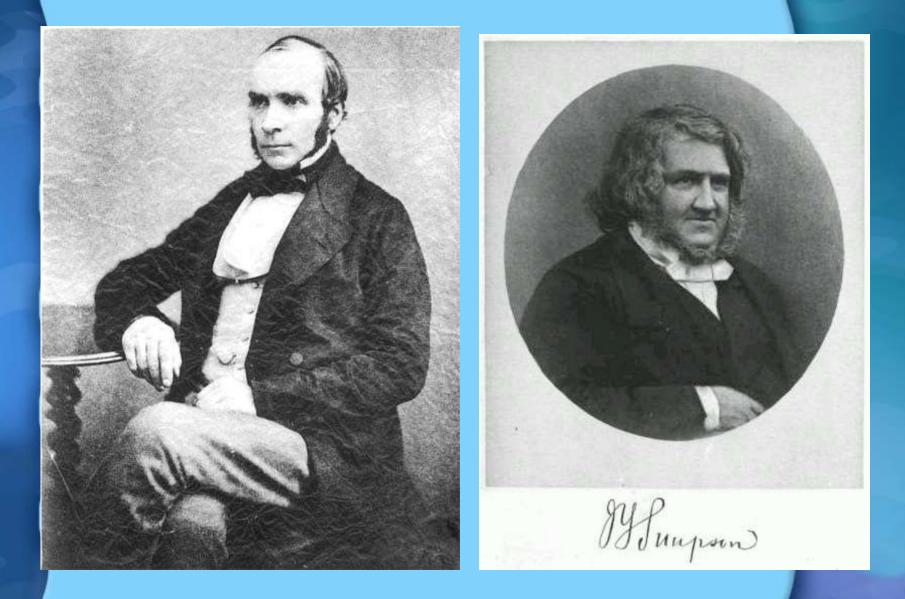
# Ether Dome Neuroendocrinology

### **Obstetric anesthesia**

 James Young Simpson – obstetrician, Edimburgh, 4.XI.1847, autoanesthesia
 Answers to the Religious Objections Advanced against the Employment of Anaesthetic Agents in MIdwifery and Surgery and Obstetrics

John Snow

Dr Snow gave that blessed chlorophorm and the effcect was soothing, quieting, and delightful beyond measure. Queen Victoria



#### **Queen Victoria**



# **GA objectives**

Analgezia

#### Hypnosis

Vital and neurovegetative stabilty, block of the vegetative response to pain

#### Muscle relaxation

# **GA** steps

- Induction
- maintenance
- Awakening

 Postoperative analgesia– Recovery room, PACU

# Induction Monitoring!!!!

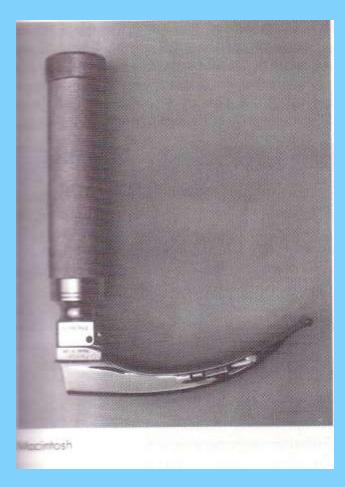
#### • Hypnotics

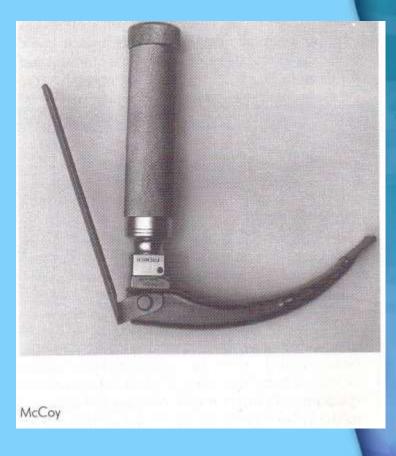
- Barbiturates thiopental, metohexital
- Benzodiazepine
- Propofol
- Etomidate

**Preoxigenation !!!** 

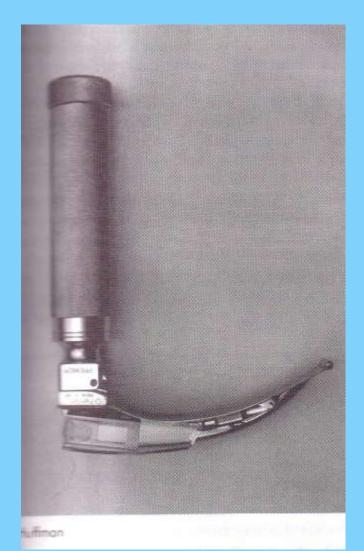
- Rapid sequence induction crush induction
- Relaxation priming, timing

