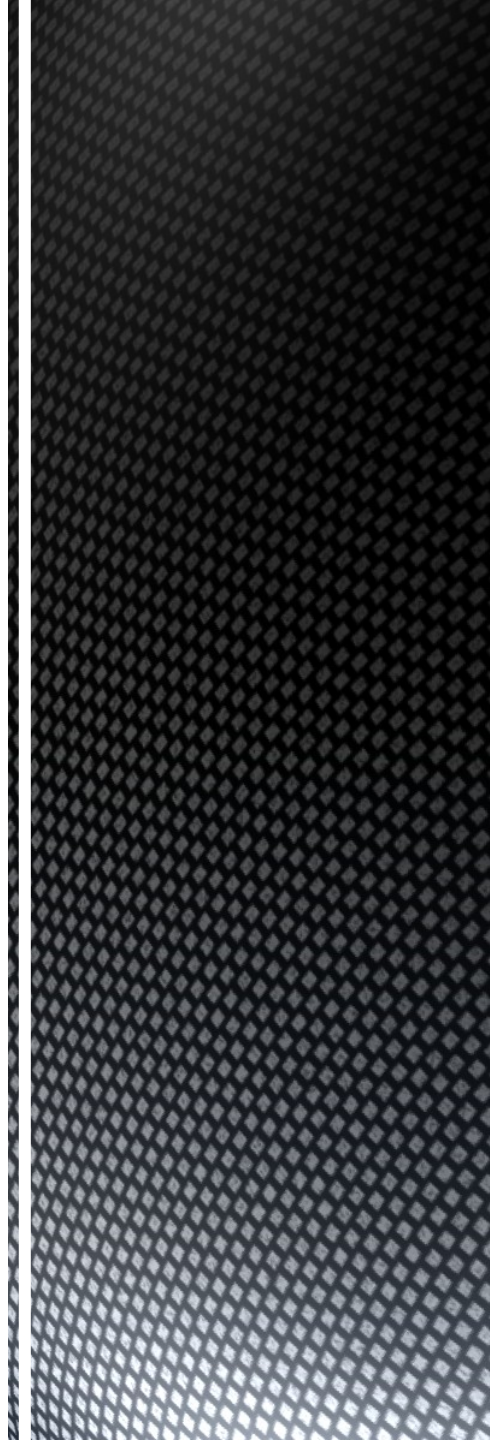


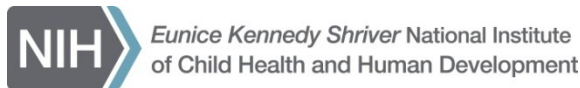
Promoting Healthy Socioemotional Development: A Neuropsychologically Informed Approach

Erin B. Tone. Ph.D.



- Conflicts of interest: None
- Research funding:
 - National Institute of Child Health and Development (NICHD)
 - Anxiety and Depression Association of America (ADAA)
 - Georgia State University Brains and Behavior Program

Conflicts of Interest and Funding Acknowledgement



- How do emotion and emotion regulation skills develop?
- How can the dynamic interplay among intrapersonal and interpersonal characteristics lead to deviations from typical developmental trajectories?
- How can we effectively target the emotion dysregulation that can emerge in atypical emotional development?



- We begin early to experience and express emotion



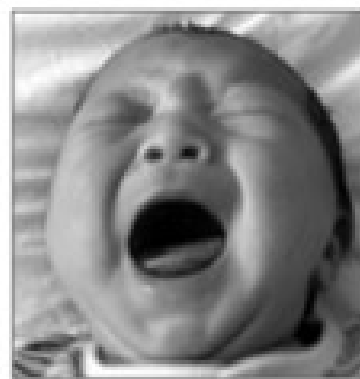
- We do so in ways that others can accurately read and interpret . . .



