# Nervous system development in Vertebrates

Caroline Borday

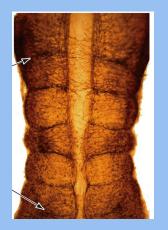
23<sup>th</sup> september 2025



- Nervous system description
  - Nervous tissue

Nervous system development
Neurulation



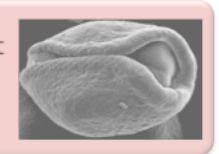


- > Neural induction
- Anteroposterior regionalization
  Rhombencephalon
  - Dorsoventral regionalization
- Migration ans synaptogenesis



- Nervous system description
  - Nervous tissue

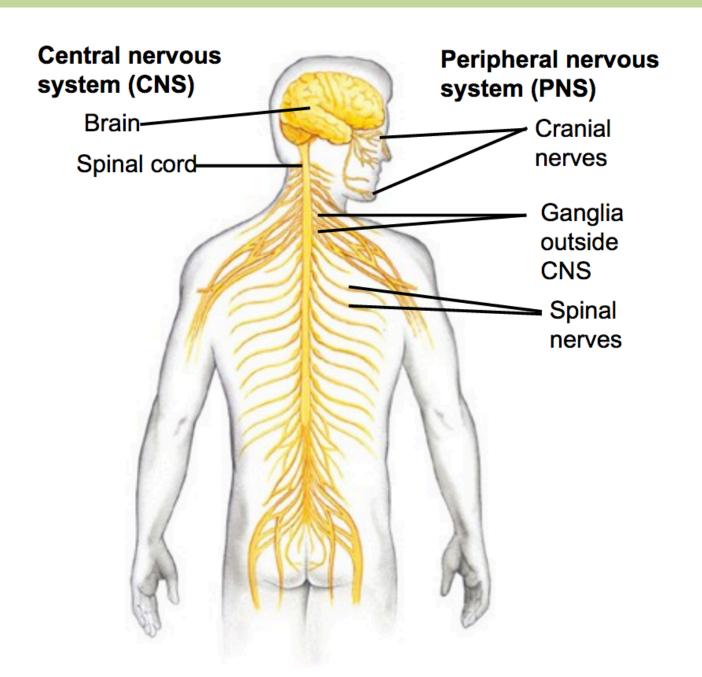
Nervous system development
Neurulation



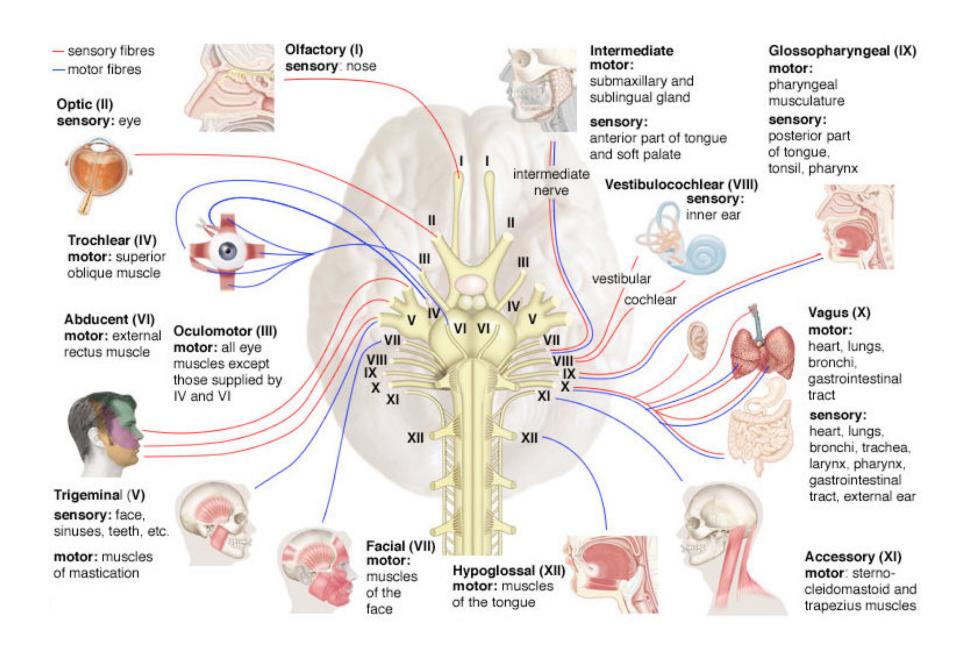


- Neural induction
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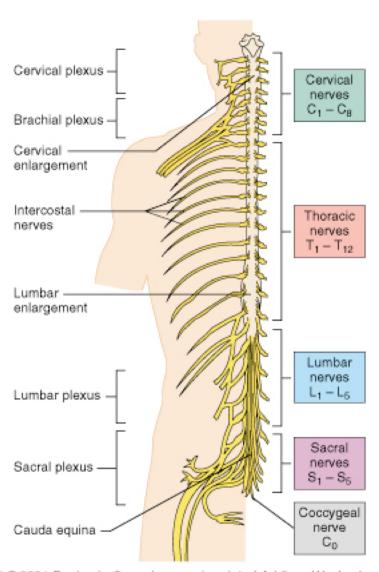
#### The nervous system: physiological point of view



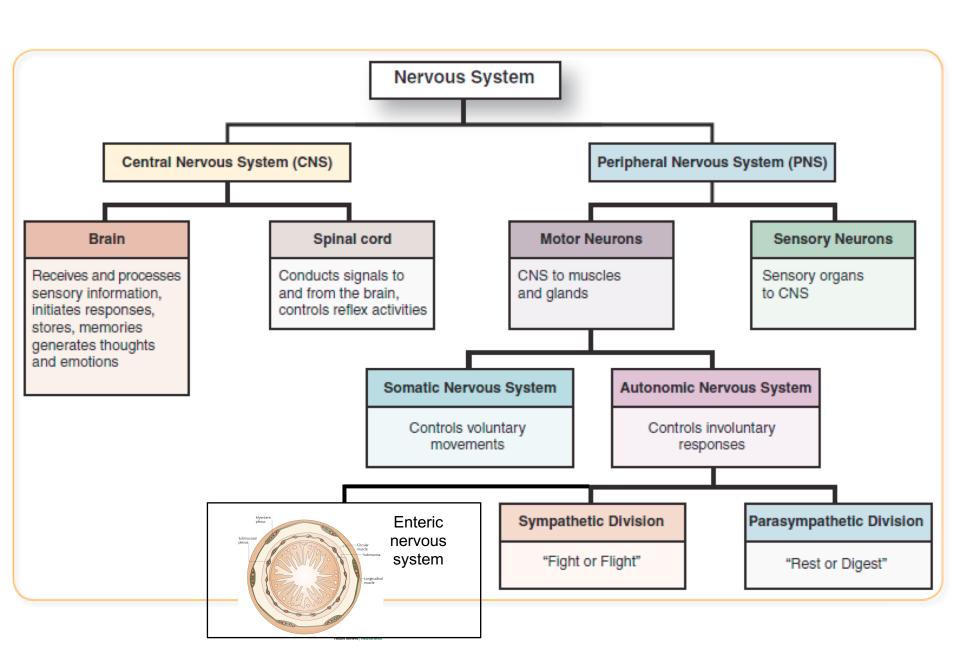
#### **Cranial nerves**



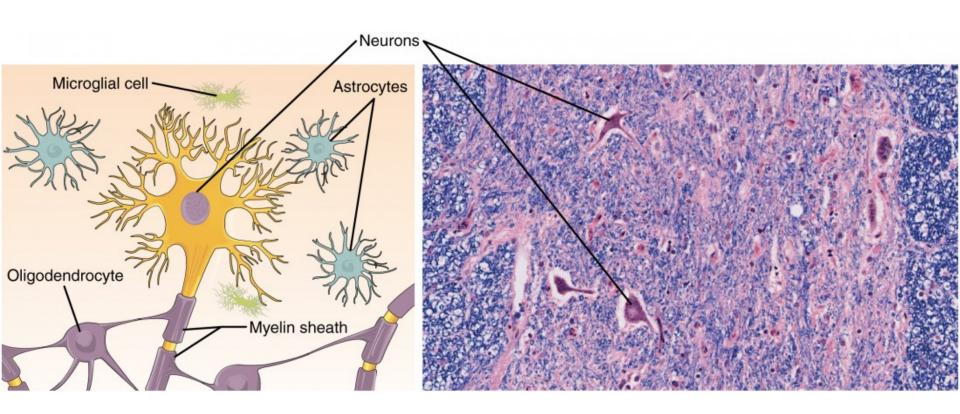
#### **Spinal nerves**



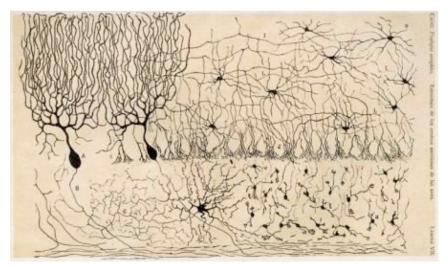
Copyright © 2001 Benjamin Cummings, an imprint of Addison Wesley Longman, Inc.



#### **Nervous tissue**



#### **Neuronal diversity**

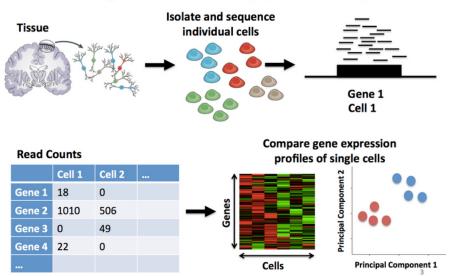


Ramon y Cajal, 1905

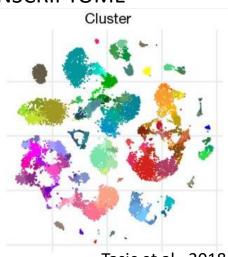
- Morphology
- Connectivity
- > Electrical properties
- Neurotransmitter

...