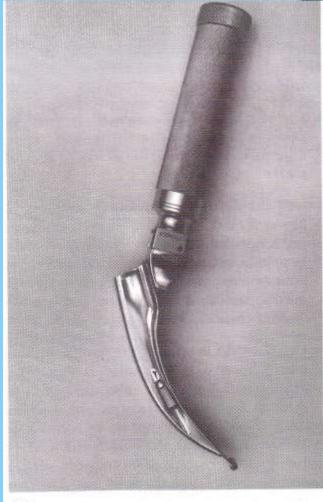
#### Laringoscopes





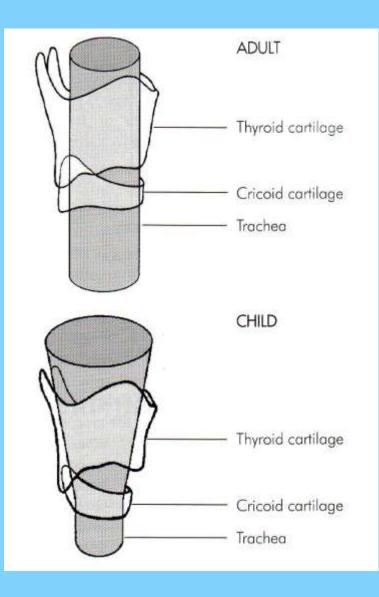
# Laringoscoape



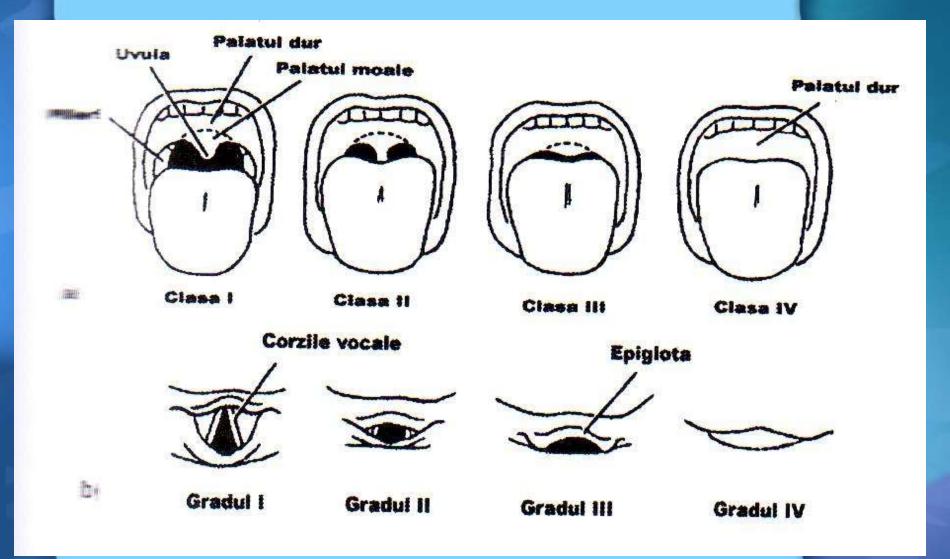


Polio

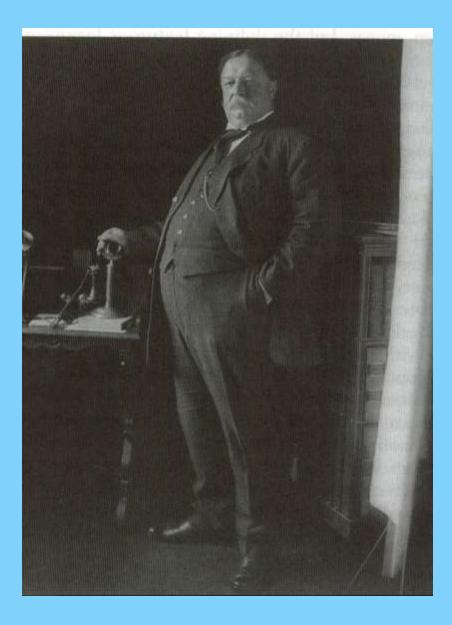
#### The adult and child larynx



### Intubation difficlutyMallampati; Cormack și Lehane



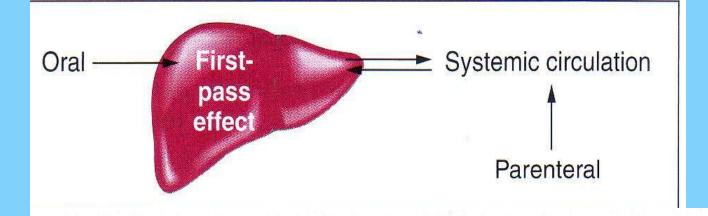
# Taft





# **Tissue compartments**

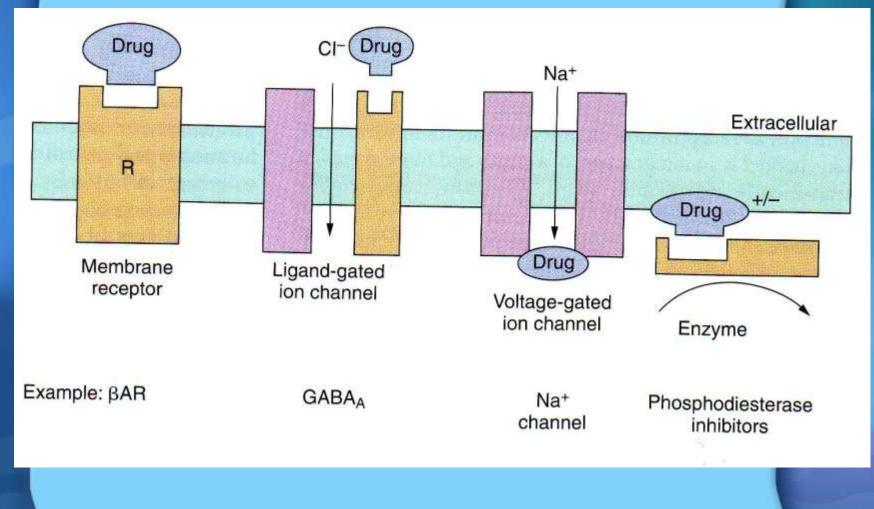
Table 8-4 Body Tissue Compartments				
Compartment	Body Mass (% of a 70-kg Adult)	Blood Flow (% of Cardiac Output)		
Vessel-rich group	10	75		
Muscle group	50	19		
Fat group	20	5		
Vessel-poor group	20	1		



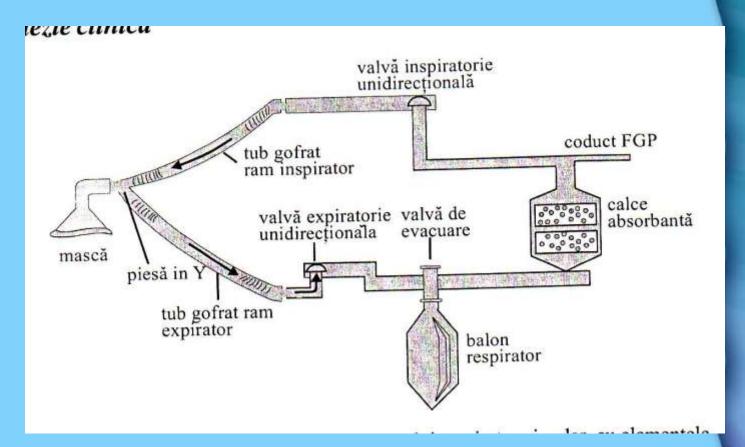
A PERSON AND A PER		A REAL PROPERTY OF THE PROPERT		CENAR OF EVENING STUDY LODGE SHEET		
	and the state of the provide state of the	and the second state of th	CONTRACTOR OF CONTRACTOR OF CONTRACTOR	The second se	CONTRACTOR DESCRIPTION	Contraction of the second seco
					AND DO THE LAST	
	1 1000 and 10 19 19		Y ADDER 199 11 191 191 191			
	STREET AND ADDRESS AND		A T ANDRESS AND ADDRESS ADDRES	9 1 UNG200 UN 9 2		tments
A State of the second	Description of the statement	Construction of the second states and the second s	Dr. ANNING MADE TO AN A STORE	and the second sec	Configurate and Description of the	
	SALE DE DESERVER DE		AT HOUSE BEAT MICH. N. R. M.		And the second s	