Happy



Sad



Furious



Terrified

Yilin's Emotion Chart



Happy



Sad



Petulant



Lonely



Amused



Skeptical



Furious



Wistful



Confused



Bored



Sarcastic



Regretful



Aroused



Terrified



Proud



Mischievous

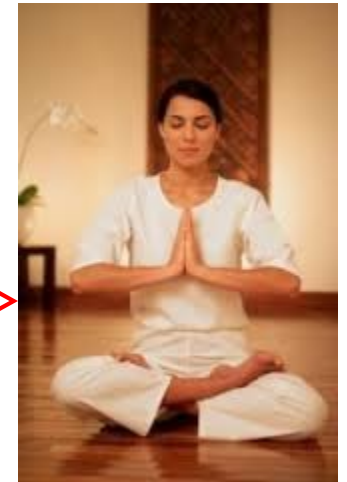
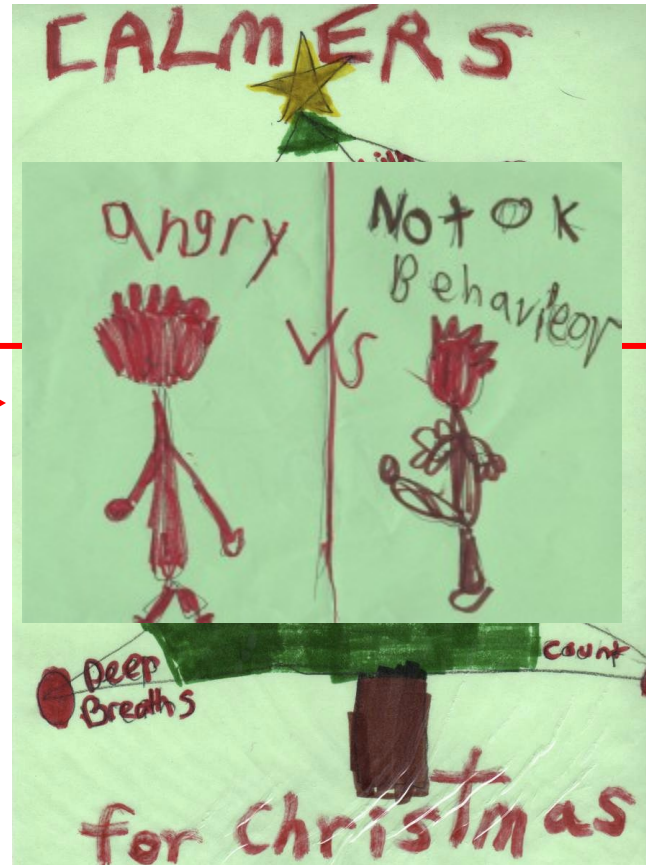
and we do so in an
incredibly nuanced
way . . .

We also begin early to read and respond to others' emotional cues . . .



- <http://www.youtube.com/watch?v=apzXGEbZht0>

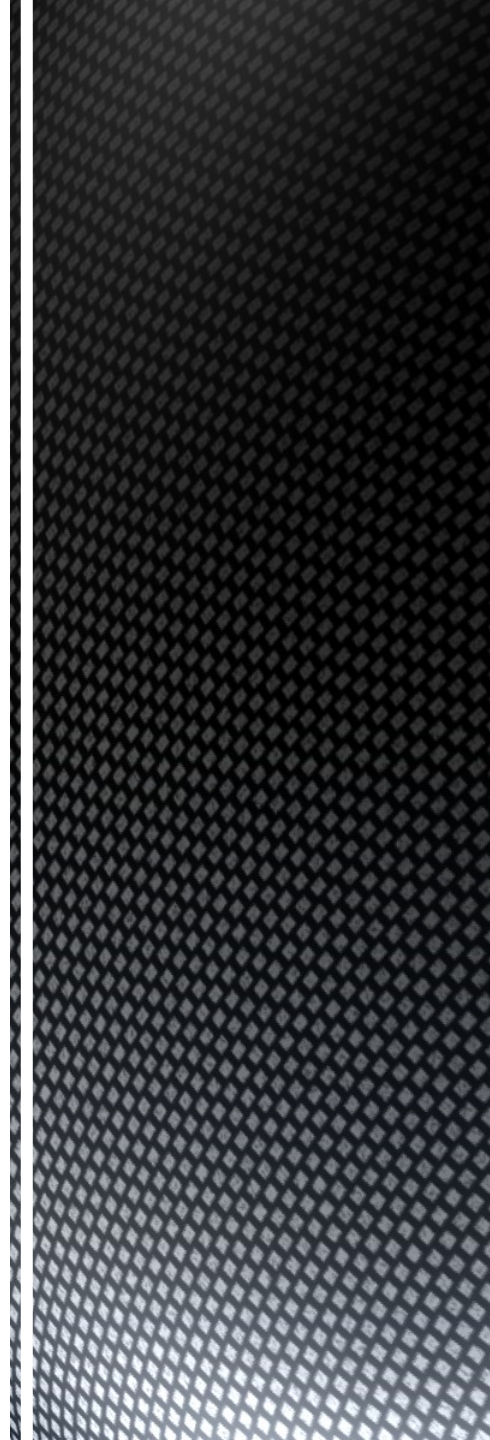
and to regulate our emotions . . . initially with ample support, increasingly on our own