#### Single-cell RNA-Seq (scRNA-Seq)

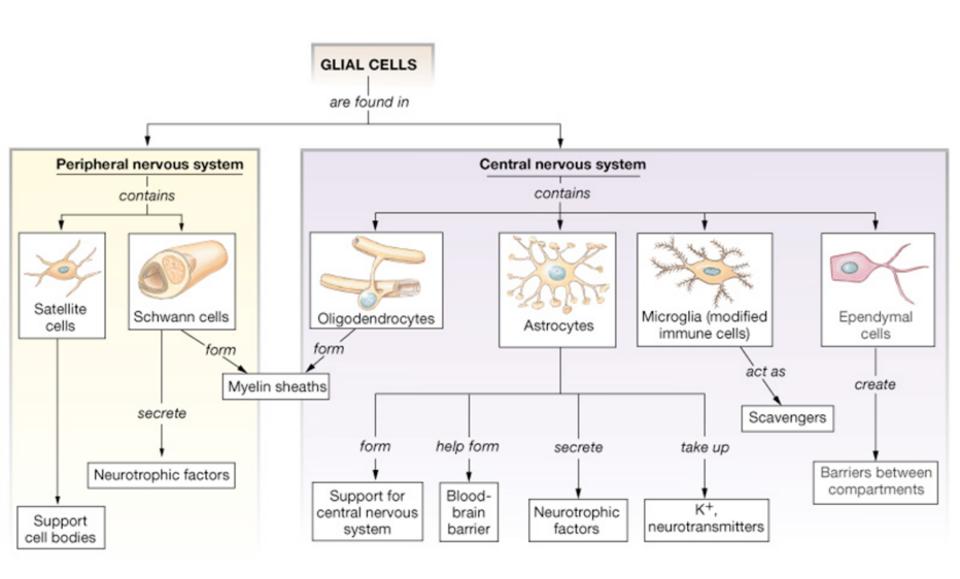


Around 130 neuronal cell types in the mouse visual cortex



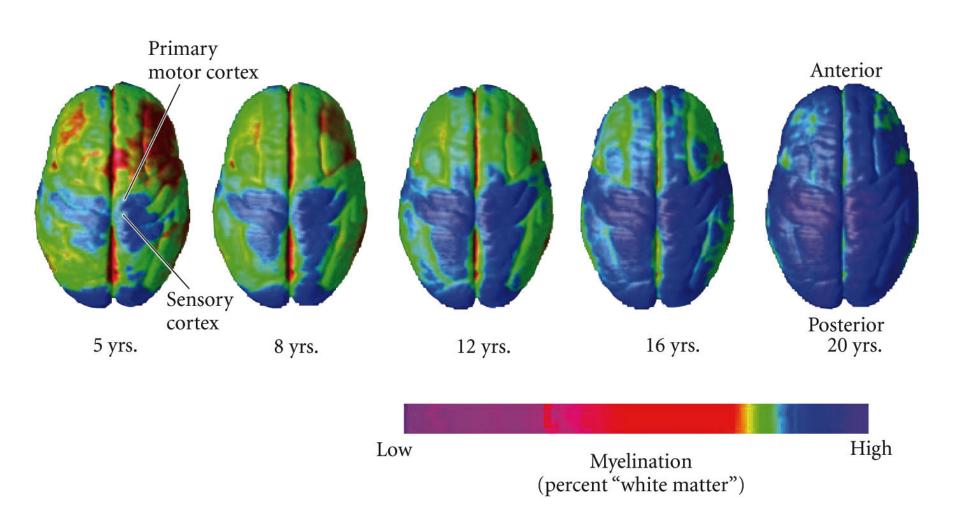
Tasic et al., 2018

#### Glial cells and their functions

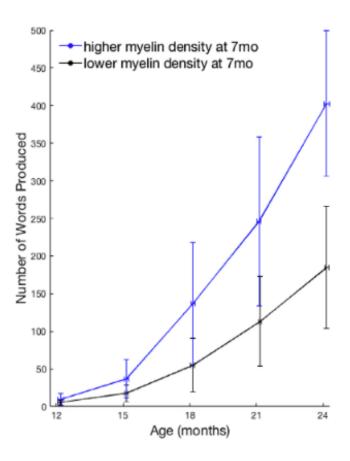


#### The myelination process

Dorsal view of the human brain showing the progression of myelination ("white matter") over the cortical surface during adolescence

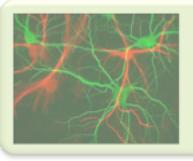


#### The myelination process



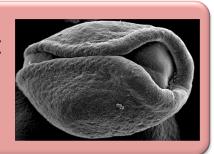


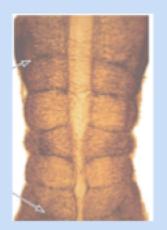
correlations between myelination at 7 months of age and vocabulary production at 24 months of age



- Nervous system description
  - Nervous tissue

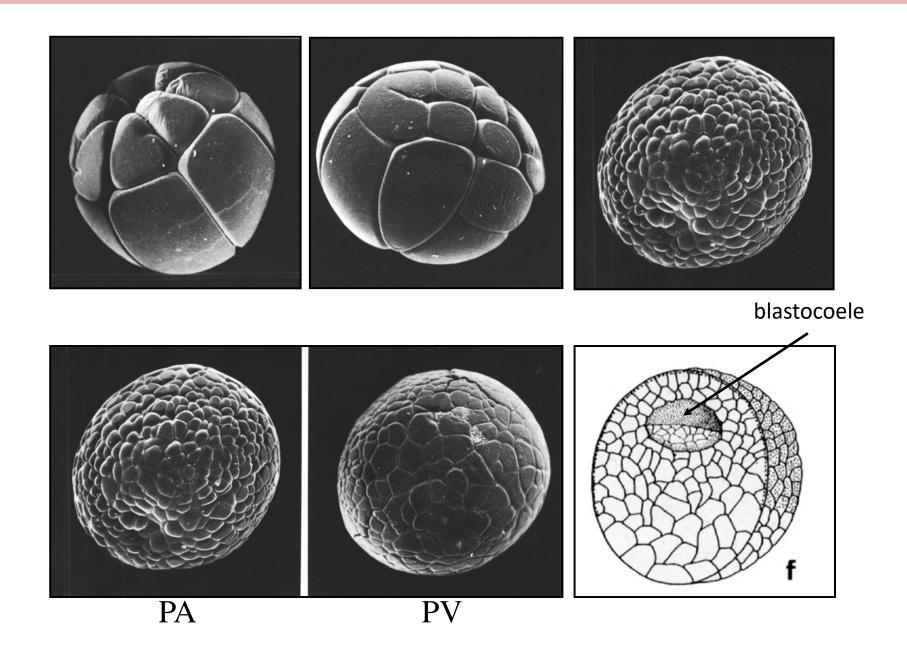
Nervous system development
Neurulation





- Neural induction
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  Rhombencephalon
  - Dorsoventral regionalization
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#### Cleavage (amphibian)



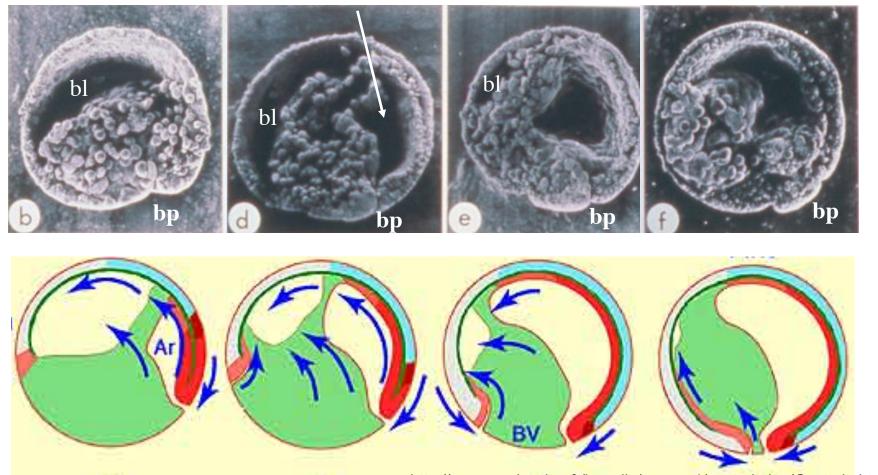
#### **Gastrulation (amphibien)**

Blastoporal superior lip В

D

#### **Gastrulation (amphibian)**

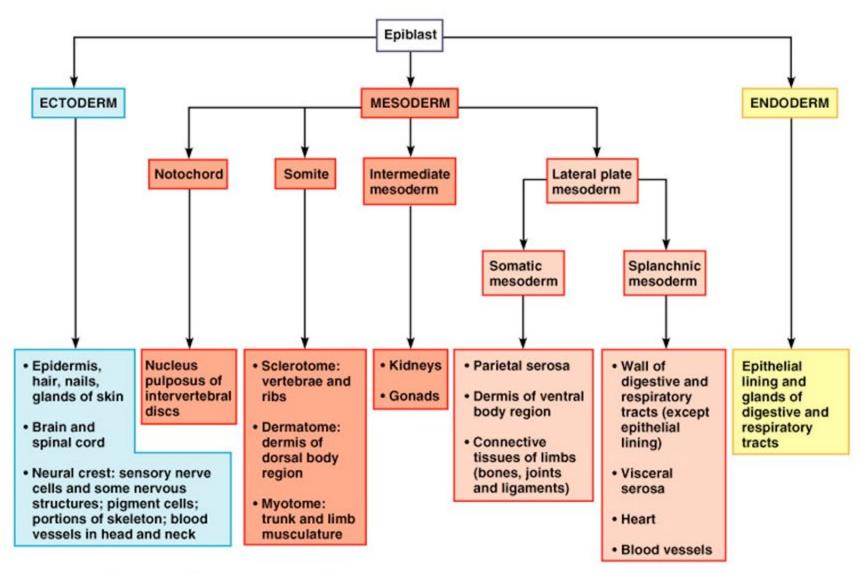
#### archenteron



http://www.snv.jussieu.fr/bmedia/xenope1/gastrulation/Gastrula.html

**Ectoderm Mesoderm Endoderm** 

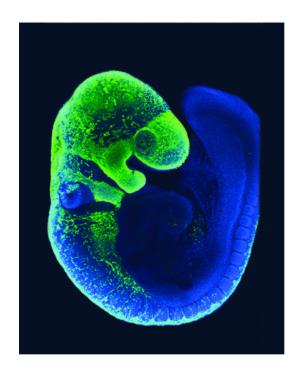
# Many structures are derived from the three embryonic germ layers during organogenesis

