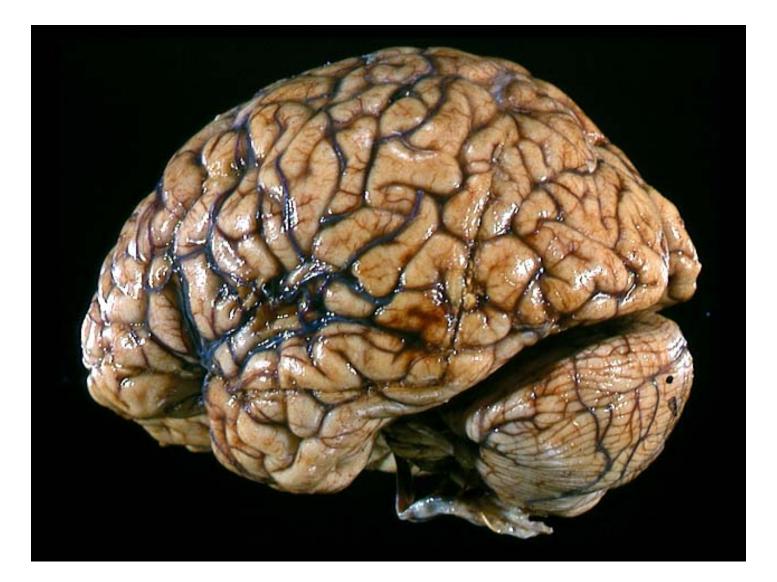
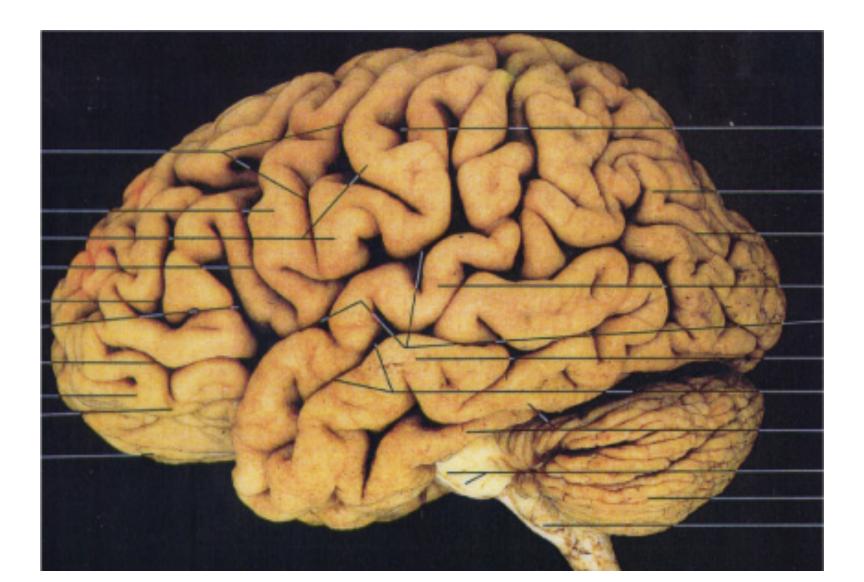




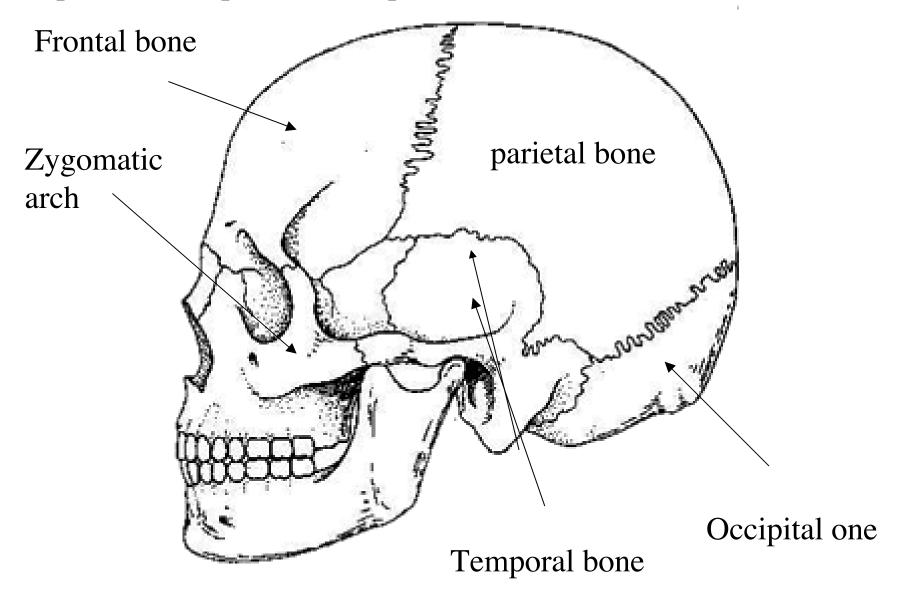
A fresh brain with dura removed. Note the numerous superficial blood vessels running in the arachnoid.



If we strip off the meninges we see the brain has a heavily folded surface or **cortex** (in latin: cortex =bark)



Recall that the main bones of the cranium are the frontal, parietal, occipital and temporal.