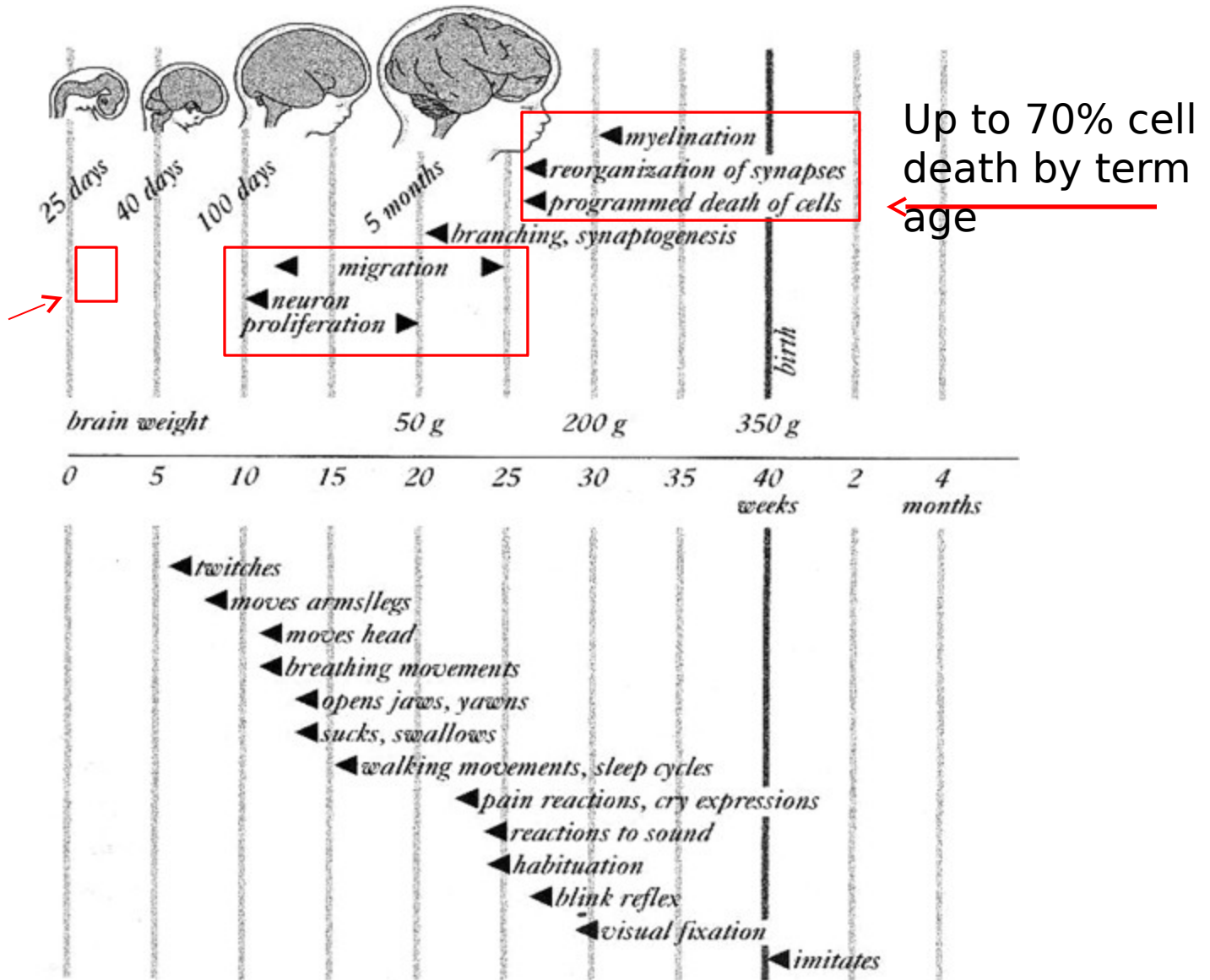
**How do emotion and
emotion regulation skills
develop?**