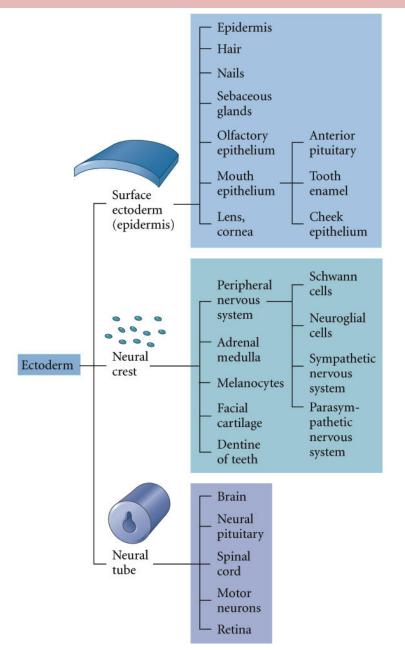


#### Major derivatives of the ectoderm

#### Three subdivisions of the ectoderm:

- Surface Ectoderm
- Neural Crest
- Neural Tube

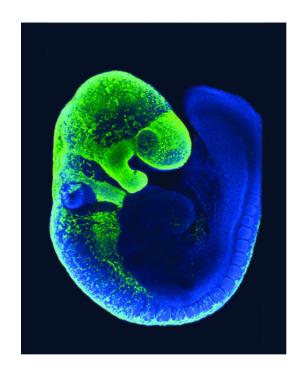


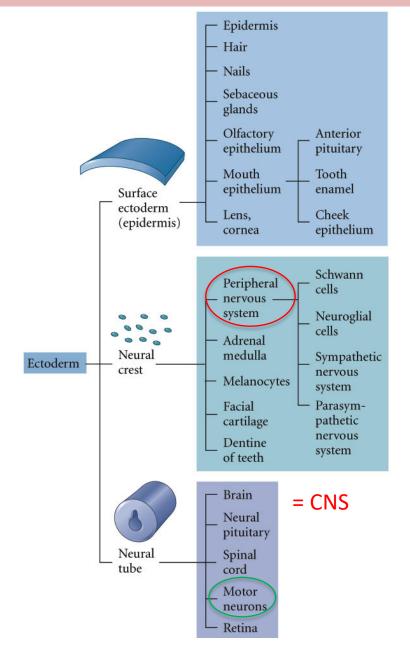


#### Major derivatives of the ectoderm

#### Three subdivisions of the ectoderm:

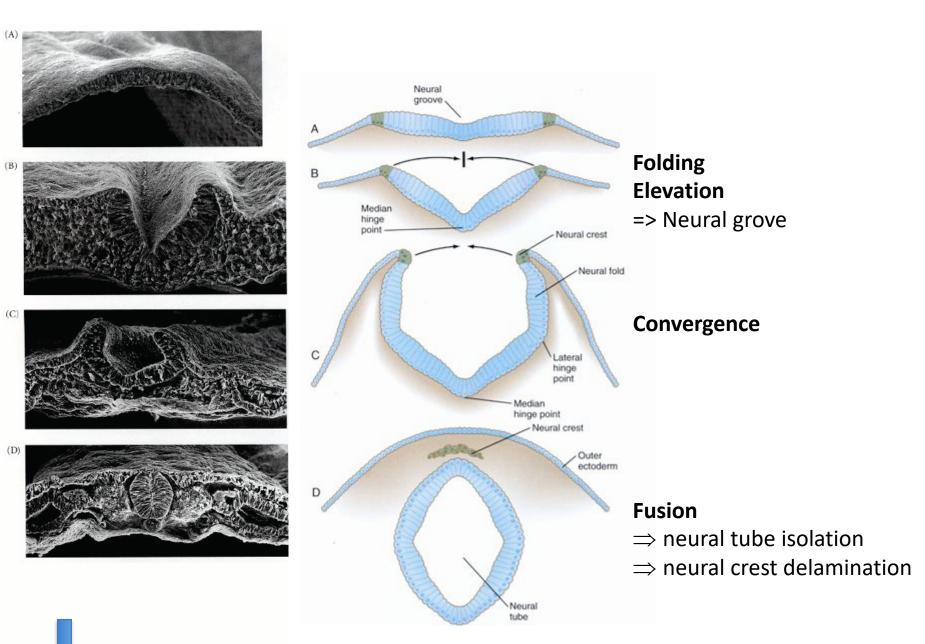
- Surface
- Neural Crest
- Neural Tube





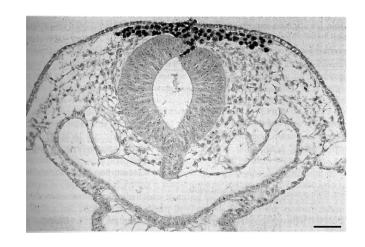
=> Embryological point of view of the nervous system subdivisions

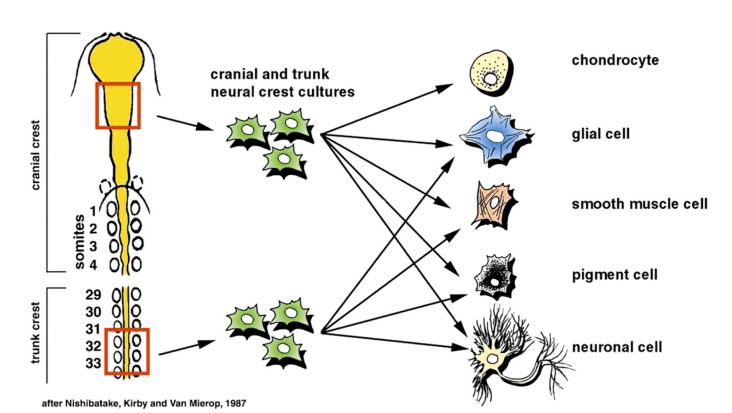
#### **Primary neurulation (chick)**



proliferation allowing growth in thickness

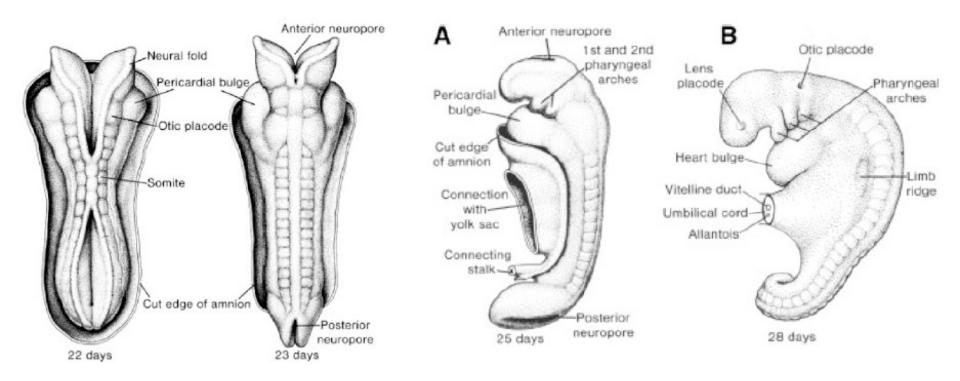
#### The neural crest cells



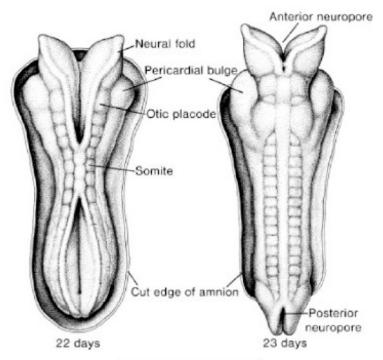


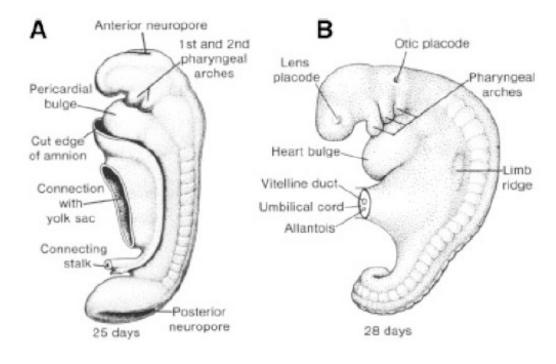
# Neurulation in the *xenopus* embryo

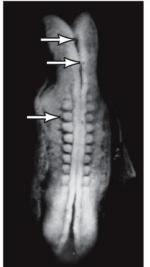
#### **Neurulation in human**

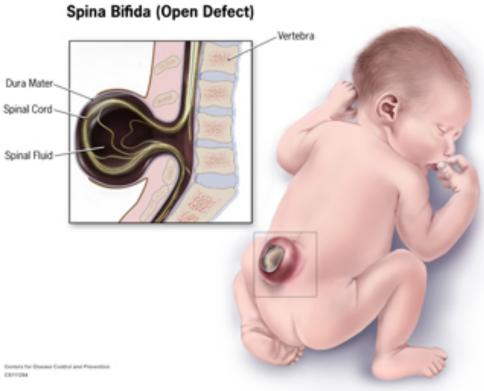


#### **Neurulation in human**











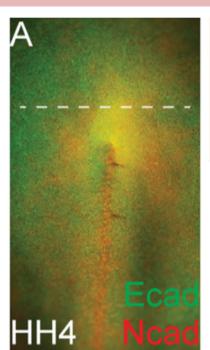
Lightest form: asymptomatic

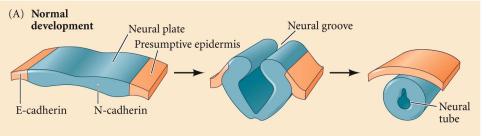


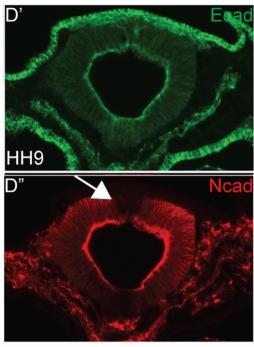
Spina Bifida = defect occurring at the posterior part of the neural tube. (prevalence: 1/2000)

- $\Rightarrow$  Surgery
- ⇒ Often times, locomotor disorders

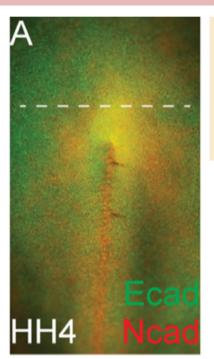
# Expression of N- and E-cadherin adhesion proteins during neurulation in *chick*