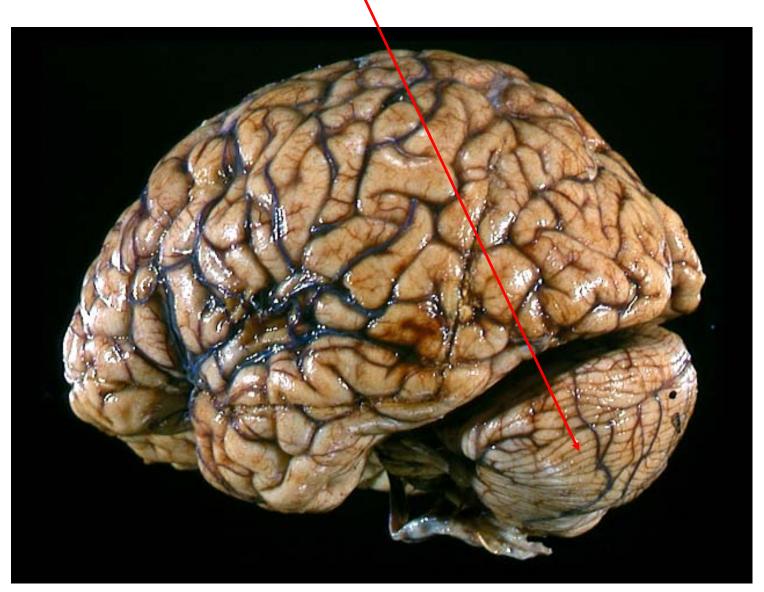
Anatomy and Functional Areas of the Brain

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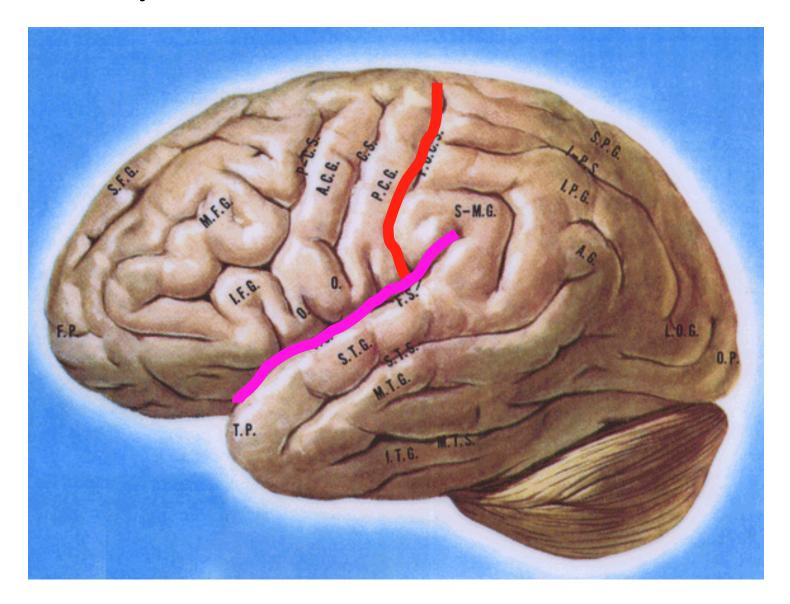




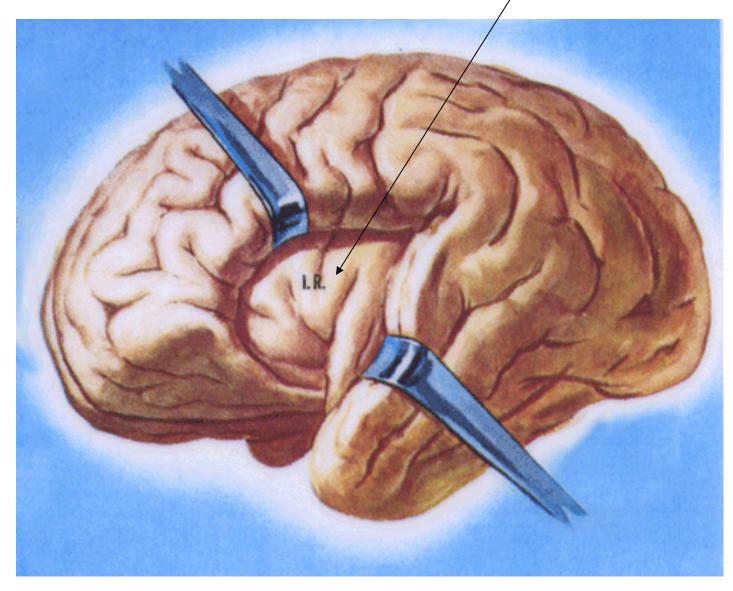
Note appearance of cerebellum; the gyri are thinner and straighter than in cortex



Most of the gyri and sulci have individual names but for now we only need to remember the central and lateral sulci.

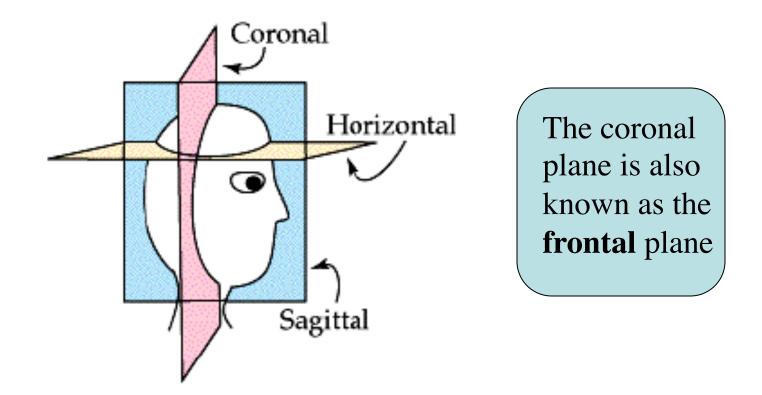


Inside the lateral fissure there is a hidden area of cortex, the **insula** or 'Island of Reil'.