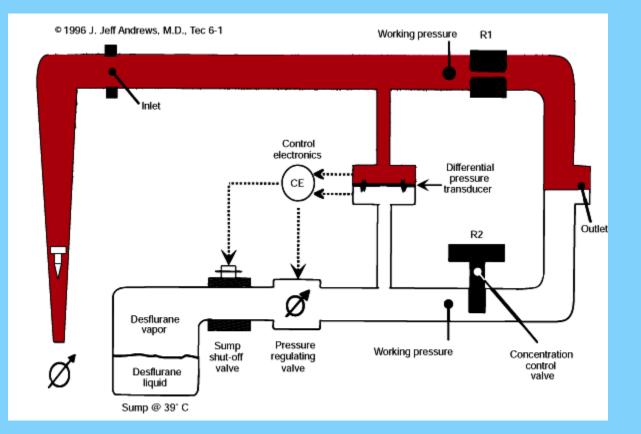
Compartment	Body Mass (% of a 70-kg Adult)	Blood Flow (% of Cardiac Output)
Vessel-rich group	10	75
Muscle group	50	19
Fat group	20	5
Vessel-poor group	20	1

# **Receptors and anesthetics**

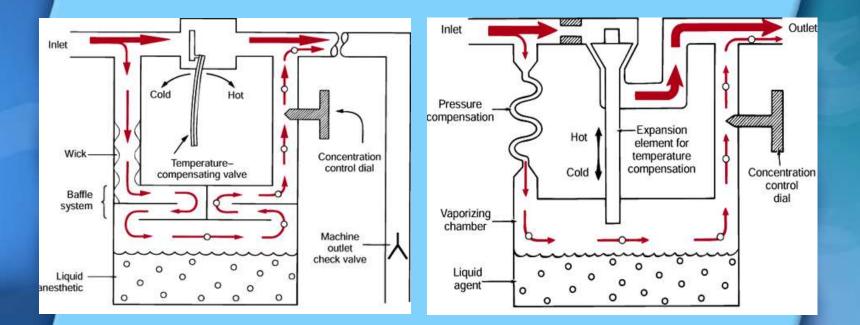


### **Circle respiratory system(Sword)**

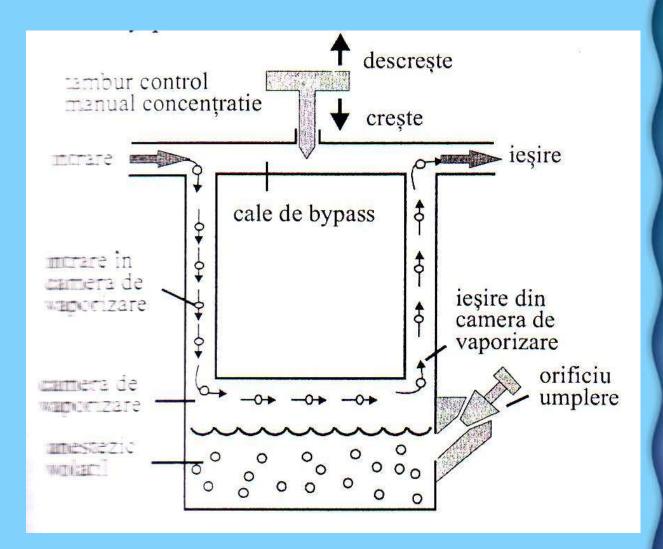


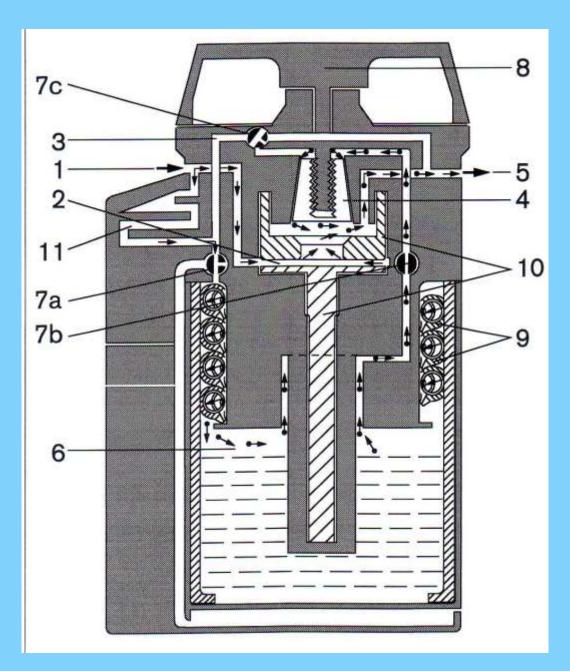






#### Vaporizors Plenum with variable by-pass

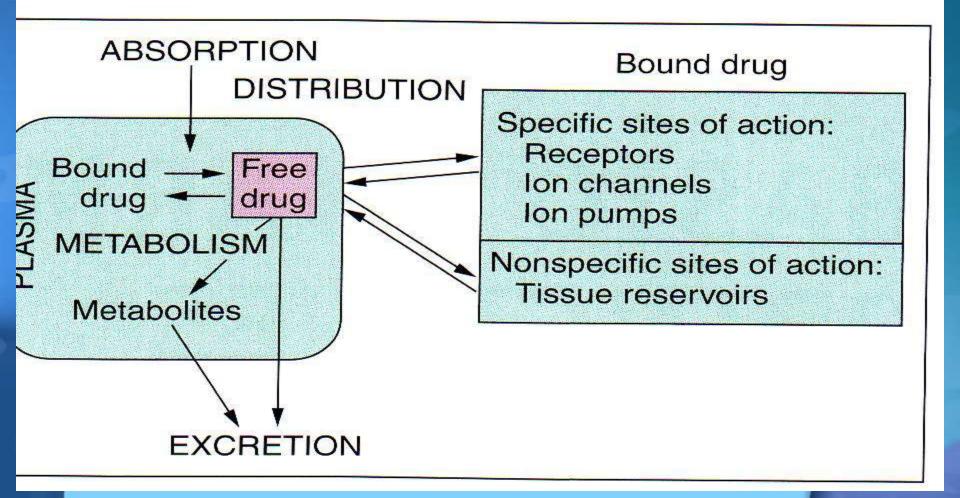




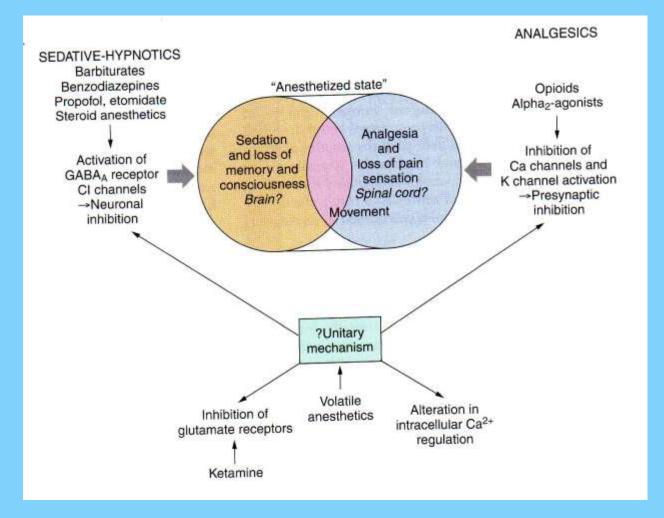
# **Anesthetic breathing systems**

Туре	Reser voir bag	Air source	rebreathing	Name, cath egory
open	No	Environmental air	No	Bag and bottle
Semi-open	No	Air+O2 from external source	partial	Bag and bottle +occlusive packing
Semi-closed wtihout absorbtion	yes	Air+O2 from external source	No	Bain, Modified Jackson-Rees, Ayre's T piece, Lack, Magill
Semi-closed with absorbtion	yes	Air+O2 from external source	partial	Co2 absorbers with leak
Closed	yes	Air+O2 from external source	complete	Co2 absorbers, no leak

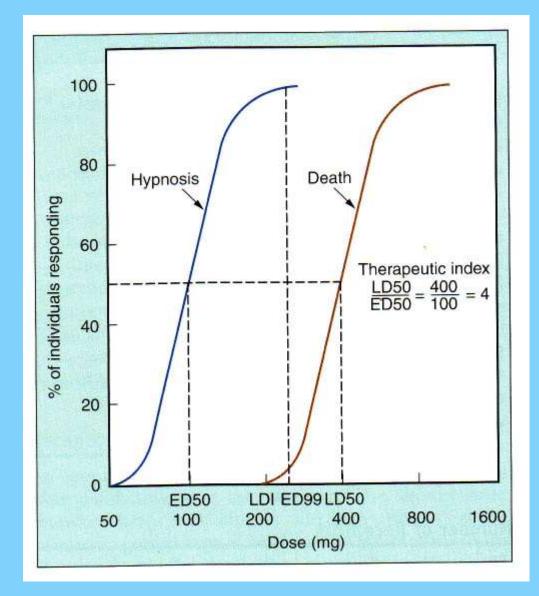
# Absorbtion, metabolism, excretion



Major routes of action of sedatives, hypnotics and analgesics specific for volatile anesthetics