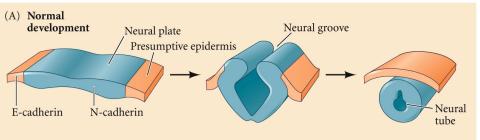




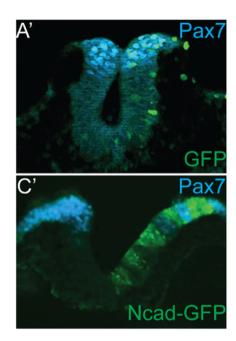


# Expression of N- and E-cadherin adhesion proteins during neurulation in *chick*

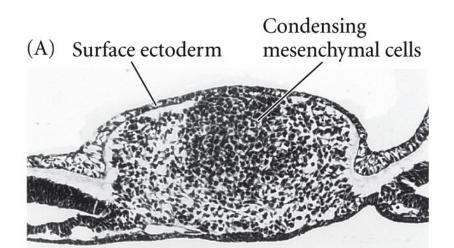


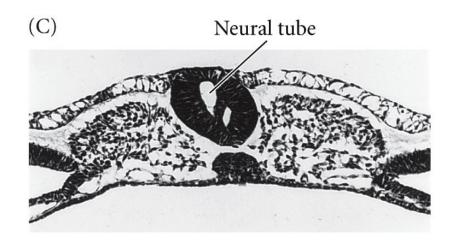


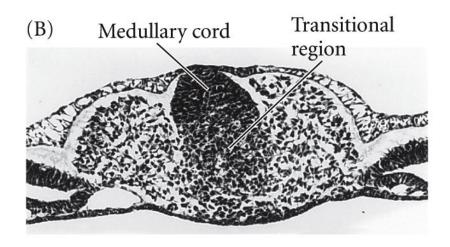


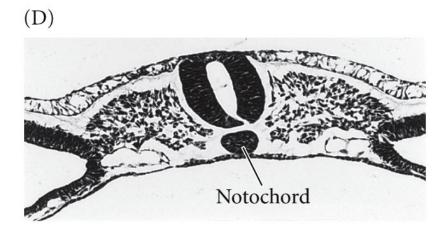


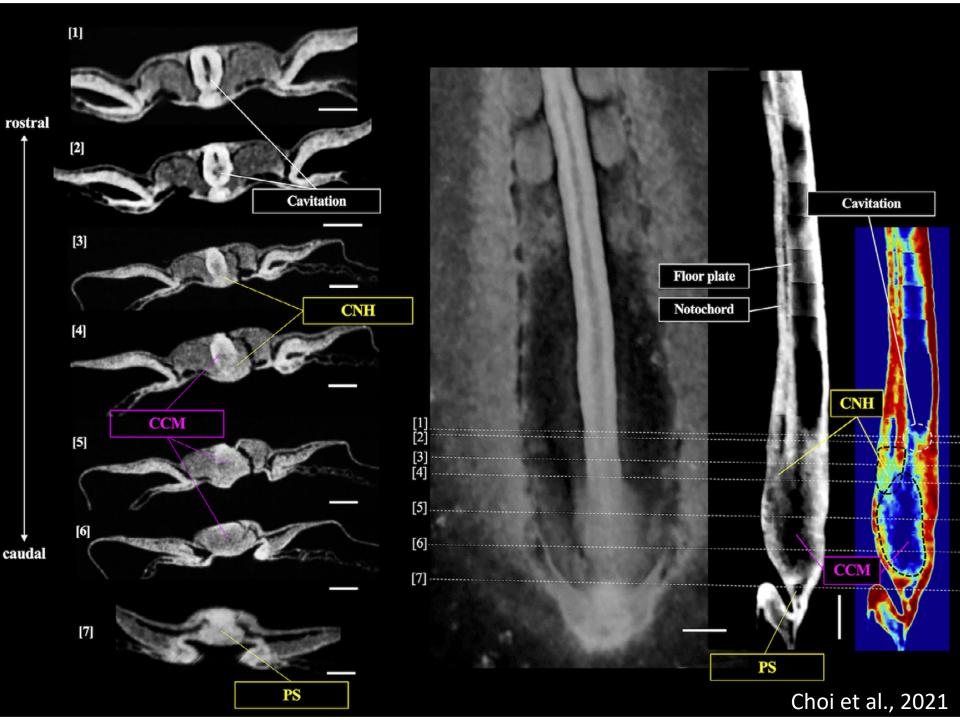
# Secondary neurulation: example in the caudal region of a 25-somite chick embryo









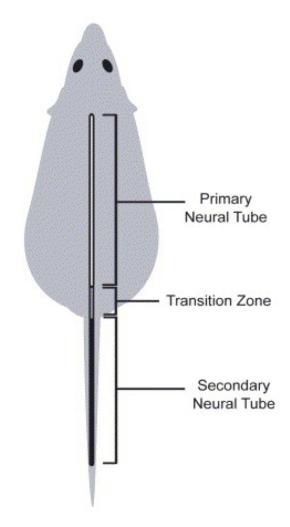


# Secondary neurulation: in Human

# Caudal eminence becomes continuous with neural canal Neural tube 20 days Lumen of caudal eminence becomes continuous with neural canal Neural canal 40 days

Source: Neurosurg Focus © 2010 American Association of Neurological Surgeons

A

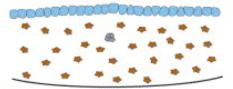


neuromesodermal common progenitor cells

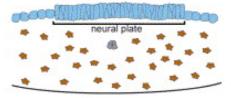
neural restricted lineage

#### **Primary Neurulation**

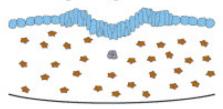
1. Initial epithelium



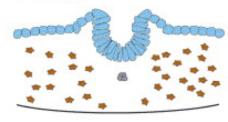
2. Columnarization



3. Rolling/folding



4. Closure

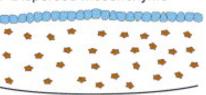


5. Neural tube complete

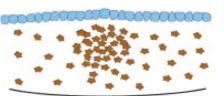


#### Secondary neurulation

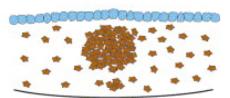
1. Dispersed mesenchyme



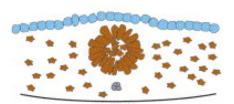
2. Mesenchymal condensation



3. Medullary cord/neural rod



4. Epithelial transition/cavitation

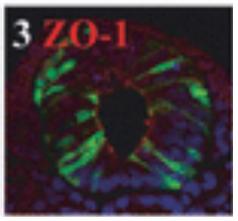


5. Neural tube complete

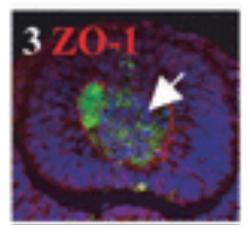


#### Mesenchymal-epithelial transition during secondary neurulation is regulated by differential roles of Cdc42 and Rac1





#### **CDC42** activation



#### **Rac1 inhibition**

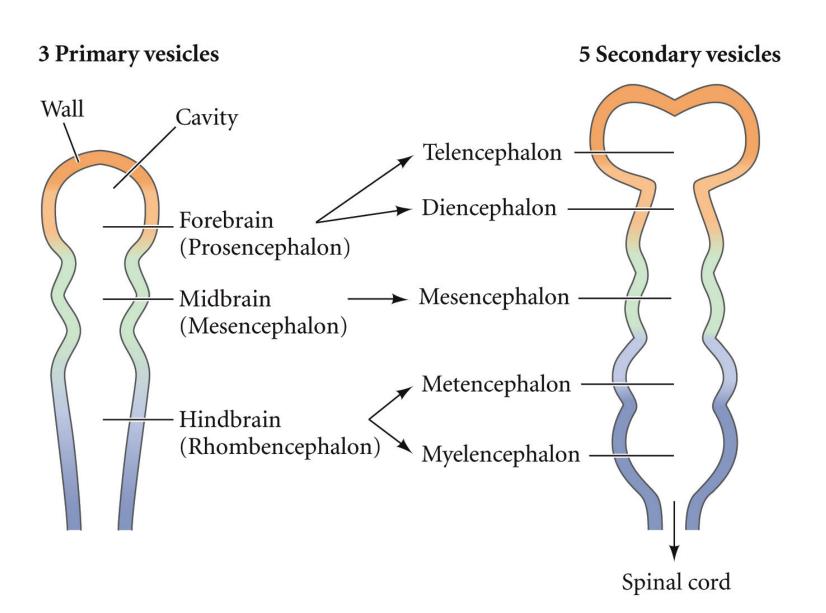


D'après Shimokita and Takahashi, 2011

Electroporated cells

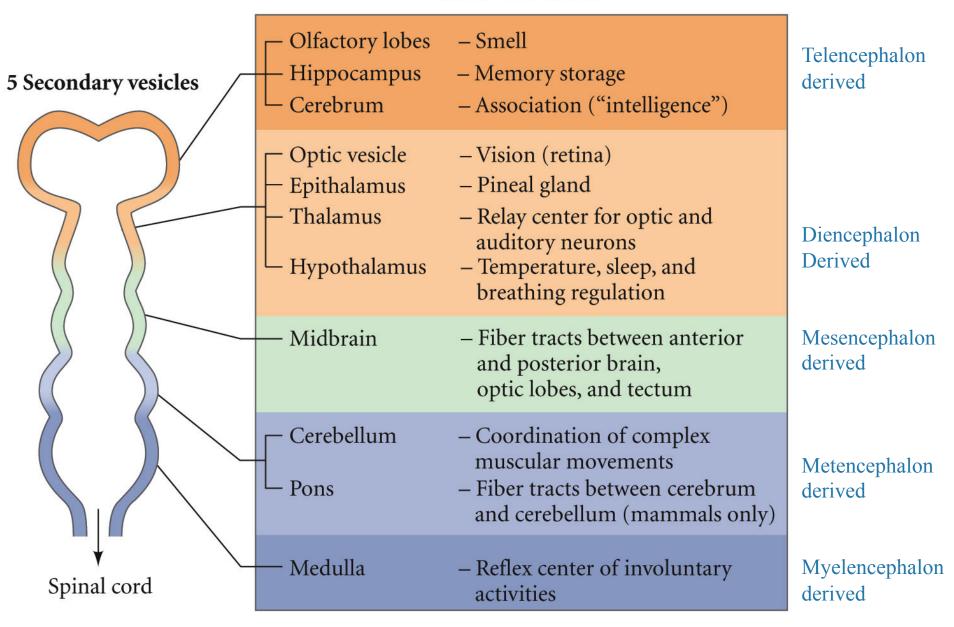
#### **Early brain development**

Proliferation to grow in thickness => Vesicule formation due to differential proliferation