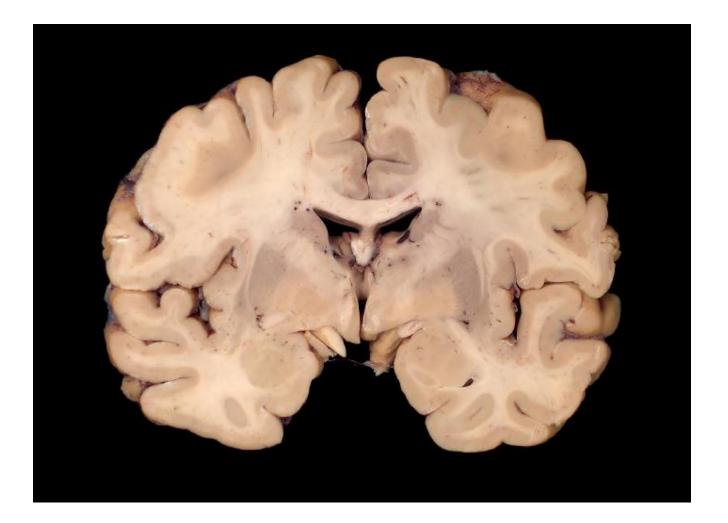




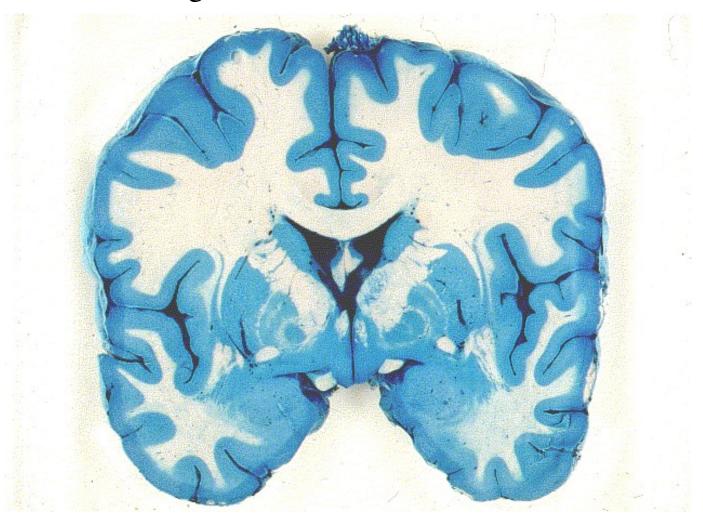
Before we can investigate the interior of the brain by cutting it up we need to **define the planes of the brain** that we will cut (section) in.



Ggoing to a higher horizontal (axial) plane is going **rostral**. Going down towards the spinal cord and cauda equina is going **caudal**. This is a brain cut in the **frontal** plane. Unstained brain tissue shows up as grey (actually pinky-grey) and white matter



Staining the brain tissue is essential to differentiate structures. In this stain cell bodies are stained blue. Thus the cortex can be seen to contain large numbers of nerve cell bodies

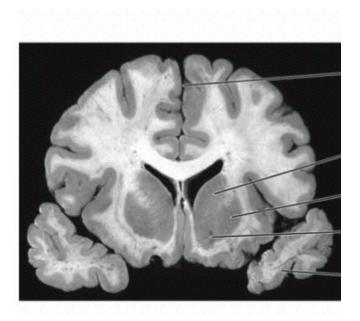


Grey matter = cell bodies & processes

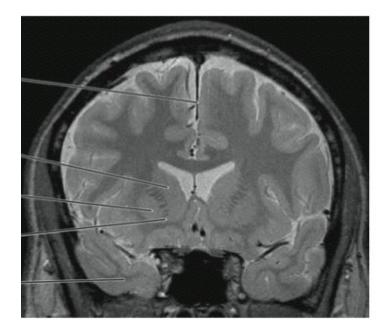
White matter = axons



Nowadays MRI enables us to see a histology-like picture in the living brain.