**What happens in the brain that
allows us to acquire these
competencies?**

Prenatal development

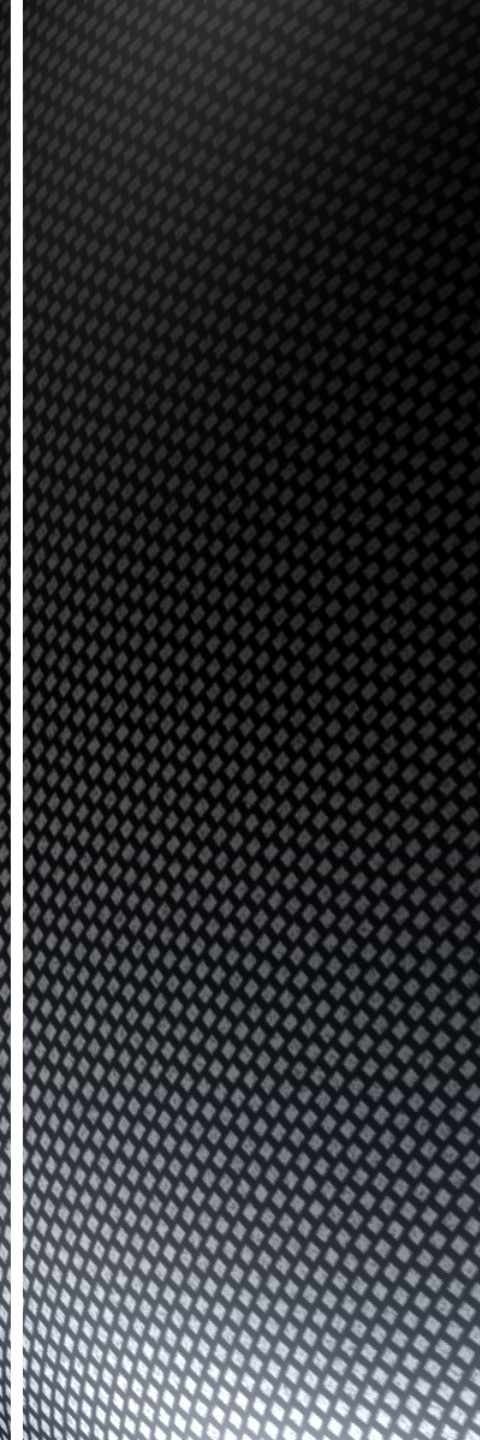


Predecessor neurons migrate to prospective human cortex (~30 days gestation)