#### Volatile, BNM



#### Altitude effect on vaporizers ↑ pressures

- Vaporizoare calibrate în concentrație

- Das density alterations→↑ resitance to flux in the vaporization chamber
- ↓vaporizer out put and concentration in vol%↓

Eg: p= 2atm  $\rightarrow$  conc vol%  $\downarrow$  to  $\frac{1}{2}$ 

- Vaporizers with flux measurement
  - ↓conc vol% or partial pressure

## **Basic principles in anesthesia**

Farmacokinetics:
Absorbtion
Distribution
Metabolization
Inhaled or PEV administered drugs
P<sub>A</sub> ↔ P<sub>a</sub> ↔ P<sub>cr</sub>

# **Basic principles in anesthesia**cont

### **Pharmacodynamics**

Receptors' response to drugs

#### Effects of mechanisms Eg: drug power

- Agonism
- Antagonism competitive or noncompetitive
- Aditive effect
- Sinergism

## **P**<sub>A</sub> Alveolar pressure

# The amount of alveolar anesthetic depends on:

- Pi partial pressure in the inspired air
- Alveolar ventilation
- Ventilatory characteristics of the anesthesia machine

### The amount of anesthetic in the arterial blood depends on:

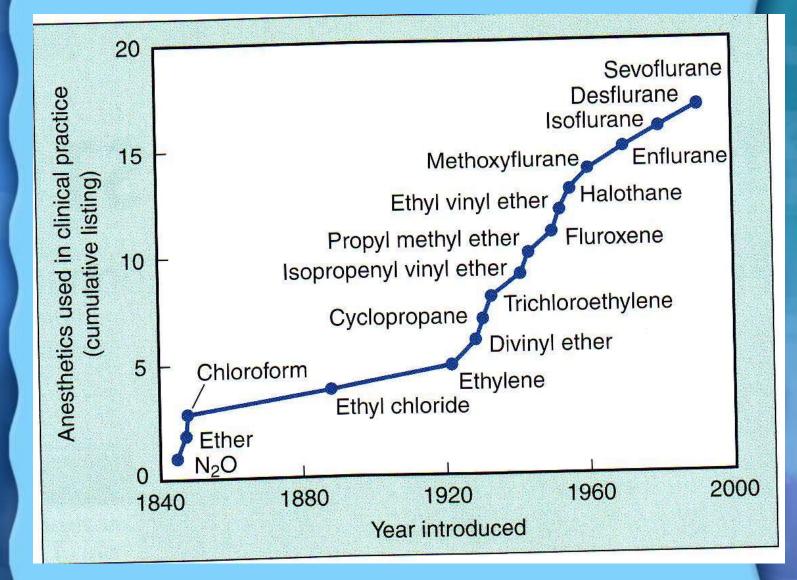
- Anesthetic solubility
- Cardiac output
- Alveolo-venous gradient