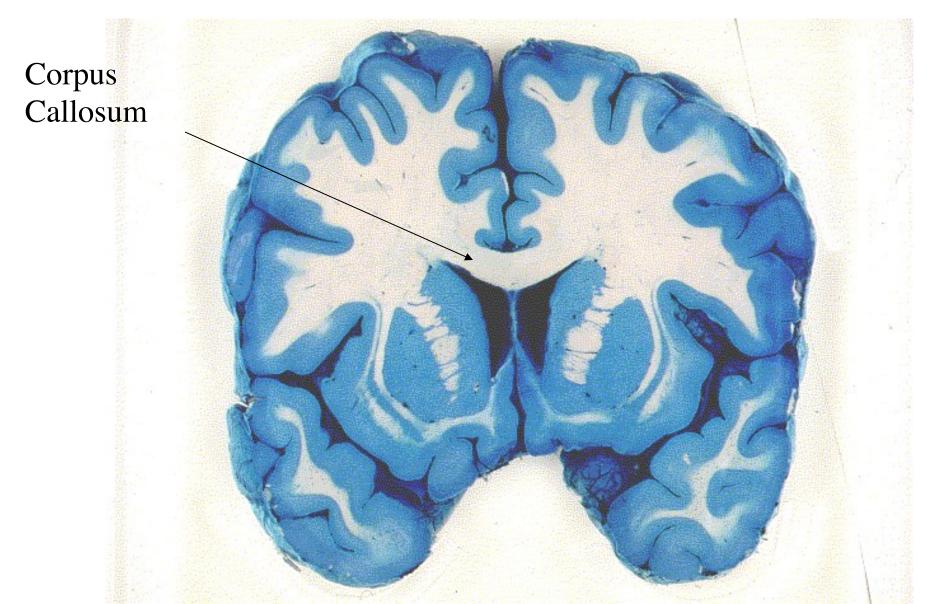


Stained post-mortem



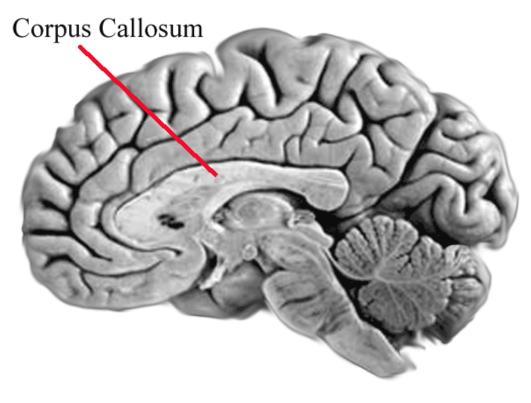
Living

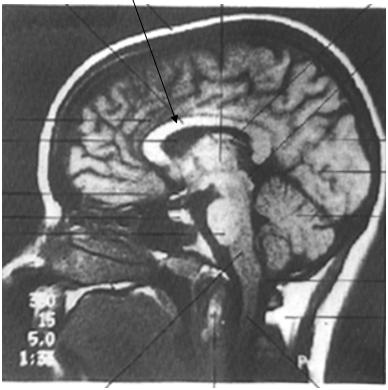
The corpus callosum is the most important landmark in the brain. It is a **bridge of axons** that joins the two hemispheres and allows communication between them.



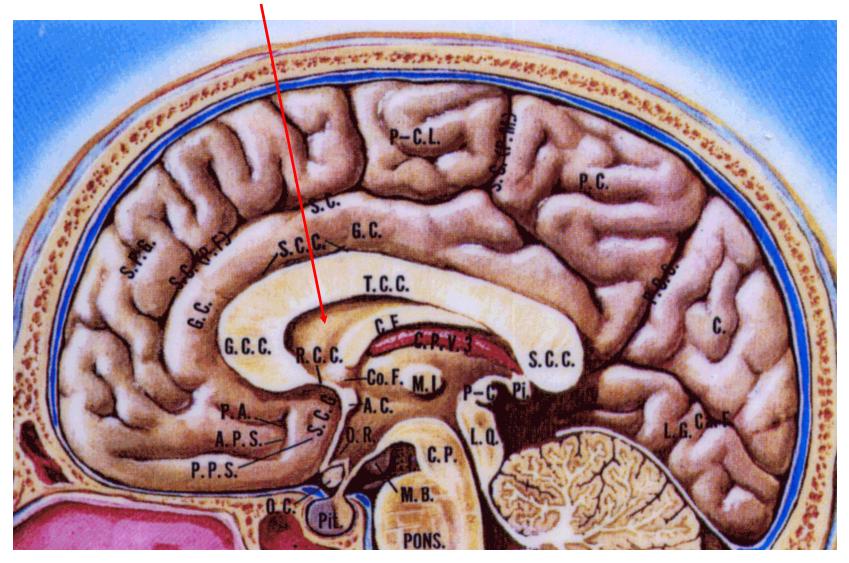
The corpus callosum is very easy to identify in the midsagittal plane (below).

Corpus callosum is easy to see on an MRI



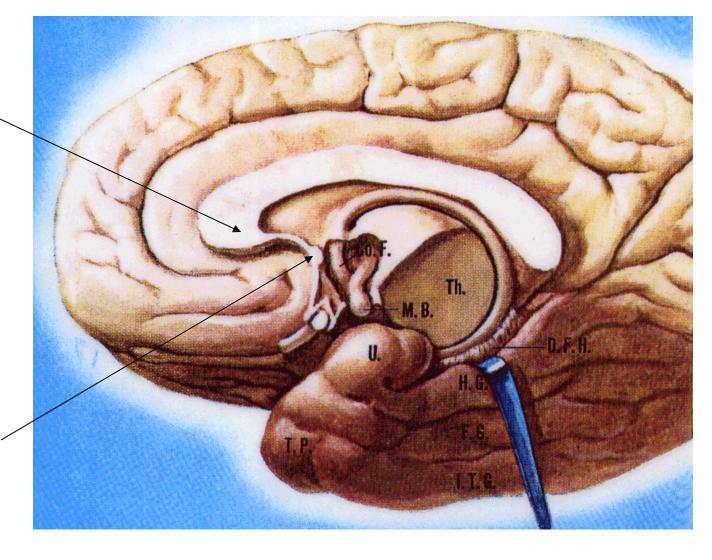


Underneath the anterior part of the corpus callosum we can see the lateral ventricles



Note that the corpus callosum folds back on itself rostrally. This region ends in the **anterior commissure**