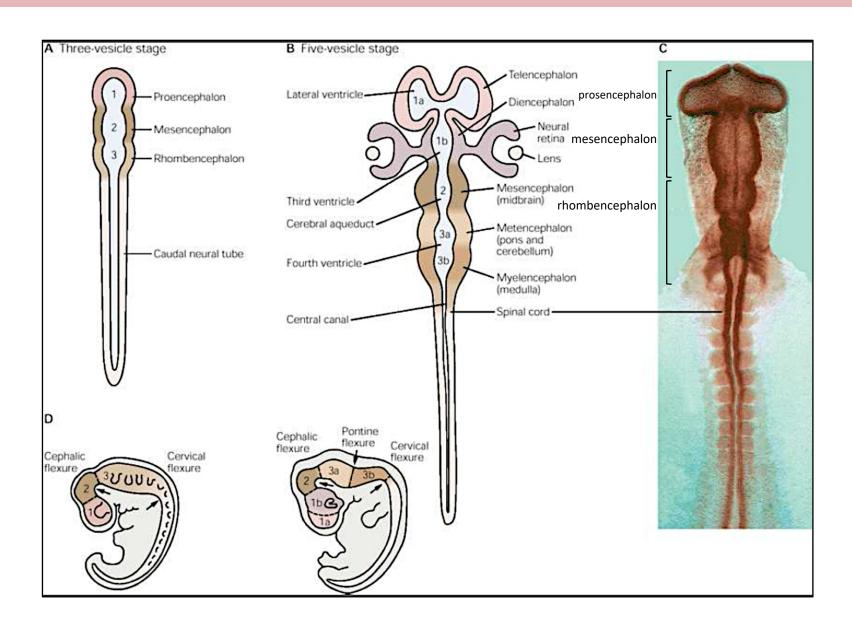


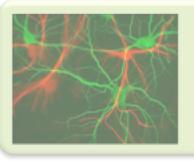
#### Early brain development

#### Adult derivatives



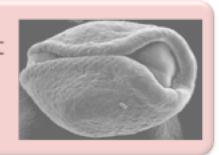
#### **Curving of the neural tube**

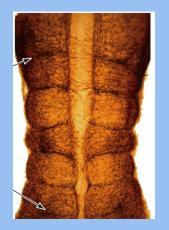




- Nervous system description
  - Nervous tissue

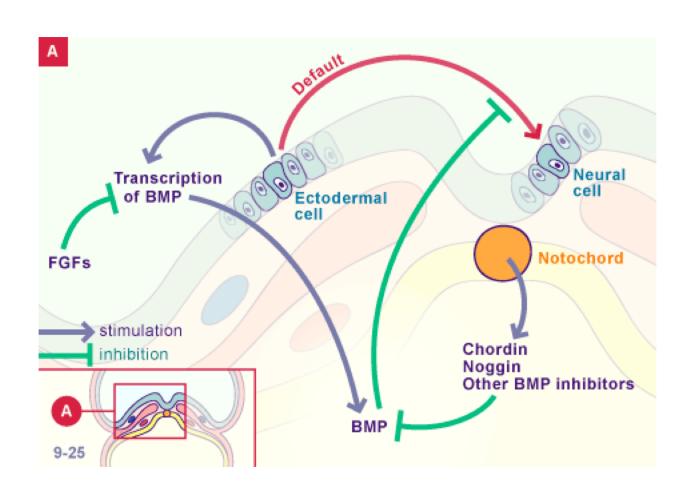
Nervous system development
Neurulation



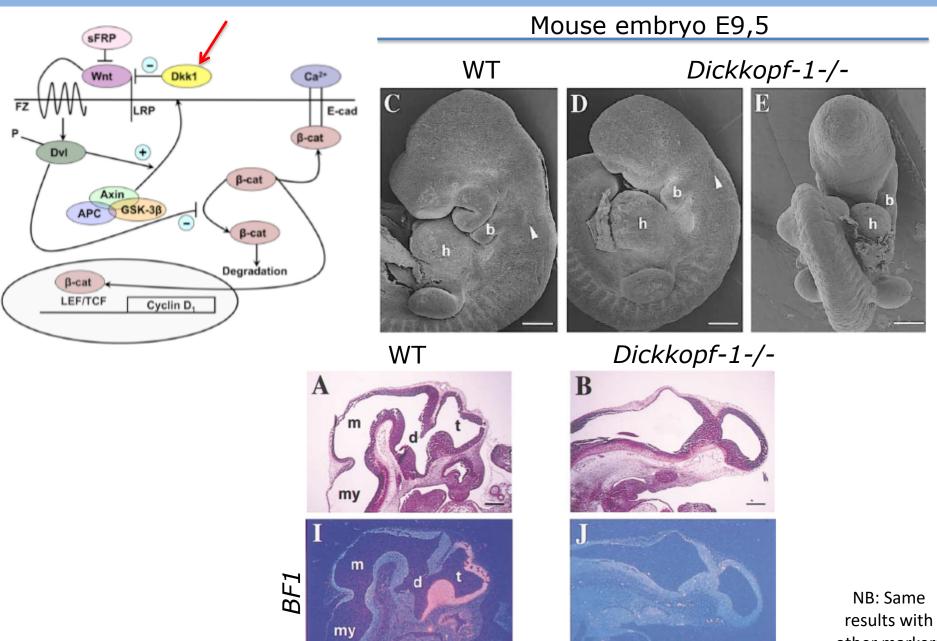


- Neural induction
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#### **Neural induction**

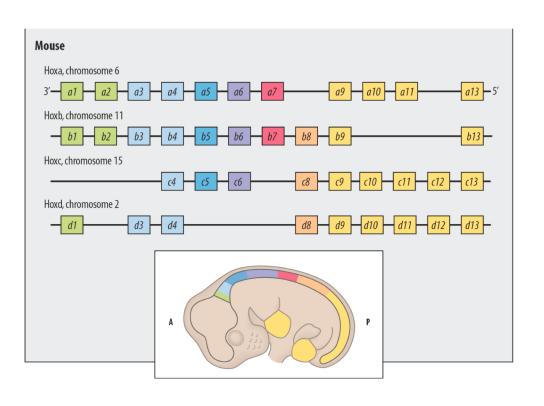


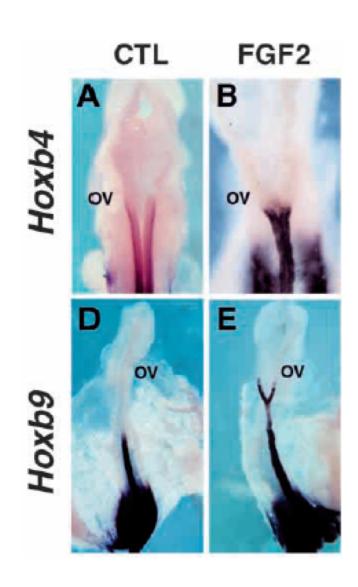
#### Wnt: posteriorization of the brain



other markers

#### Sensitivity of the neural tube to FGF

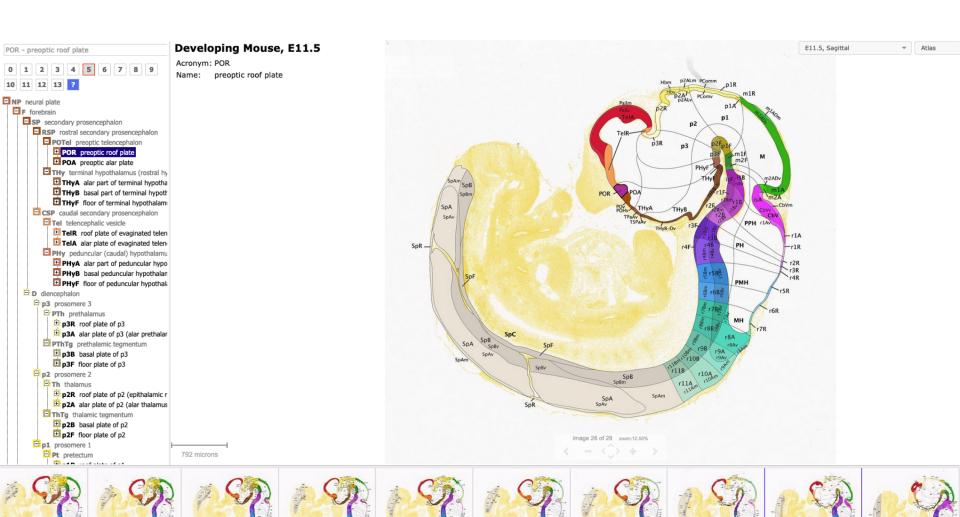




#### **Allen Brain Atlas (ABA)**

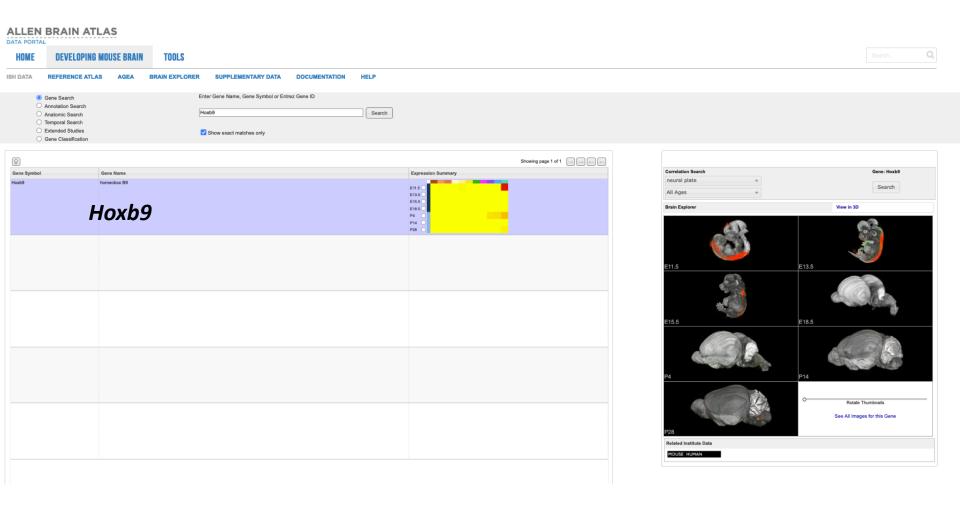
Online database with gene expression mapping in mouse and human brains

#### Allows visualization of brain structures



#### **Allen Brain Atlas (ABA)**

#### Allows visualization of specific gene expression in 2D or 3D



#### **Regionalization defects**

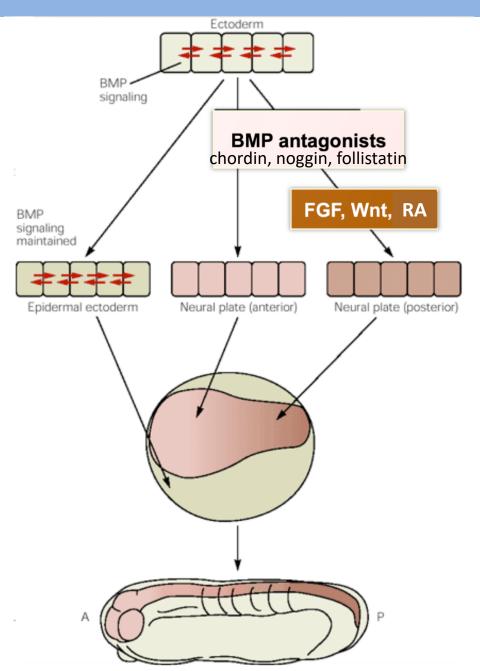
Example: Holoprosencephalias:

Anomalies during the subdivision of the prosencephalon to generate the telencephalon and the diencephalon. Moreover, the telencephalon does not divide totally into two hemispheres.