## Pi anaesthetic gas effects

### Concetration effect- High Pi at induction

2nd gas effect

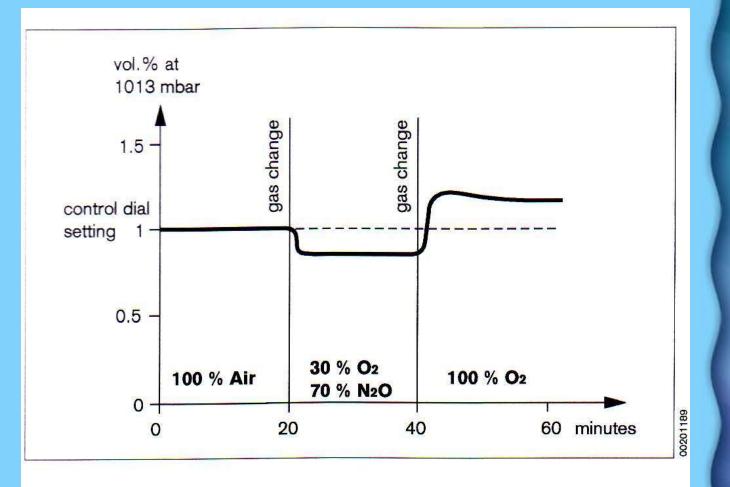
# **Volatile anesthetiss**



# P<sub>A</sub> & P<sub>br determinants</sub>

- P<sub>A</sub> = Alveolar offer overtakes from alveoli to venous pulmonary blood
- Offer:
  - $\mathsf{P}_{\mathrm{I}}$
  - $-V_{\rm A}$
  - Respiratory system characteistics
- Overtake:
  - Solubility
  - Cardiac output
  - $D_{A-v}$

# Carrier gas composition and the vaporizer output



# MAC

Factors increasing MAC	Factors decreasing MAC
Drugs: ephedrine, amphetamines, cocaine, ethanol	Benzodiazepines, iv anesthetics, opioids, lithium
Age	Age - increasingly
Hyper, hypothermia	Anemia, hypercarbia, hypothermia

# MAC / MAC awake

#### Table 1. Structured Interview

- 1. What is the last thing you remember before going to sleep?
- 2. What is the first thing you remember waking up?
- 3. Do you remember anything between going to sleep and waking up?
- 4. Did you dream during your procedure?
- 5. What was the worst thing about your operation?

#### Table 2. Awareness Categorization

- No awareness: no reported awareness or a vague description, or what had been reported had a high probability of occurring in the immediate pre- or postoperative period; i.e., music, people talking, dressing application
- 2. Dreaming, possibly associated with awareness
- 3. Possible awareness: patient unable to recall any event definitely indicative of awareness
- Awareness: recalled event was confirmed by attending personnel, or the investigators were convinced that the memory was real, but no confirmation could be obtained

# MAC / MAC awake

#### Table 1. Structured Interview

- 1. What is the last thing you remember before going to sleep?
- 2. What is the first thing you remember waking up?
- 3. Do you remember anything between going to sleep and



procedure? out your operation?

#### Table 2. Awareness Categorization

- 1. No awareness: no reported awareness or a vague description, or what had been reported had a high probability of occurring in the immediate pre- or postoperative period; i.e., music, people talking, dressing application
- 2. Dreaming, possibly associated with awareness
- 3. Possible awareness: patient unable to recall any event definitely indicative of awareness
- 4. Awareness: recalled event was confirmed by attending personnel, or the investigators were convinced that the memory was real, but no confirmation could be obtained

# **Table 5–1** Characteristics of Nonionized and Ionized Drug Molecules

Characteristic	Nonionized	Ionized
Pharmacologic effect	Active	Inactive
Solubility	Lipids	Water
Cross lipid barriers (renal tubules, gastrointestinal tract, placenta, blood-brain barrier)	Yes	No
Renal excretion	No	Yes
Hepatic metabolism	Yes	No

### **Spinal anesthesia**

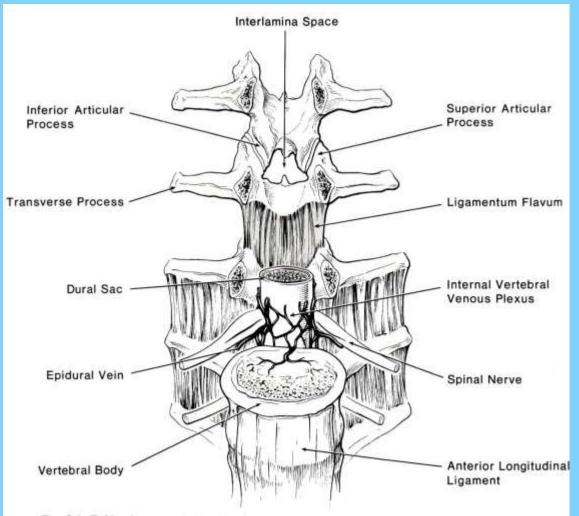
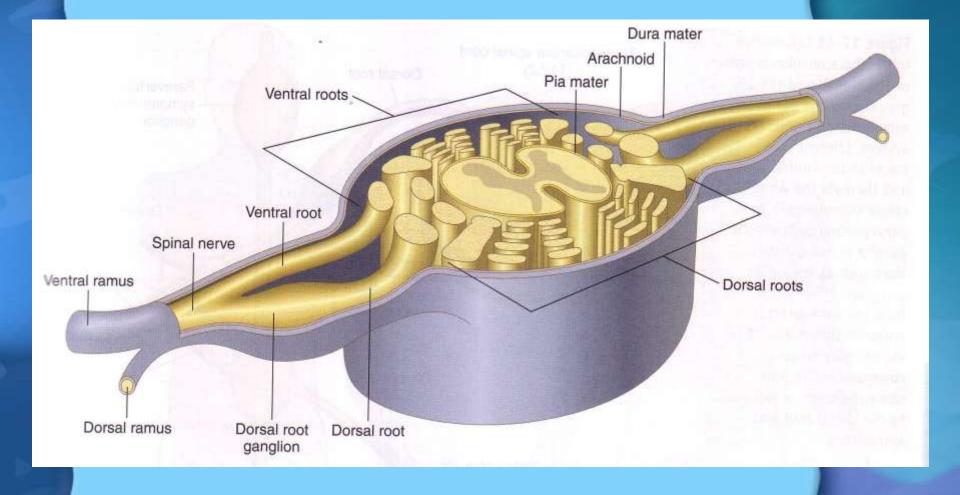
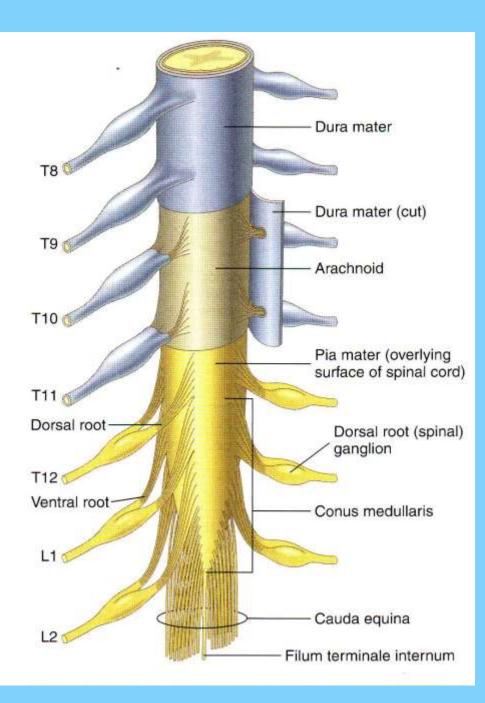


