

# Primary oligodendrocyte precursor cell culture

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## SCOPE OF THE METHOD

<b>The Method relates to</b>	Animal health, Human health
<b>The Method is situated in</b>	Basic Research
<b>Type of method</b>	In vitro - Ex vivo
<b>This method makes use of</b>	Animal derived cells / tissues / organs
<b>Species from which cells/tissues/organs are derived</b>	Mus Musculus
<b>Type of cells/tissues/organs</b>	Brain (cortex)

## DESCRIPTION

### Method keywords

oligodendrocyte

shakeoff

cell culture  
isolation

### **Scientific area keywords**

basic research  
fundamental research  
differentiation  
neuroscience

### **Method description**

This method describes the steps from a living mouse to a single cell solution of primary oligodendrocyte precursor cells.

### **Lab equipment**

### **Method status**

Internally validated

### **PROS, CONS & FUTURE POTENTIAL**

#### **Advantages**

Primary cultures give rise to a condition more similar although not identical to the *in vivo* situation when compared to cell line experiments ;

Methodologically feasible ;

Highly reproducible ;

Astrocytes can be simultaneously isolated ;

Oligodendrogenesis can be evaluated purely.

## Challenges

Interspecies differences ;

Terminal experiment for the lab animal ;

Time consumable (2 weeks to reach an OPC culture, additional time required to reach oligodendrocyte stage).

## Future & Other applications

The protocol can be adapted and used in other animal species.

## REFERENCES, ASSOCIATED DOCUMENTS AND OTHER INFORMATION

### Associated documents

[Primary OPC isolation mouse .docx](#)

## PARTNERS AND COLLABORATIONS

### Organisation

**Name of the organisation** Hasselt University

**Department** Biomed Neuro-Immune Connection and Repair

**Country** Belgium

**Geographical Area** Flemish Region

*Coordinated by*



*Financed by*