'Folded back' region



Anterior commissure

ALILA MEDICAL MEDIA

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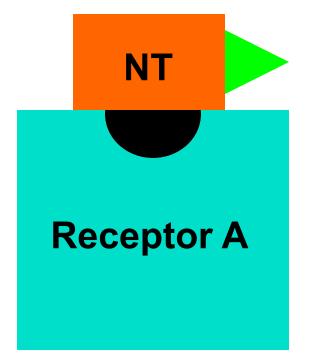
CRITERIA

- NT found in axon terminals
- NT released by action potentials
- Synthesis identified
- External application mimic normal Response
- Pharmacology same for normal and externally applied NT ~

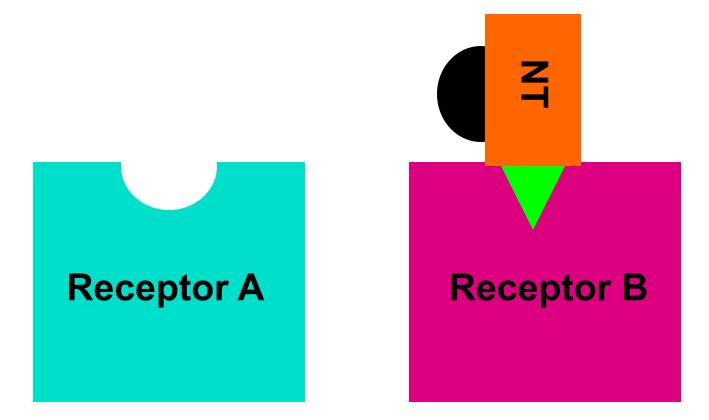
Lock & Key Model

NT binds to receptor NT = key**Receptor = lock** Receptor changes shape determines if EPSP or IPSP receptor subtypes ∎ NOT NT ~

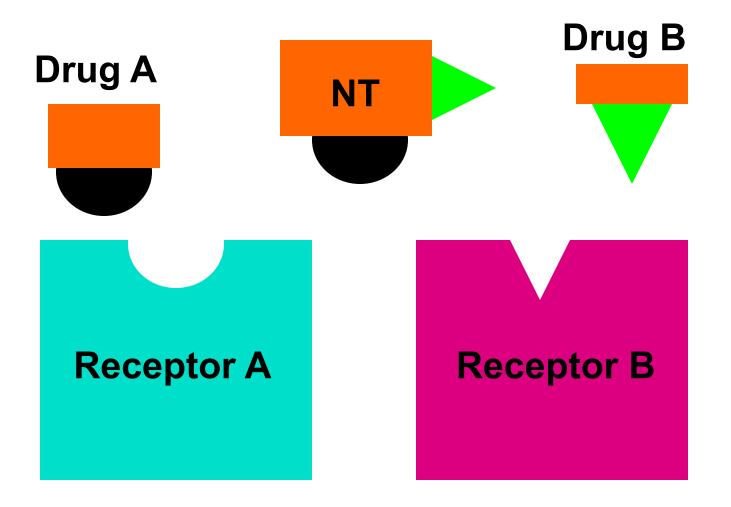
ligand binds to receptoractivation: + or - ~



Same NT can bind to different -R different part of NT ~



Specificity of drugs



The Chemical Synapse Images courtesy of McGraw-Hill Higher Education

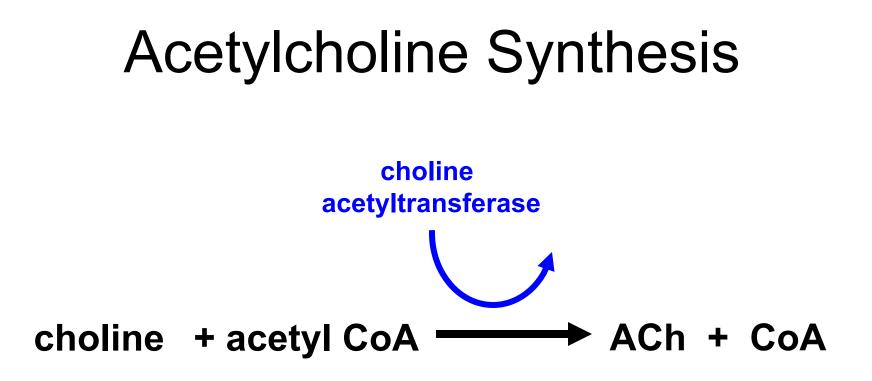
Check us out on twitter and FB. @StudentHelp4AP

Acetylcholine - ACh

- Most abundant NT in Peripheral N.S.
 - also found in Central N.S.
- Precursor = choline
 - nutrient
- Degraded by acetylcholinesterase-
 - AChE

Membrane bound - pre- & postsynaptic

- Nicotinic receptor ionotropic
- Muscarinic receptor metabotropic ~



Ach - Distrubution

- Peripheral N.S.
- Excites somatic muscle
- Autonomic NS
 - Ganglia
 - Parasympathetic NS Neuroeffector junction
- Central N.S. widespread
 - Hippocampus
 - Hypothalamus ~

Cholinergic Agonists

Direct

- Muscarine
- Nicotine
- small doses
- Indirect
 - AChE Inhibitors ~

AChE inhibitors

- Physostigmine
- Organophosphates irreversible
 - DFP
 - Soman & Sarin
 - Malathion*
- Agonist or Antagonist? indirect agonist ~