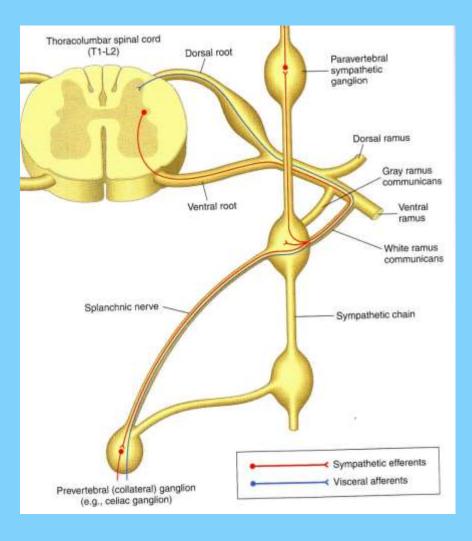
Fig. 8-6. Epidural space, relationships from anterior view. Note "interlaminar space" at one level and covered by ligamentum flavum at the level below. Epidural veins are in continuity with veins draining vertebral body ("internal vertebral venous plexus"). (Macintosh, R.R.: Lumbar Puncture and Spinal Analgesia. Edinburgh, E. & S. Livingstone, 1957)

#### The spine

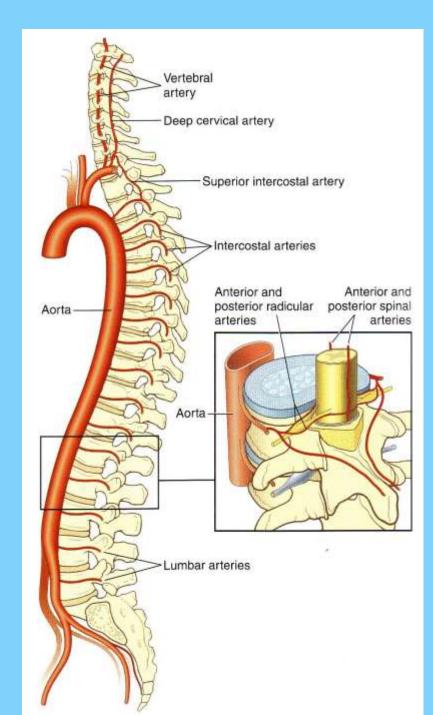




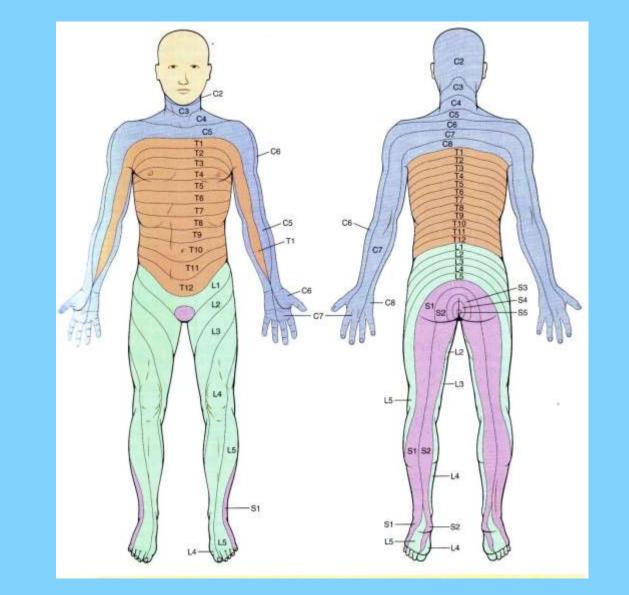
#### The spine



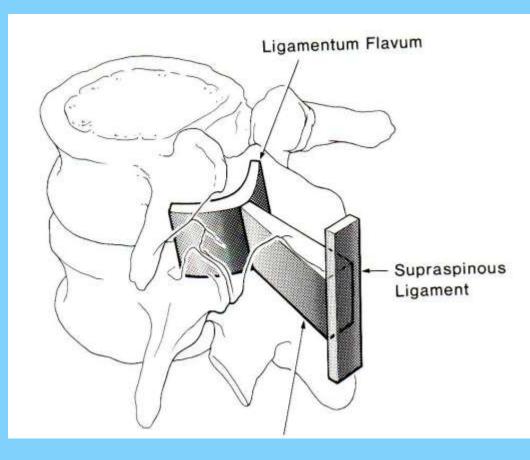
#### Spine vessels



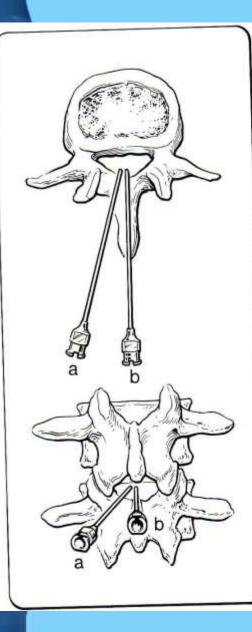
#### Dermatoams Spinal sensorial innervation areas



