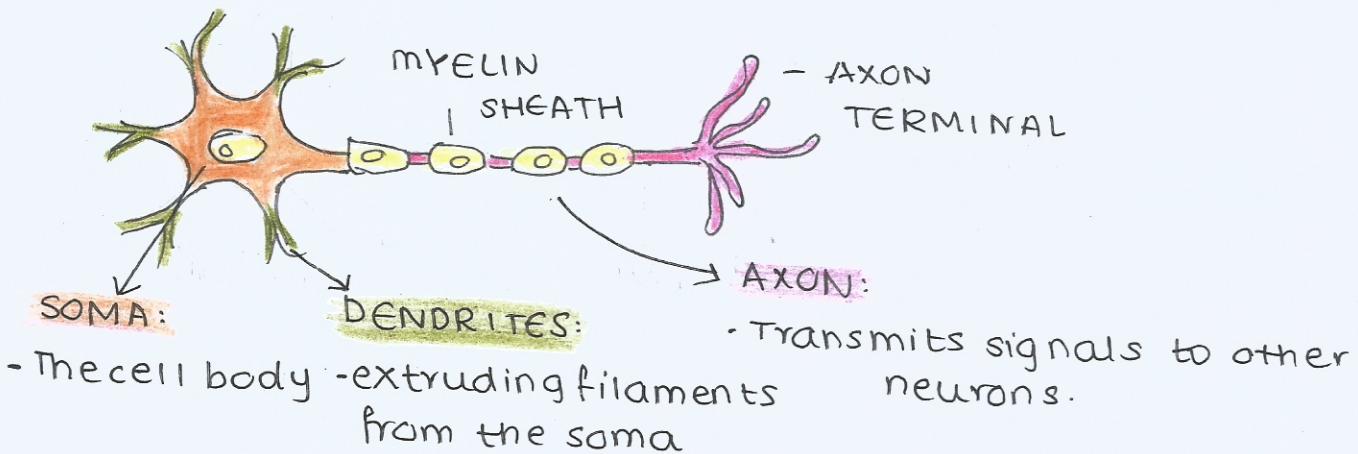


# NEUROTRANSMITTERS AND BEHAVIOUR

## Nervous system processes

Nervous system is a system neurons.

### Neuron



Dendrites and soma receive the signals from other neurons.

### Synapse:

- It is a structure that connects two neurons.
- Axon of one neuron approaches the dendrite/soma of another neuron → synapse is formed.

### Threshold of excitation

- It is the level of excitation that is needed for a neuron to react properly to a stimulus. If the sum of excitation exceeds this threshold, the neuron generates a brief pulse called action potential.

### Neurotransmitters

- They are chemical messengers.
- Constantly synthesized in the neuron
- Stored in the axon terminal.

The electrical pulse reaches the end of axon



at the synaptic gap the mechanism of transmission becomes chemical.

When the action potential reaches the end of an axon ,



A neurotransmitter is released into the synaptic gap.

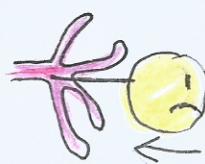
When the neurotransmitter is present in the synaptic gap, it can either

get metabolized  
→ gets destroyed

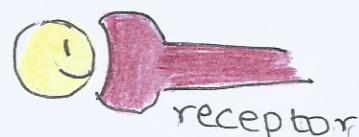


neurotransmitter

reuptake  
→ gets pulled back into the axon terminal



post-synaptic membrane  
→ reach the post-synaptic membrane and bind to the receptors on its surface



## (TWO) TYPES OF NEUROTRANSMITTERS

### Excitatory

- They allow the impulse to pass the synapse.
- produces stimulating effects

### Inhibitory

- They prevent the impulse to cross the synapse.
- produces calming effects

When neurotransmitters are out of their optimal ranges, they can cause mental disorders.

These neurotransmitters are affected by

- Agonists

- Antagonists

**Agonists:** enhance the action of a neurotransmitter

**Antagonists:** counteract a neurotransmitter (prevent a signal from being passed further)

**SSRIs** → Selective serotonin reuptake inhibitors

- ▷ They block the reuptake of the serotonin from the synaptic gap.
- ▷ This increases the concentration of serotonin in the synapse.

### EFFECT OF SEROTONIN ON PROSOCIAL BEHAVIOUR

Crockett et al (2010)

**Aim:** To investigate the effect of serotonin on prosocial behaviour.

**SEROTONIN** - Inhibitory neurotransmitter



- Helps in regulating sleep cycles and
- sustaining stable mood.



**Sample:** 30 healthy subjects (mean age 26)

Repeated measures design with 2 conditions:

Condition one

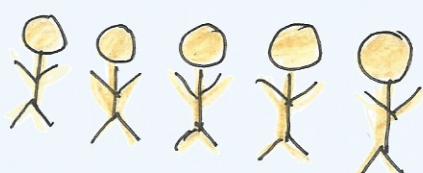
- participants were given a dose of citalopram (SSRI).