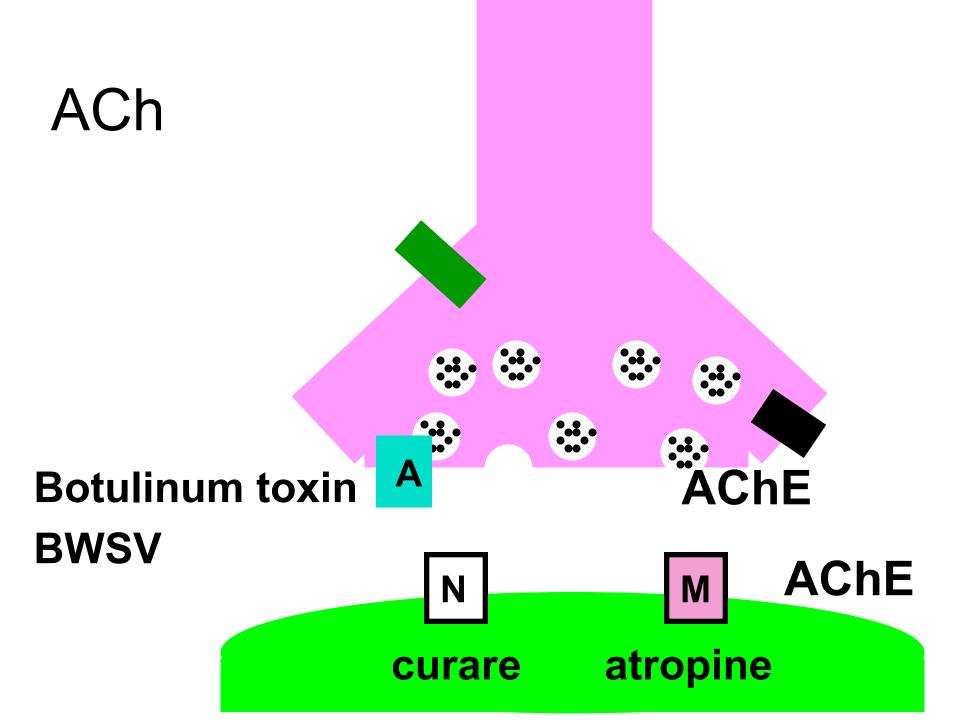
Cholinergic Antagonists

Direct

Nicotinic - Curare Muscarinic - Atropine Scopolamine

Indirect

Botulinum Toxin Black Widow Spider Venom ~



Monamines

- Amino acid precursors
 - single amine group
- 2 groups

Catecholamines - catechol ring Indolamine - indole ring

Affected by many of same drugs ~

Monoamines

Catecholamines

Dopamine - DA

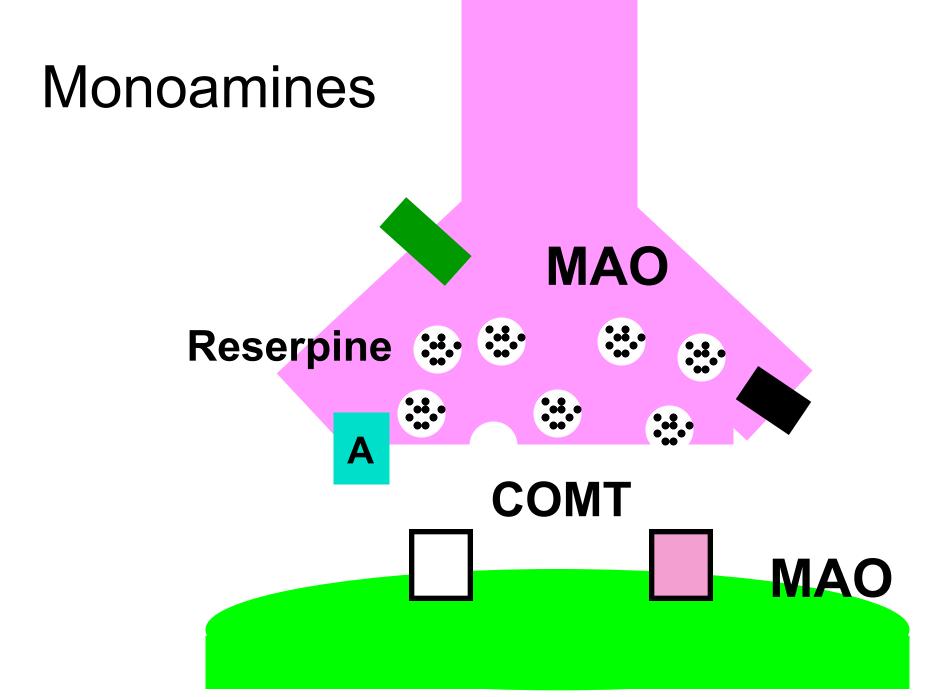
- Dopaminergic
- **Norepinephrine NE**
- Noradrenergic
- **Epinephrine E**
- Adrenergic ~

■ <u>Indolamines</u>

- Serotonin 5-HT
 - Serotonergic

Monoamines

Terminated by... reuptake monoamine oxidase - MAO catechol-O-methyltranferase - COMT also in liver Reservine ----> leaky vesicles depletes monoamines ~



Indirect Monoamine Agonists

MAOIs

Iproniazid

- Reuptake blockers
 - Tricyclic antidepressants
 Imipramine

Desipramine

Cocaine & Amphetamine ~

Dopamine

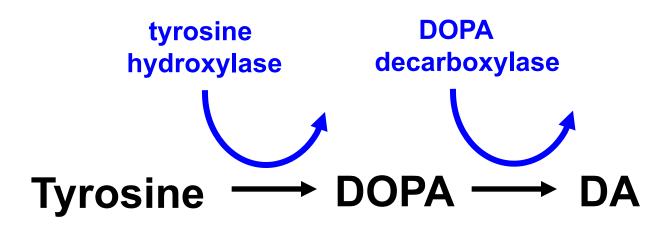
- Only in central nervous system mostly inhibitory systems
- Reward
- Schizophrenia
- Movement
 - Nigrostriatal Pathway
- At least 5 DA-R types: D_1 , D_2 , etc. ~

Dopaminergic Drugs

Agonist

- L-dopa
- Antagonists
 - Chlorpromazine
 - **D**₁
 - Haloperidol
 - D₂ ~

Dopamine Synthesis



Norepinephrine

Peripheral N.S.