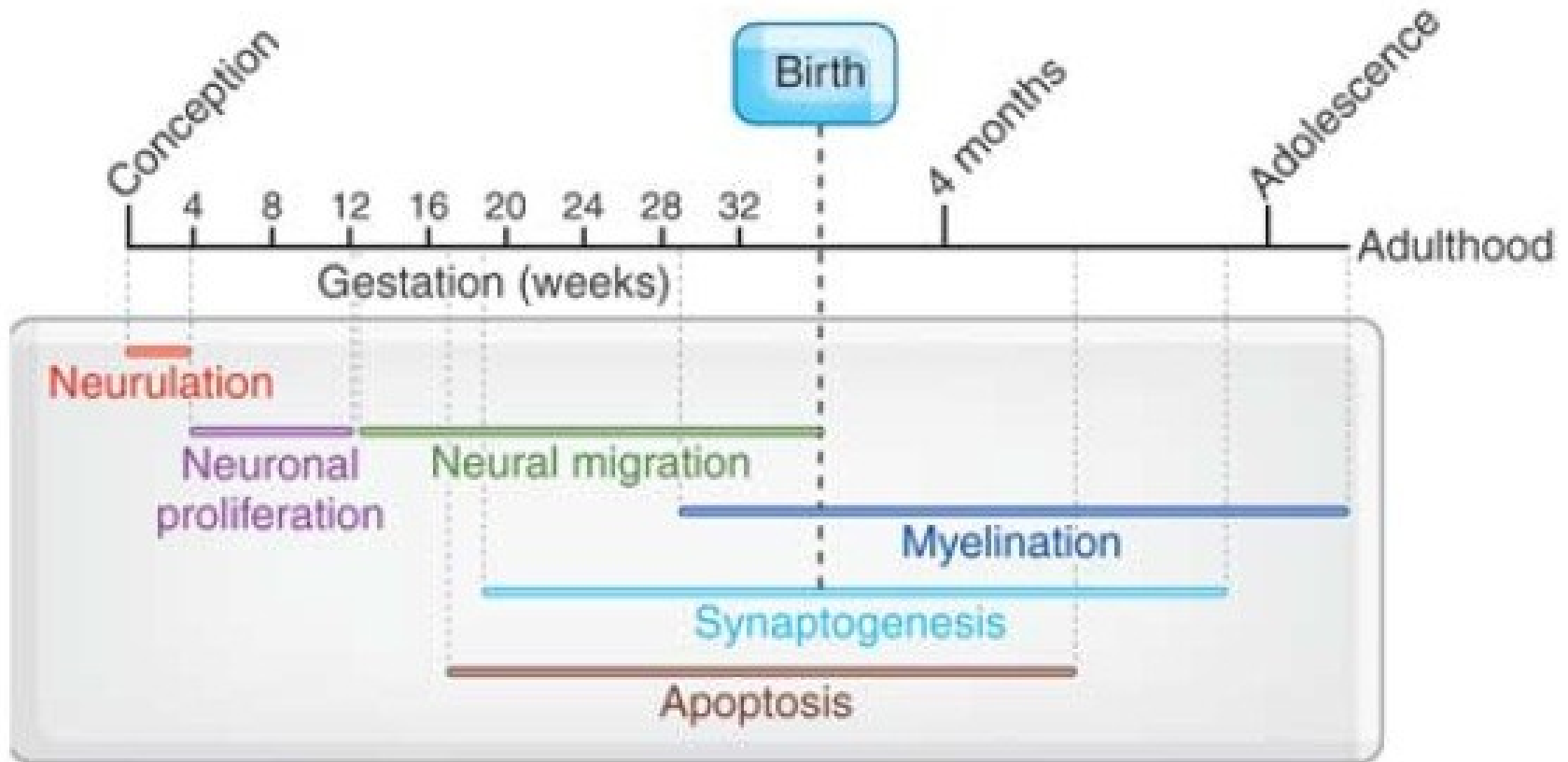


The development of the brain displayed in a schematic form.
 Source: The Newborn Brain. Edited by Lagercrantz H, Hanson M, Evrard P, Rodeck C. Cambridge University Press, Cambridge, UK. 2002, p. xii

Development in
Infancy,
Childhood, &
Adolescence



Cellular Level Development



Tau GZ, Peterson BS (2010). "Normal Development of Brain Circuits". *Neuropsychopharmacology* **35** (1): 147-168. doi:10.1038/npp.2009.115. PMC 3055433.PMID 19794405.