Condition two

- participants were given a placebo (a harmless substance with no effect)

**Procedure:**

- Participants were given a series of moral dilemmas;
- They were asked to choose between a Utilitarian outcome  
→ saving 5 lives.



Personal

pushing a man off  
the bridge to stop  
a train & save 5  
people

Arvesive harmful  
actions

diverting the  
track by pulling  
a level to  
save 5 but  
kill 1.

**Results:** The responses in the impersonal version were unaffected by citalopram.

### Condition 1

→ people were less likely to push the man off the bridge in the personal scenario.

### Conclusion:

- Serotonin reduces acceptability of personal harm and promotes pro-social behaviour.

### Limitation:

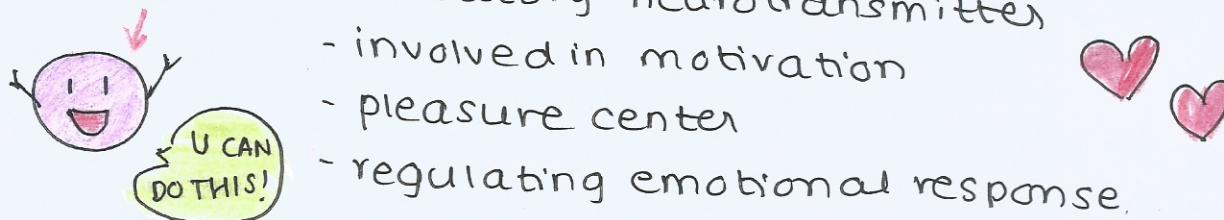
- The citalopram intake caused nausea in the participants.
- The participants could have figured if they were on citalopram or not which might have affected their answers.

## EFFECT OF DOPAMINE ON ROMANTIC LOVE - Fisher, Aron and Brown (2005)

**Aim:** To study the neural mechanisms of romantic love.

### DOPAMINE:

- excitatory neurotransmitter
- involved in motivation
- pleasure center
- regulating emotional response



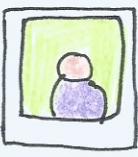
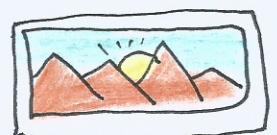
**Sample:** 10 men and 7 women - intensely in love but not with each other.

- Mean age: 21 years
- Mean duration of being in love: 7 months

### Procedure:

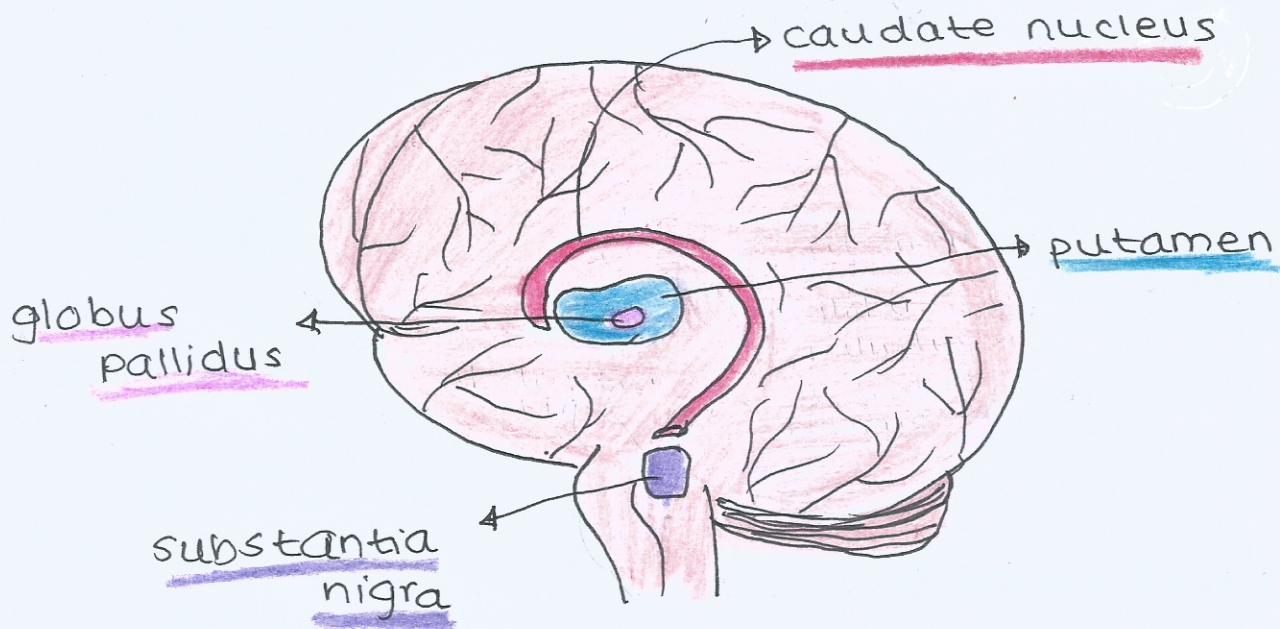
- Participants were placed inside a fMRI and were asked to look at photographs of their loved ones while their brains were being scanned.