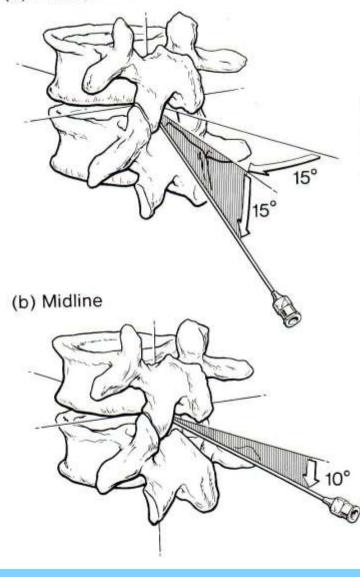
### Anatomy of the vertebral ligg

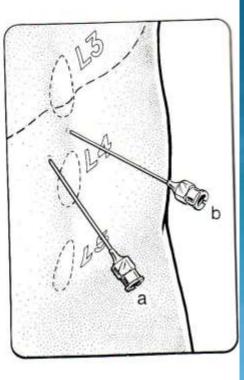


#### Median and paramedian (lateral) approach

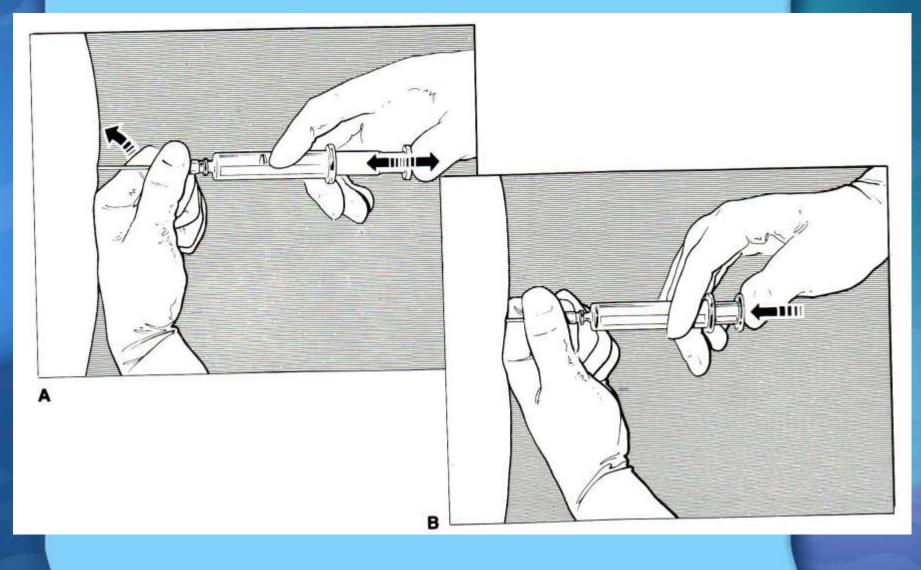


(a) Paraspinous

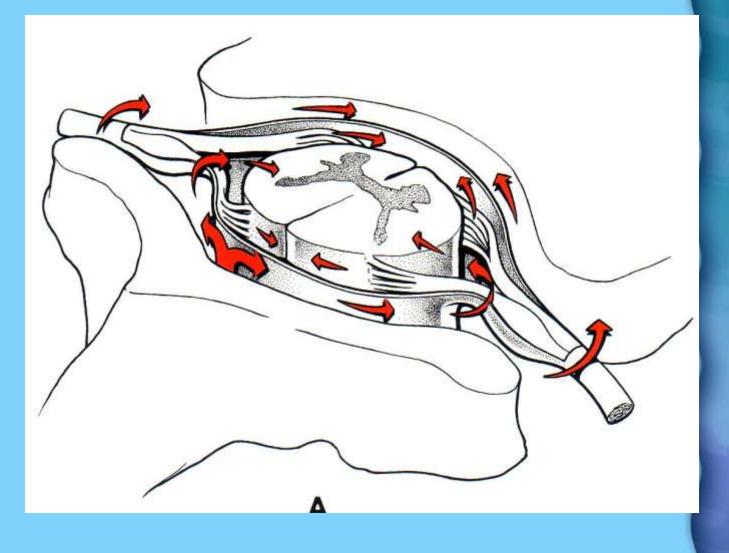




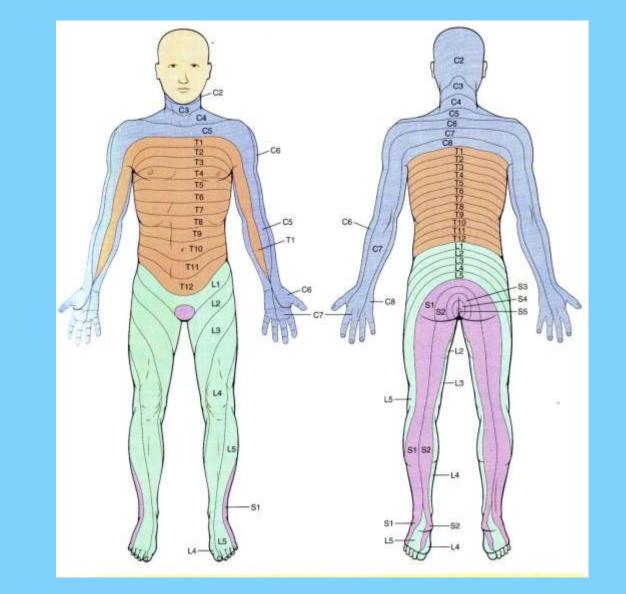
#### **Epidural anesthesia–Bromage**



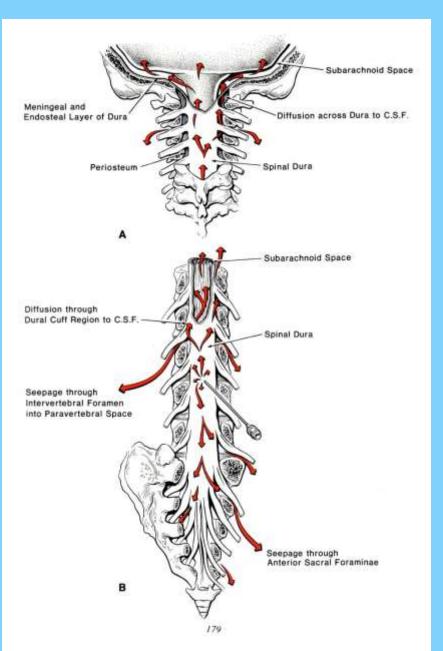
# Horizontal diffusion of local anesthetic during epidural injection