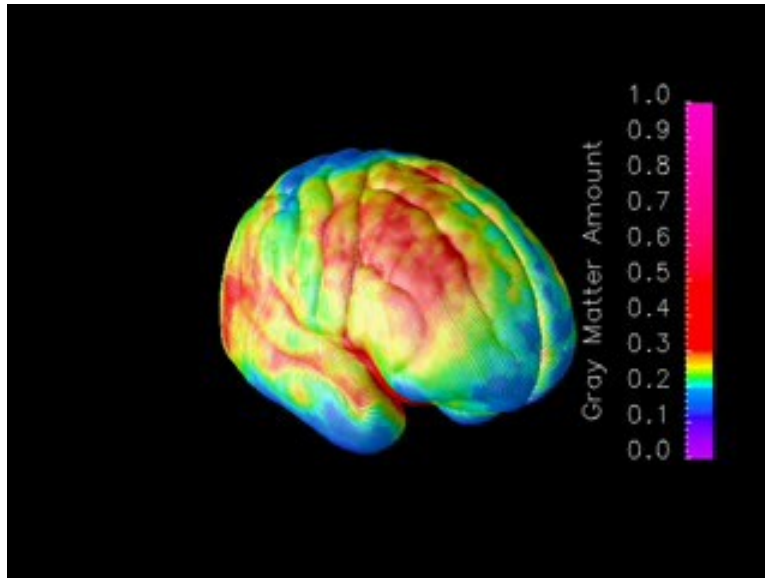
Structural Level Development: Gray matter

Big Picture:

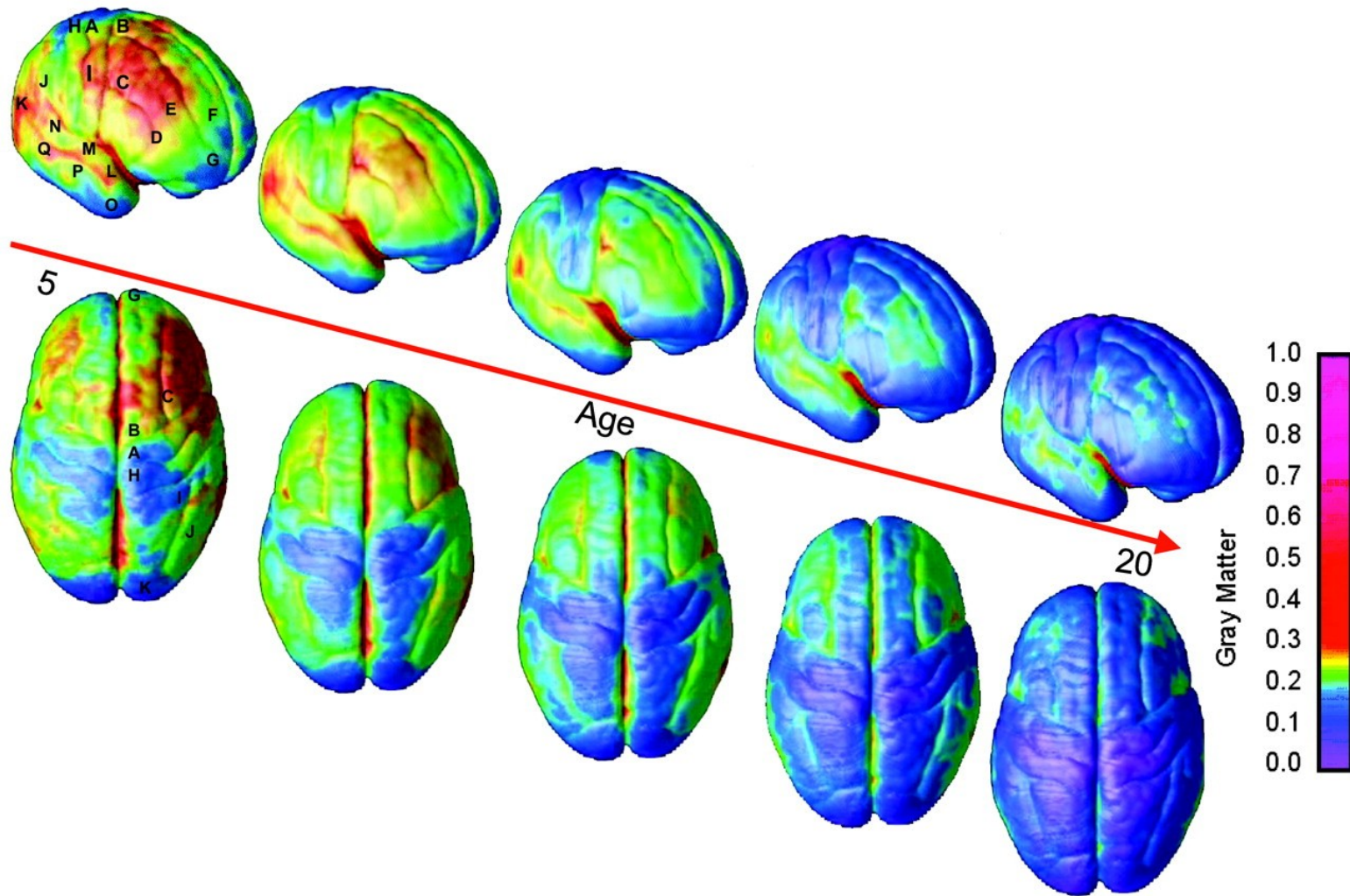
Higher-order association cortices mature after somatosensory and visual cortices