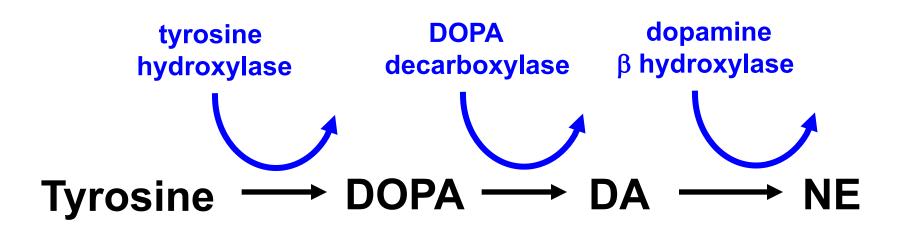
- Sympathetic neuroeffector junction
- Adrenal glands
- Central N.S.
 - Hypothalamus
 - Locus coeruleus
- Alpha & Beta receptor subtypes
 - NE_{α} & NE_{β} ~

Noradrenergic Drugs

Agonists

- Mescaline
- Ephedrine
- Antagonist
 - Propranalol -
 - beta receptors ~

Norepinephrine Synthesis



Serotonin

NOT a catecholamine Peripheral 98% in blood & smooth muscle ■ Central N.S. Raphe nucleus Hypothalamus **R** subtypes: $5HT_1 \& 5HT_2 \sim$

Sertonergic Drugs

Agonists

• SSRIs

Selective Serotonin Reuptake Inhibitors

- Buspirone
- MDMA

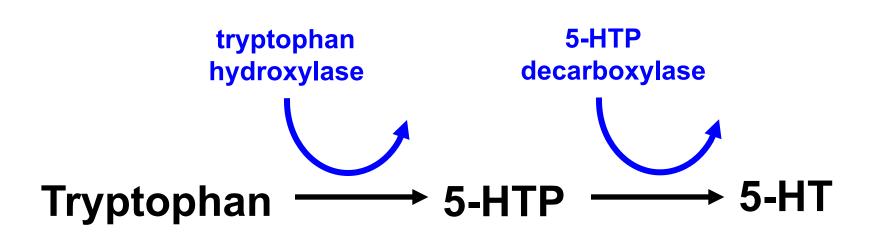
Ecstacy ~

Sertonergic Drugs

Antagonists

- Psilocybin
- LSD in CNS
- Inverse agonist
 - ketanserin ~

Serotonin Synthesis



Gamma-aminobutyric acid

- GABA GABAergic
- Major NT in brain inhibitory system
- Receptor subtypes
 - **GABA_A controls CI- channel**
 - GABA_B controls K+ channel
- Precursor = glutamate ~

GABAergic Drugs

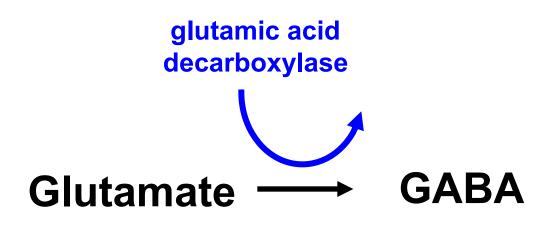
Agonists

- Benzodiazepines
- Barbiturates
- Ethyl alcohol (ETOH)
- Antagonists
 - Picrotoxin
- Inverse agonist
 - Ro 15-4513
 - •ß CCM ~

GABA Synthesis & Reuptake

From Krebs cycle metabolism of glucose in mitochondria From Glial cells GABA ---> Glutamate ---> Glutamine Glutamine into neurons ■ After release GABA back into glia ~

GABA Synthesis



Neuropeptide

- Chains of amino acids
- Synthesis in soma
- Often neuromodulators
 - alters sensitivity of neurons
 - slower, longer-lasting effects
- Substance P pain signaling
- Endorphins analgesia, euphoria ~

Endorphins

Opioids Dynorphin met-enkephalin leu-enkephalin **Beta-endorphin** Receptor subtypes: mu1, mu2, kappa, delta, omega ~

Endorphins (cont.)