#### Dermatomes Sensorial innervation areas

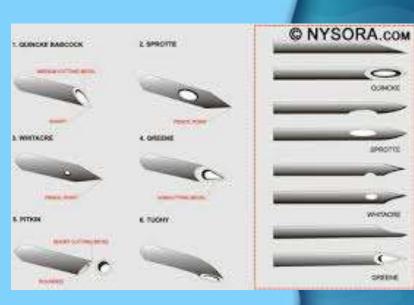


#### **Vertical epidural diffusion of LA**

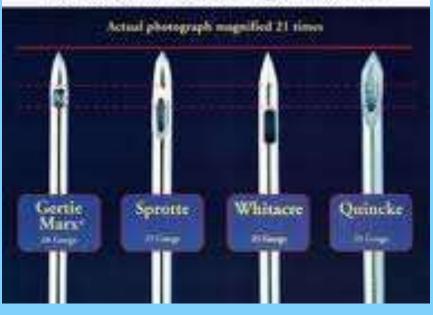


#### Spinal needles' tips





The Gertie Marx<sup>\*</sup> Needle for Regional Anesthesia



### LA characteristics

Strong and effcient in small concentrations

- Good penetrability
- Rapid action installation
- Long duration of action
- Reduced systemic toxicity
- Nonirritant, non neurogenic injuries
- Reversible
- Easy to sterilize

#### Karl Koller, 1894, cocaine



# LA

Weak bases **Esthers** : procaine, cocaine, clorprocaine, tetracaine T1/2 pl minuts **Plasma Colinestheraze Metabolized to PAB Amides:** lidocaine, mepivacaine, bupivacaine, etidocaine, ropivacaine N-dazalkilation, liver hydrolisis T1/2: several hrs

#### LA action emchanism

 Block the nervous conduction by lowering the AP that cannot reach the threshhold value

• Direct interaction with Na<sup>+channel receptors</sup>: passive diffsuion in neutral state (more rapid), ionized - linkage to Na<sup>+channels</sup>

#### Installation speed, duration & intensityAL

- Liposolubility: increases power (easily traverse of nervous membranes)
- Linkage to plasmatic proteins prolongs effects
- Small PKa = more rapid effect (electroneutralitaty favors diffusion)
- Acidity delays installation by reducing the nr of nonionized molecules

#### **Power and toxicity**

LA	Anesth etic power	CNS toxicity	Maximal doses mg	Maximal dosis with epinefring mg	
Procaine	1	1	400	600	
Lidocaine	2	3	300	500	
Mepivacaine		2	300	500	
Bupivacaine	14	12	175	200	

### LA' effects on nervous fibers

Ту	pe	Myelinis ation	Diameter µ	Function	
A	-α	++	6-22	Motor efferents, proprioceptor a	fferents
A	-β	++	6-22	Motor efferents, proprioceptor a	fferents
A	-γ	++	3-6	Muscular spin effecrents	
A	-δ	++	1-4	Pain, touch, temperature affere	nts
	B	+	< <b>3</b>	Preganglia autonomous	
	С	-	0,3-1,3	Motor efferents, proprioceptor a autonomous postganglia	fferents

# **Clinical sequence of conduction anesthesia**

- 1. Sympathetic blockade: sympathetic vasodilation, 1 skin temperature
- 2. Los of pain and thermal sensation
- 3. Loss of proprioception
- 4. Tactile and pressure sensations lost
- 5. Motor paralysis

## **Complications - severity**

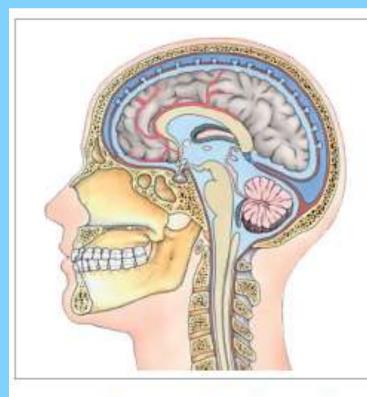
AccidentsIncidentsComplications

PreIntraPostoperative



High Position on Wall Street

#### What is the normal plasma osmolality??



Brain 1,340 mL + 40 mL (3%) Blood 120 mL - 40 mL (-33 %) CSF 140 mL - 40 mL (-29 %)

Total 1,600 mL

Intracranial compartment responses to a change in plasma osmolality: A decrease in plasma osmolality by approximately 3%, say, from 288 to 280 mosmol/kg H<sub>2</sub>O, invariably results in an increase in brain volume by 3%, causing a decrease in blood and/or CSF volume by as much as 30%. Osmolarity vs. osmolality

	Pla Electrolytes (mmol/l)	sma Osmotically active species (mosmol/l)	Ringer's acetate (mmol/l)	0.9 % NaCl (mmol/l)
Na+	142	142	130	154
K.	4,5	4,5	5	
Ca <sup>2+</sup>	2,5	1,3*	1	
Mg <sup>2+</sup>	1,25	0,7*	1	
CL-	103	103	112	154
HCO3-	24	24		
Phosphate <sup>2-</sup>	1	1		
Sulfate <sup>2-</sup>	0,5	0,5		
Organic acids	1,5	1,5	27	
Proteinate-	20	1		
Glucose		5		
Urea		5		
Σ	Σ = 291		$\Sigma = 276$	$\Sigma = 308$
Theoretical osmolarity (mosmol/l)	2	91	276	308
Water content (%)	94		99,7	99,7
Theoretical osmolality (mosmol/kg H <sub>2</sub> O)	310		276	308
Osmotic coefficient	0,926		0,926	0,926
Actual osmolality (mosmol/kg H <sub>2</sub> O)	287		256	286
Measured osmolality** (mosmol/kg H <sub>2</sub> 0)	2	88	256	286

\* Because of protein binding

\*\* Freezing point depression