Phylogenetically older structures mature earlier than do newer ones

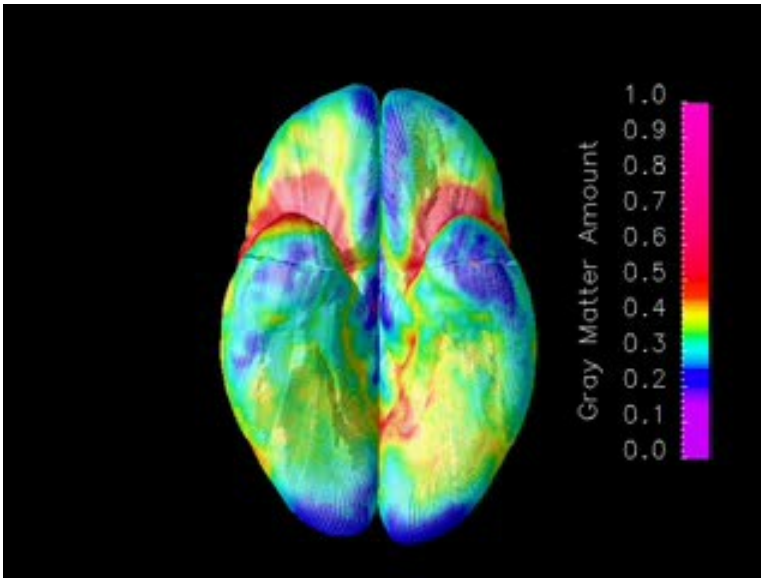
Around puberty, sustained GM loss begins



Right lateral and top views of GM maturation over the cortical surface
(n=13; each scanned 3-4 times at 2 year intervals)