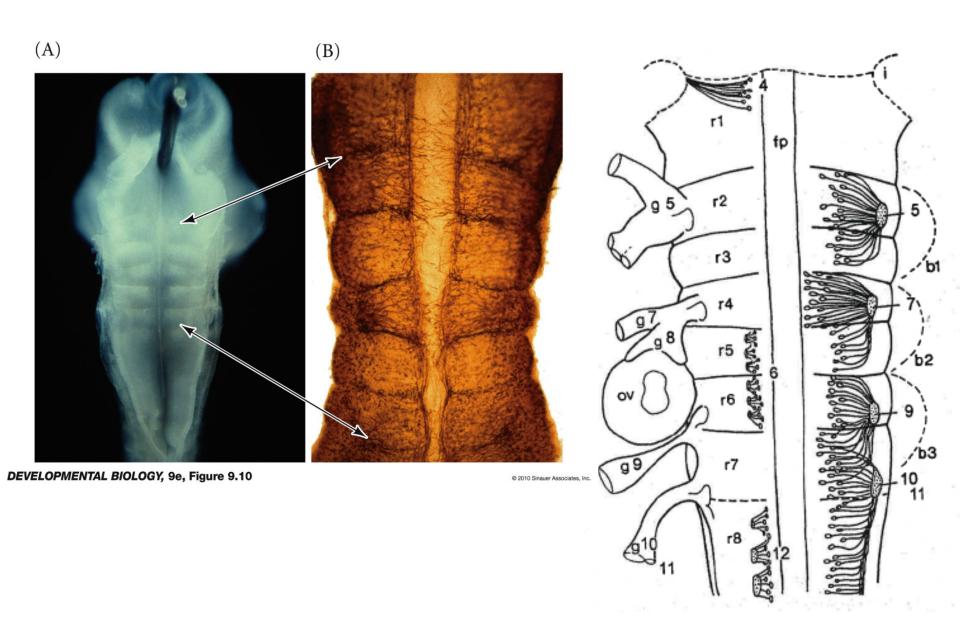
There are variable degrees of malformation.

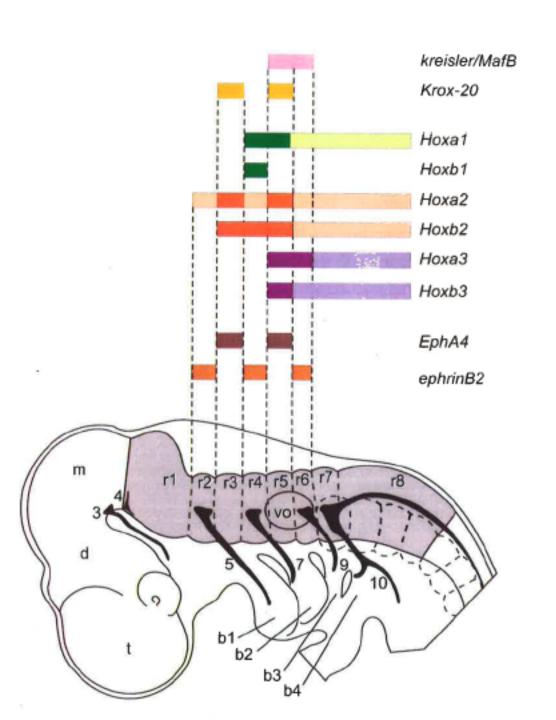


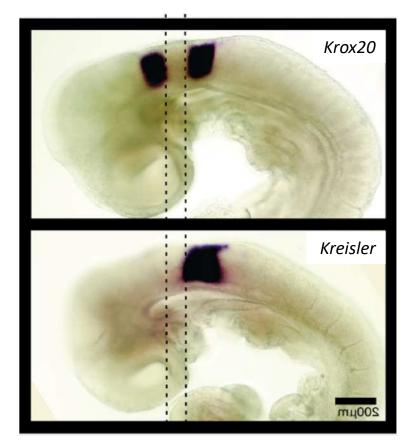
#### To sum up



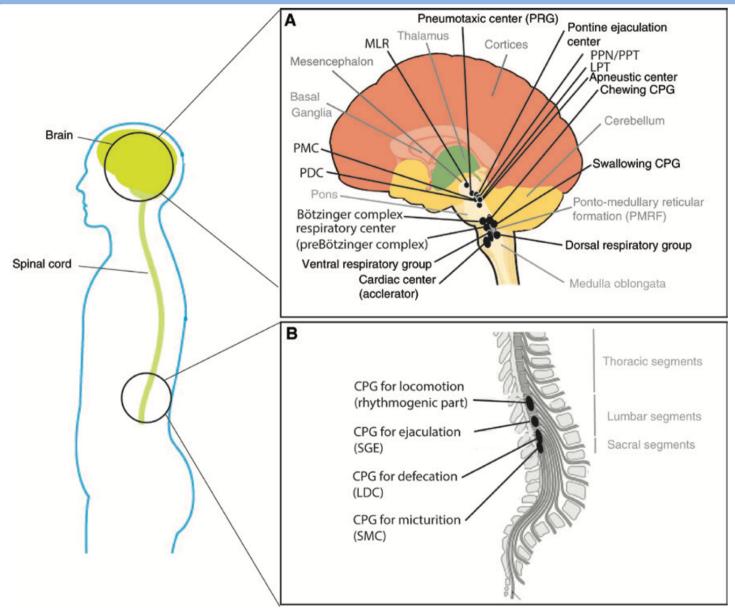
#### Hindbrain: an example of regionalisation



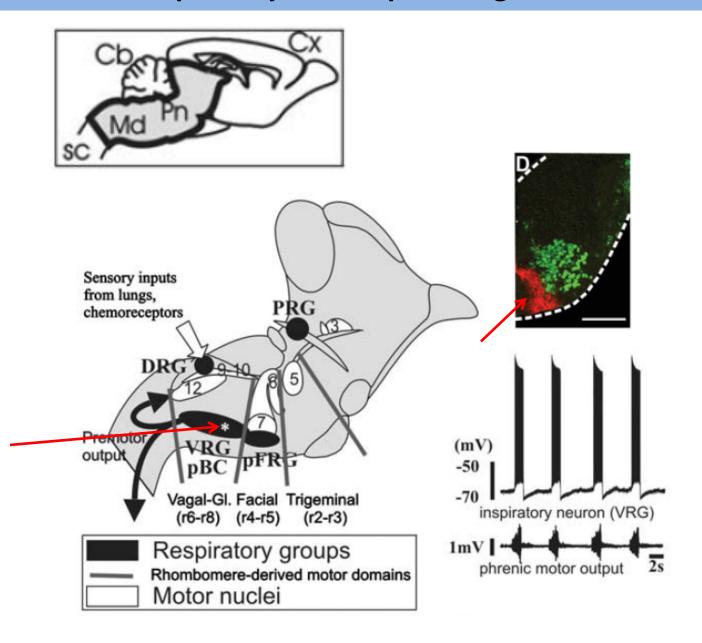




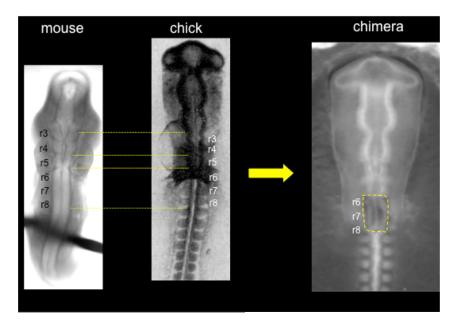
## The main central pattern generators (CPGs) in the brainstem and the spinal cord

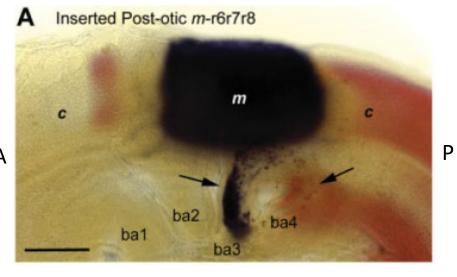


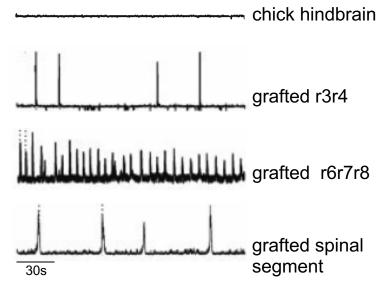
## The PreBötzinger complex: The respiratory central pattern generator



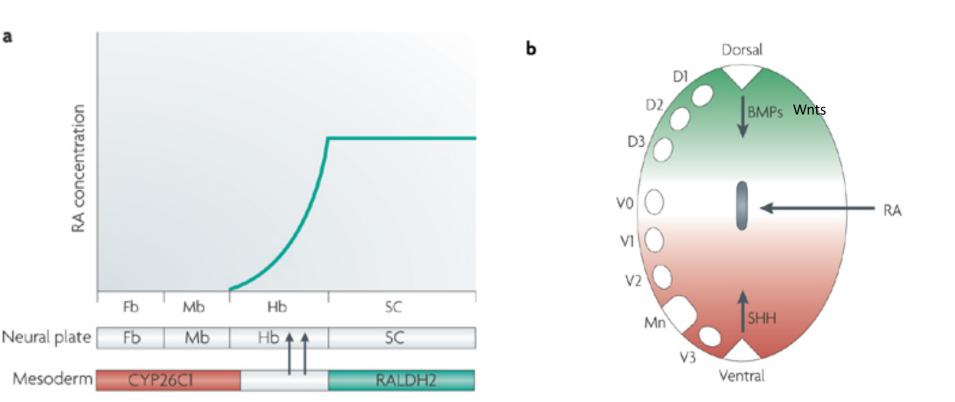
#### r6r7r8: Genetic program leading to the PreBötzinger complex