Four stages: (Repeated 6 six time → 720 seconds total)

1. For 30 seconds each participant viewed a photograph of his or her beloved person.
2. Participants were given a 40-second filler activity which was to count back from a given number.  100, 99, 98...
3. For 30 seconds more, participants viewed a photograph of an emotionally neutral acquaintance.
4. Final stage - 20 seconds of counting back from a number. 50, 49... 

### Results:

- Specific parts of the brain were activated when the pictures of their loved ones were shown.
- Caudate nucleus and Ventral Tegmental Area were primarily activated.
- They are rich in dopamine and are a key part of the **Dopaminergic pathway**.
  - It is a system that generates and transmits dopamine. It also increases dopamine related activity in the brain.



## THE ROLE OF SEROTONIN IN DEPRESSION

Low levels of serotonin in the brain play a causal role in developing depression.

Clinical trials included 2 experimental groups.

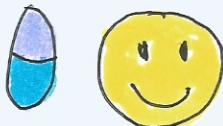
Group-1: Given drugs that affect the levels of serotonin.

Group-2: Given a placebo which had no effect on the serotonin level.

Results: In group-1 the drugs (SSRIs) lead to a reduction of symptoms of depression.

### LIMITATIONS:

- ▷ The drugs affect neurotransmitters immediately but the behavioural affects take time.
- ▷ Not all patients benefit from drugs. The link between serotonin and depression isn't universal.
- Depression was linked to 5-HTT gene which is a serotonin transporter gene.
- This gene determines a person's vulnerability to developing depression.



## THE ROLE OF DOPAMINE IN PARKINSON'S

DISEASE Freed et al (2001)

Aim: To study the role of dopamine in people with parkinson's disease.

PARKINSON'S DISEASE: It is a degenerative disorder that affects the motor functions of the nervous system.

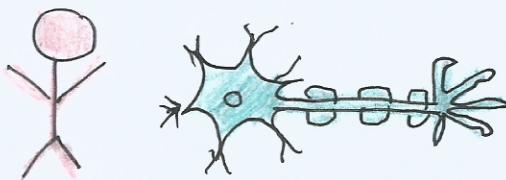
Sample: 40 patients (34-57 years old) who had severe parkinson's disease for the mean duration of 14 years.

Procedure: Randomly divided into 2 groups

GROUP-1  
(experimental)

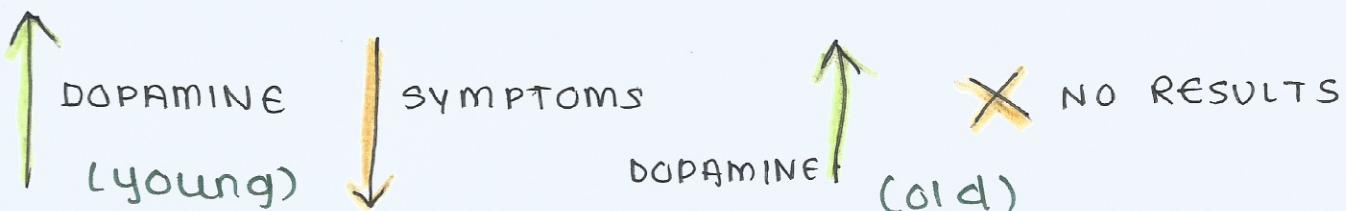
GROUP-2  
(control group)

- transplant of nerve cells • sham surgery
- In the 1st group, nerve tissues that contained dopamine producing neurons were transplanted into their putamen.
- These nerve tissues were taken from embryos who were aborted 7-8 weeks after conception.
- Putamen → part of limbic system that helps in movement regulation.
- In the 2nd group, holes were drilled in the skull but the dura wasn't penetrated.
- The surgeries for both groups were identical otherwise.



### Results:

- PET scans showed increase growth of dopamine-producing cells in the putamen.
- Reduction of symptoms by 28% in the transplant group. (only in young patients - 60 or younger)
- No improvements in the older patients (aged over 60)



### Conclusion:

- The transplant of dopamine producing neurons resulted in some improvement in the young patients and not the older ones.
- Less response to treatment in the older patients might be because of lower neuroplasticity of the brain.

### Protocols followed:

- The consent form describing the risks and benefits were approved by the ethics committee.
- Written consent form was used for the women who donated the fetal tissues from abortions.
- Clinical observations and brainscans were followed up for a year.