Agonists

- morphine
- heroin
- codeine
- Antagonists
 - naloxone
 - naltrexone ~

Other NTs

- Excitatory amino acids
 - Glutamate & Aspartate

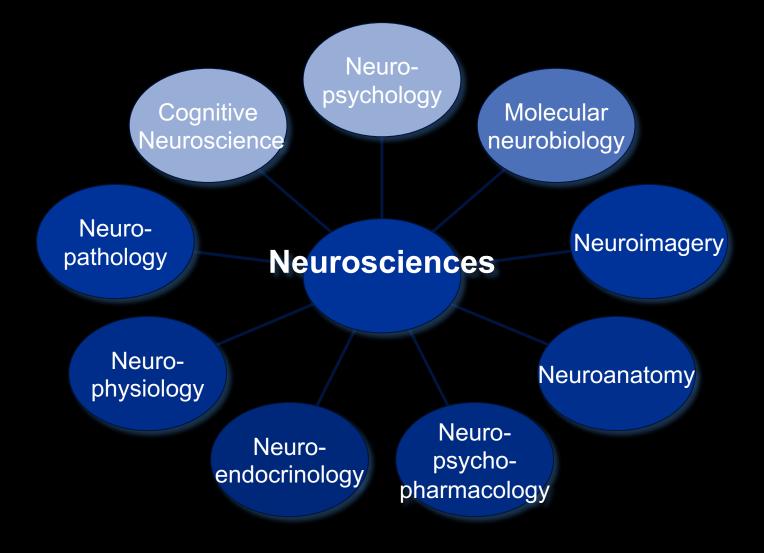
Histamine

- Inflammatory Response
- Nitric Oxide It's a gas
 - Carbon Monoxide?
- Anandamide

ligand for THC-R ~

Why would anyone want to be a neuroscientist?

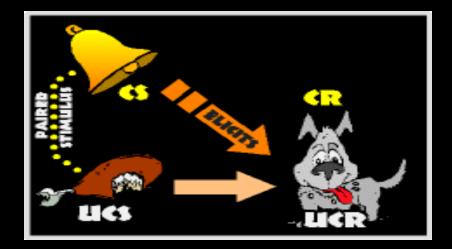
The scientific study of the nervous system and its relationship to cognition and behaviour.



Associative Learning Mechanisms

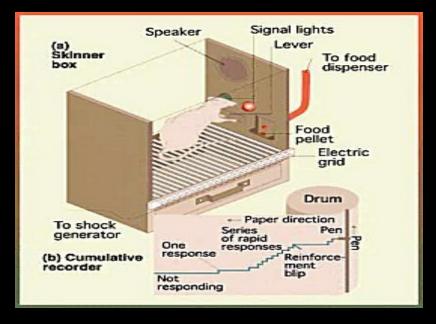
Classical Conditioning

- Discovered by Ivan Pavlov (1920s)
- Passive learning

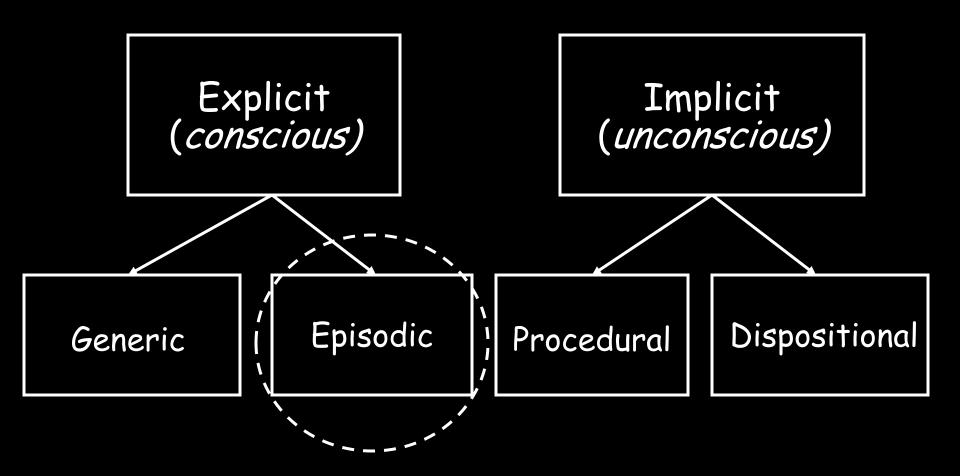


Operant Conditioning

- Discovered by B.F.
 Skinner (1960s)
- Active learning



Types of Memory

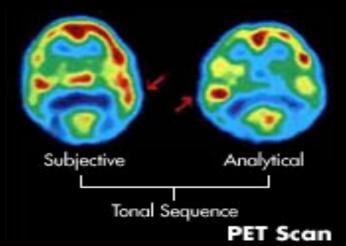


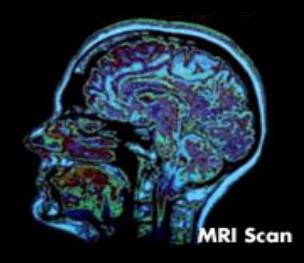
How is the Brain studied?

- Case Studies
- Gene Splicing
- Imaging
 - PET
 - EEG (electrical current detection)
- Transcranial Magnetic Stimulation (causes

temporary disruption of a brain region

- MRI
- fMRI
- -CAT





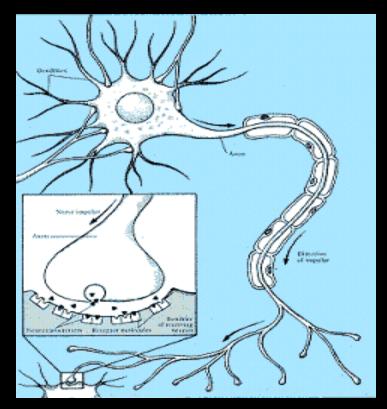
How is learning and memory enabled?

Signal Transduction:

 Inter-Neuron communication occurs via Neurotransmitters at the synaptic gap

Neural Plasticity:

 Through experience, Neurons can change the way they function



Long Term Potentiation:

 Cellular mechanism through which associations can be detected and recorded in the brain