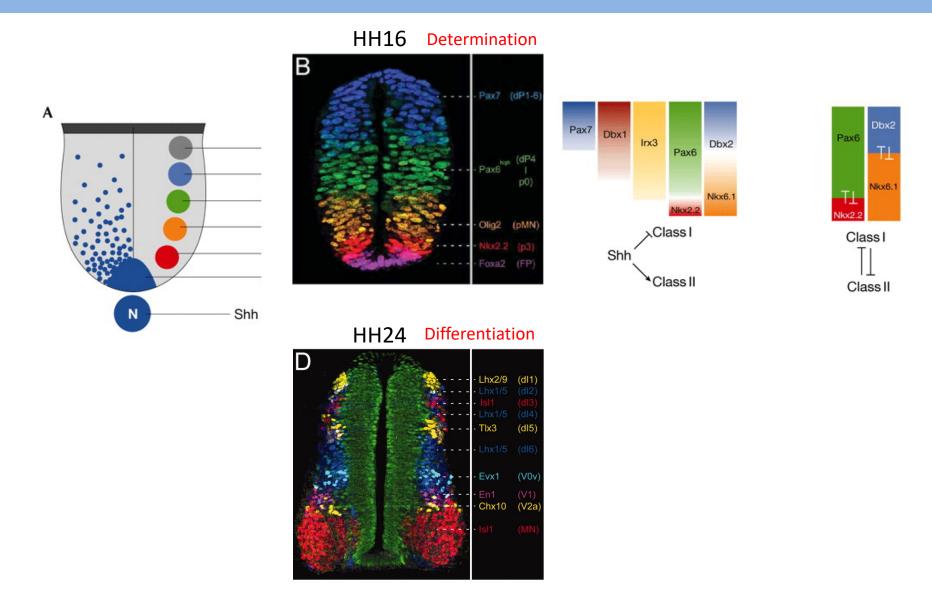




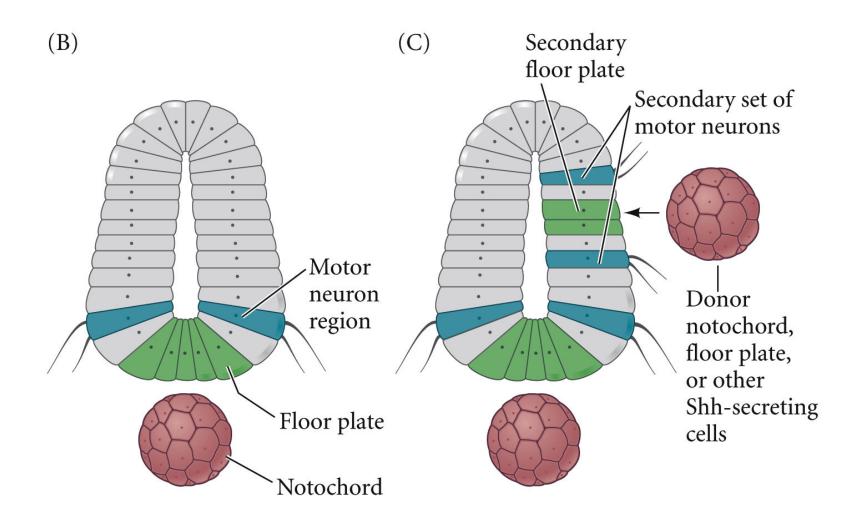
#### Control of the dorso-ventral patterning

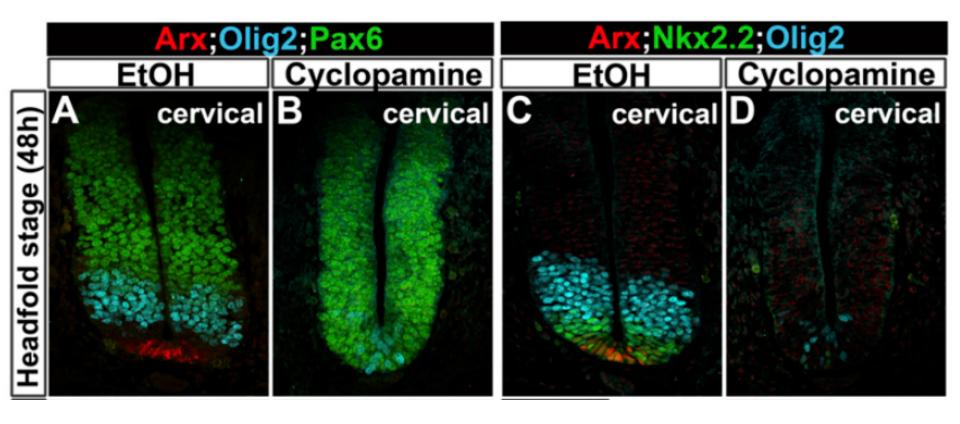


### Cascade of inductions initiated by the notochord in the ventral neural tube



Jacob and Briscoe, 2003; Le Dréau and Marti, 2012

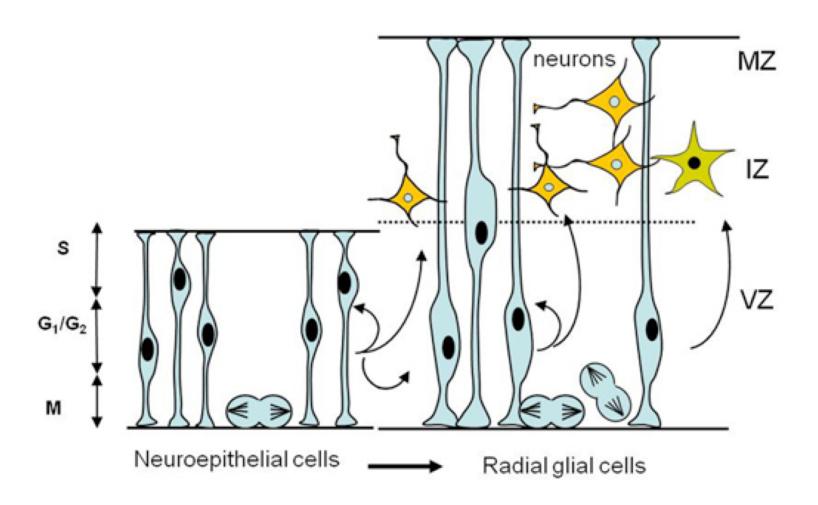






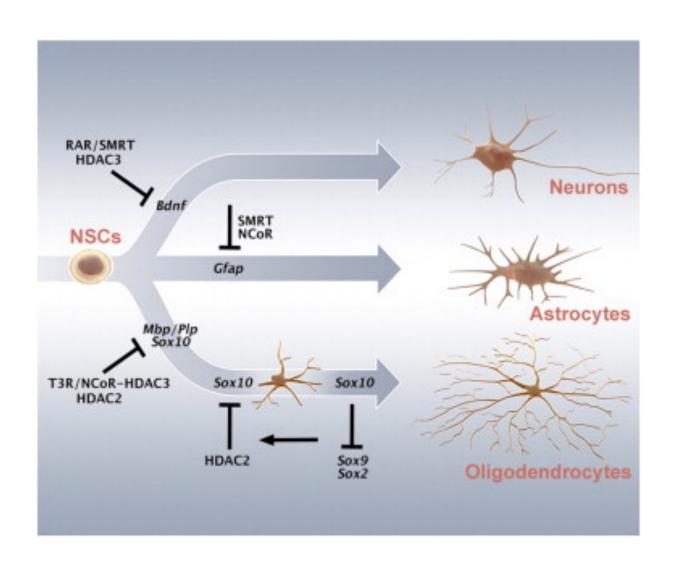
antagonizing Shh activity dorsally: a role for the Wnt canonical pathway signaling from the roof plate

#### **Proliferation and differentiation**



Neural tube / Spinal cord

#### **Proliferation and differentiation**

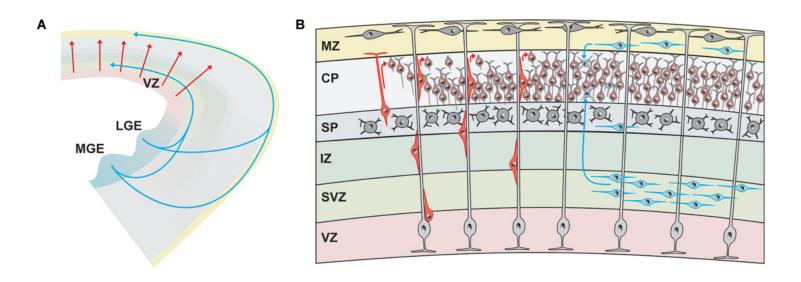


#### **Neurogenesis defects**

| Disease                         | Description                          | Symptoms   |
|---------------------------------|--------------------------------------|--|
| Megalencephaly (macroencephaly) | Brain bigger than normality          | The size of the head is increased                                  |
| Hemimegalencephaly              | One hemisphere bigger than normality | Intelectual disorders, epilepsy, paralysis on one side of the body |
| Microencephaly                  | Brain smaller than normality         | Intelectual disorders  |

#### **Neuronal migration**

Example: migration of glutamatergic and GABAergic neurons in the telencephalon



- > Intrinsic cues: Transcription factors
- Chemical cues: semaphorines and ephrins (attracting or repelling signals)
- > chemico-mechanical guiding structures: vertical fibers of radial glial cells
- > extracellular matrix protein

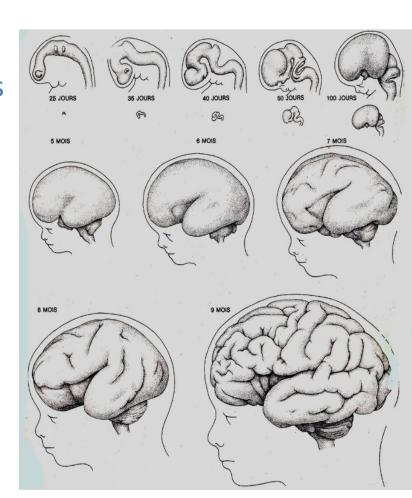
#### Migration and gyration defects

⇒ malformations of the cerebral cortex, which is not organized in six layers, as it should do.

Variable degrees: conserved organization in layers or architecture is totally disorganized

=> Gyrus are reduced compared to normal, or totally absent.

Schizencephaly, heterotopia, polymicrogyria, lyssencephaly

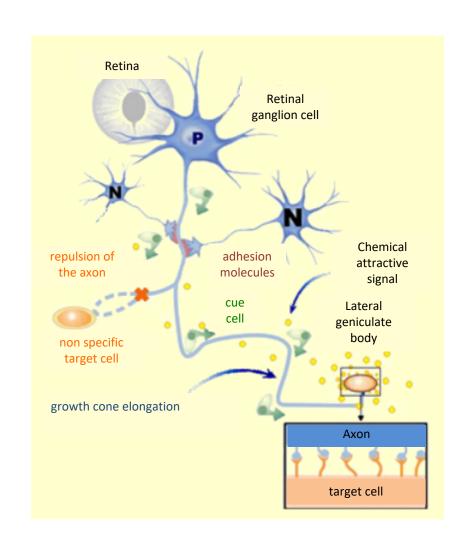


#### **Synaptogenesis**

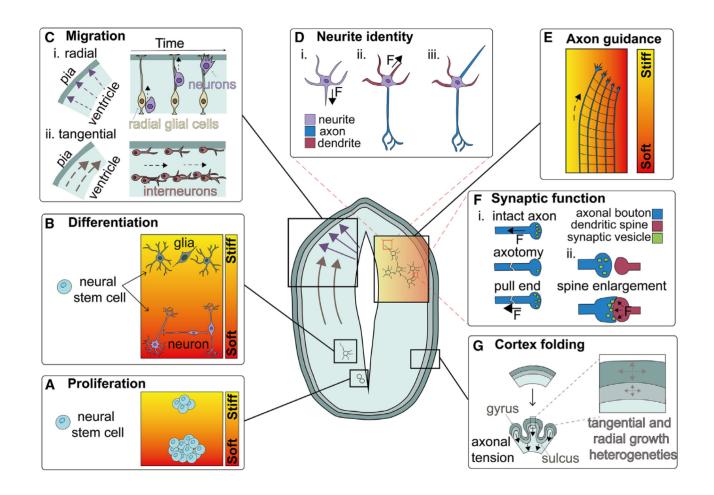
- Selective recognition of the right way and the right targets by the axon growth cone
- Formation of the synapses

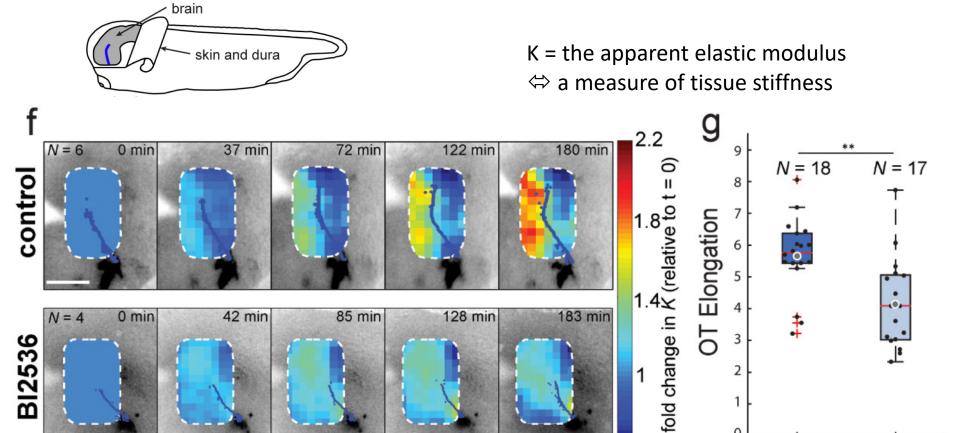
#### Then adjustments:

- Neuronal apoptosis
- Synaptic pruning: reduction in the number of synaptic connections => conservation of more efficient synaptic configurations



#### **Mechanical regulation**





BI2536: mitotic blocker

Development of a stiffness gradient in the Xenopus embryo brain

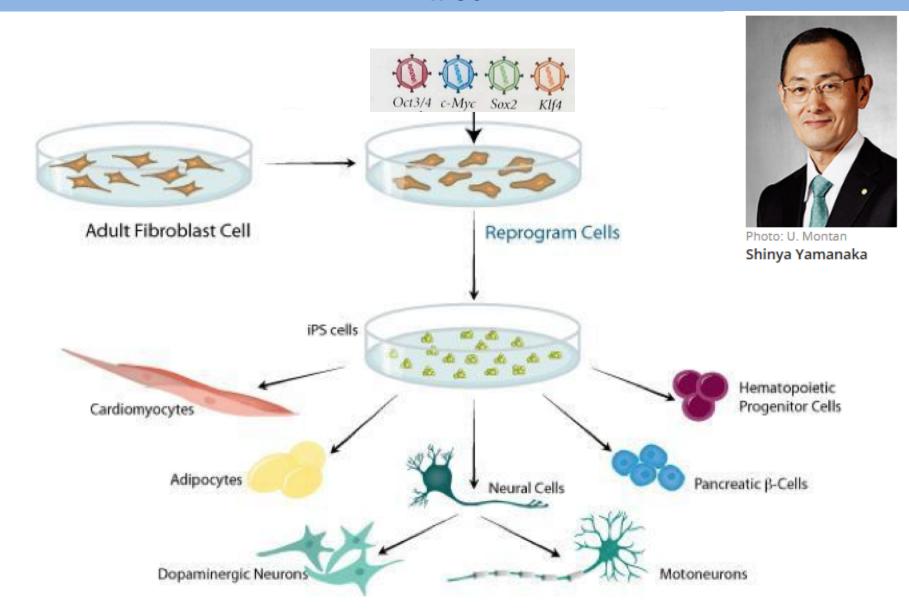
- precedes axon turning of ganglion cells
- is necessary for its correct elongation.

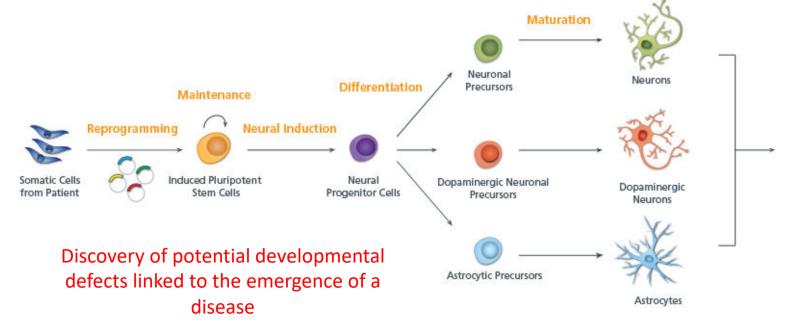
control

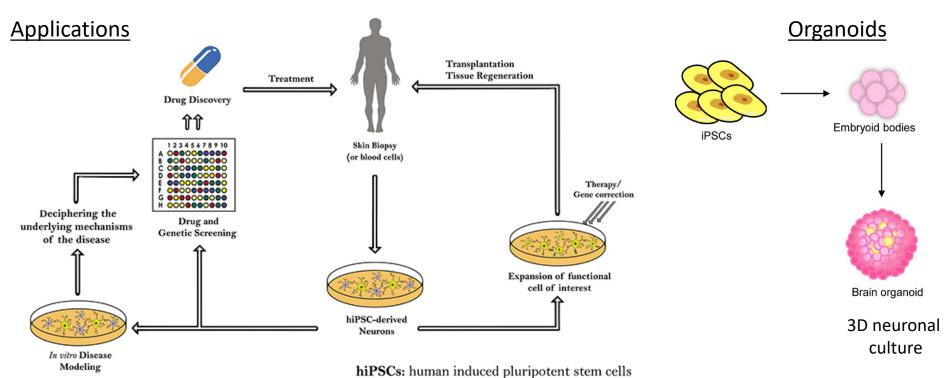
BI2536

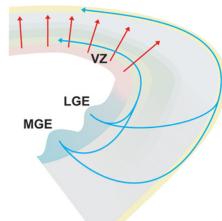
#### **Development and therapy**

#### => IPSc



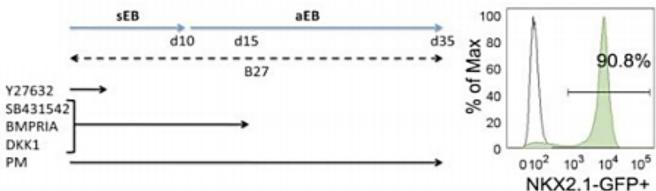






#### Inhibitory GABAergiques neurons (Nkx2.1+) are involved in autism

⇒ generation of these neurons from IPSc



BMP inhibition: SB431542 et BMPRIA

Wnt inhibition: DKK-1 (Dickkopf-1)

Shh activation: PM (purmorphamine)

(Y-27632 for iPS survey)

#### REVIEW Open Access

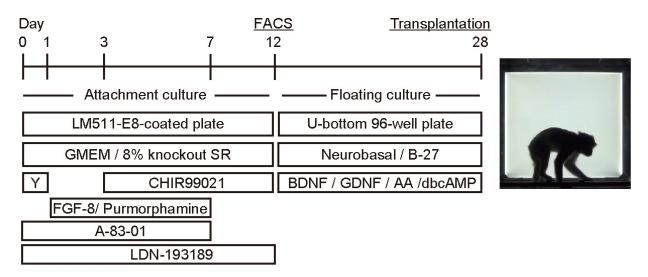
# Exploring the mechanisms underlying excitation/inhibition imbalance in human iPSC-derived models of ASD

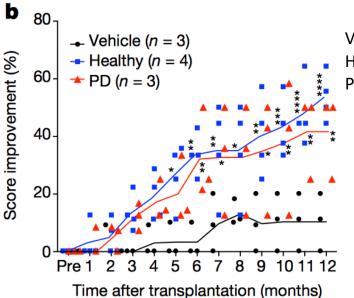


Lorenza Culotta<sup>1,3</sup> and Peter Penzes<sup>1,2,3\*</sup>

| Gene                                   | Model type  | Phenotype observed   | Reference                  |
|--|---|--|----------------------------|
| ATRX, AFF2, KCNQ2,<br>SCN2A, and ASTN2 | Homozygous deletion   | Reduced synaptic activity  | [ <u>53</u> ]              |
| CACNA1C                                | ASD-related mutations                                       | Disrupted interneurons migration   | [ <u>54]</u>               |
| CNTN5 and EHMT2+                       | Heterozygous deletion                                       | Hyperexcitability.   | [ <u>55</u> ]              |
| CNTNAP2                                | Heterozygous deletion                                       | Increased neuronal network activity  | [ <u>56</u> ]              |
| FMR1                                   | Heterozygous deletion                                       | Impaired retinoic acid (RA)-dependent homeostatic synaptic plasticity  | [ <u>57</u> ]              |
| MECP2                                  | Heterozygous deletion or duplication                        | Altered synaptic density, altered calcium signaling; altered neuronal firing rate and synchronization; delayed GABA switch                                       | [58,59,60,61]              |
| NLGN4                                  | Gene overexpression and ASD-related mutations               | Increased excitatory synapse density, altered synaptic strength  | [ <u>62</u> , <u>63</u> ]  |
| NRXN1a                                 | Homozygous and heterozygous deletion, ASD-related mutations | Impaired synaptic strength, altered synaptic calcium signaling   | [ <u>64,65,66</u> ]        |
| SHANK2                                 | Heterozygous deletion and ASD-related mutations             | Hyperconnectivity, enhanced branching complexity, increased synapse density  | [ <u>67</u> ]              |
| SHANK3                                 | Heterozygous deletion and ASD-related mutations             | Hypoconnectivity, reduced synaptogenesis, and dendritic arborization; impaired neuronal excitability and excitatory synaptic transmission; impaired HCN channels | [68,69,70,71,72,7<br>3,74] |
| TSC1/2                                 | Homozygous and heterozygous deletion                        | Altered neuronal excitability and activity, altered synchrony (cortical neurons); hypoexcitability (cerebellar Purkinje cells)                                   | [75,76,77,78]              |
| Other ASD models                       |   | Aberrant neuronal maturation, altered neuronal differentiation and synaptic formation  | [ <u>79</u> , <u>80</u> ]  |

## Parkinson's disease: degeneration of midbrain dopaminergic neurons ⇒ Human iPS cell-derived dopaminergic neurons function in a primate Parkinson's disease model





Vehicle: monkeys that received control injection

Healthy: monkeys transplanted with cells derived from healthy individuals

PD: monkeys transplanted with cells derived from PD patients