Gogtay N et al. PNAS 2004;101:8174-8179

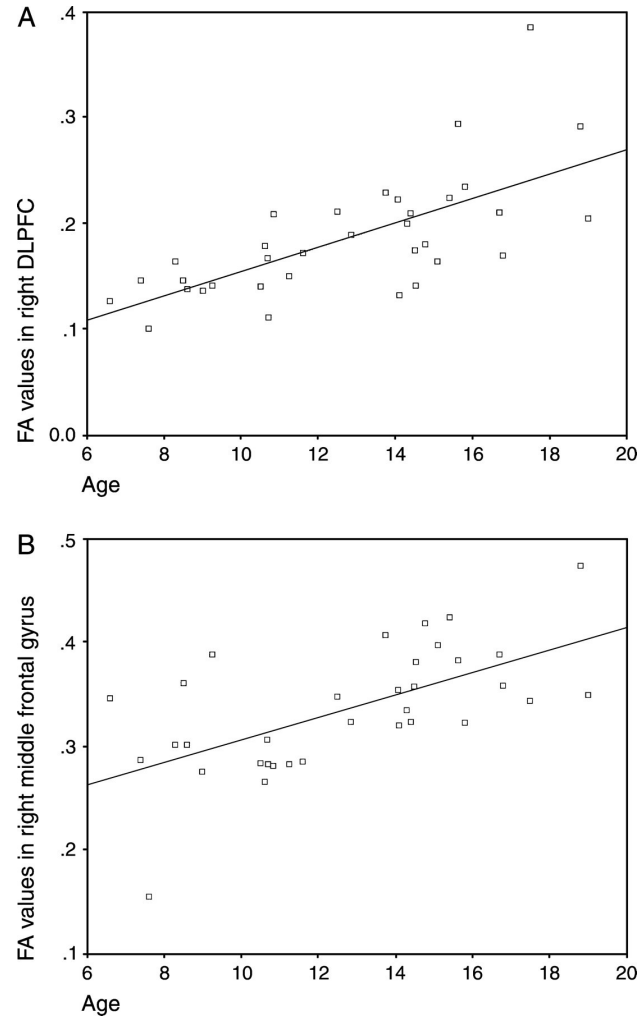


Medial inferior temporal & caudal/medial inferior frontal regions mature early, with little change thereafter

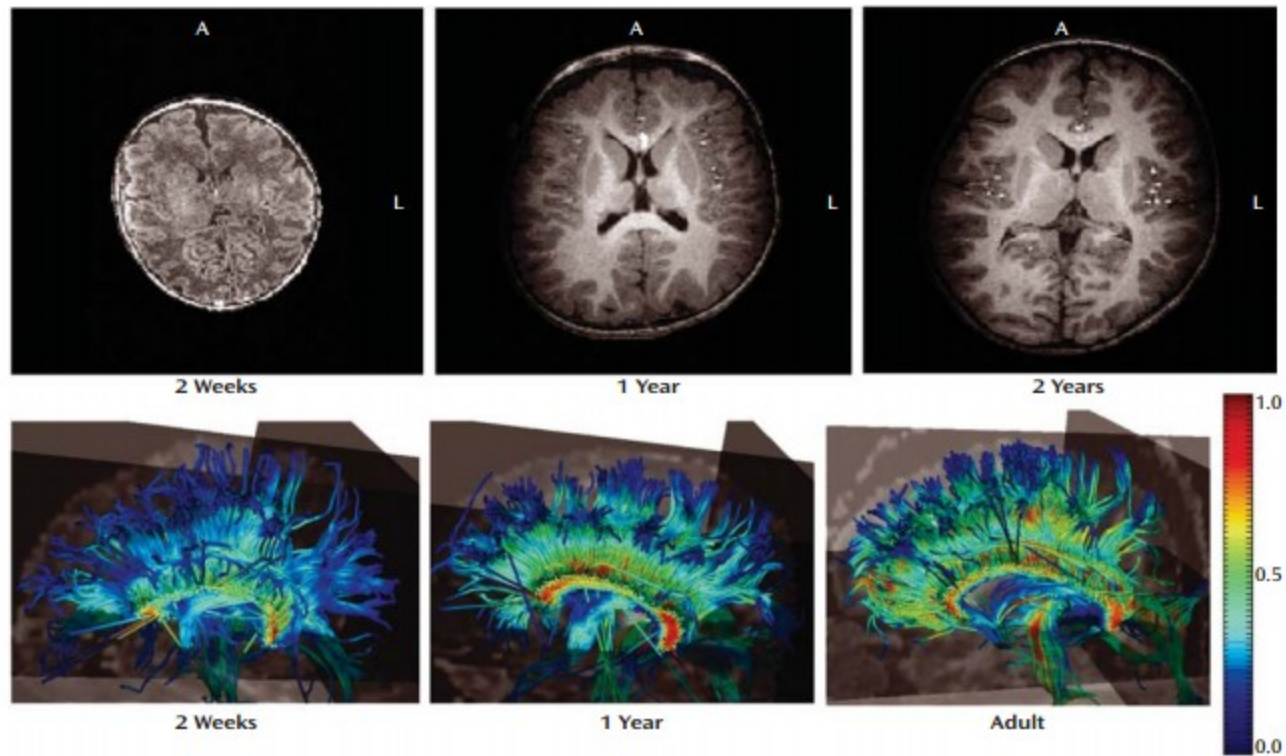
Orbitofrontal regions continue to mature throughout adolescence

Structural level Development: White Matter

Non-cortical white matter circuitry becomes more coherent, or more myelinated, with age



Barnea-Goraly N et al. *Cereb. Cortex* 2005;15:1848-1854

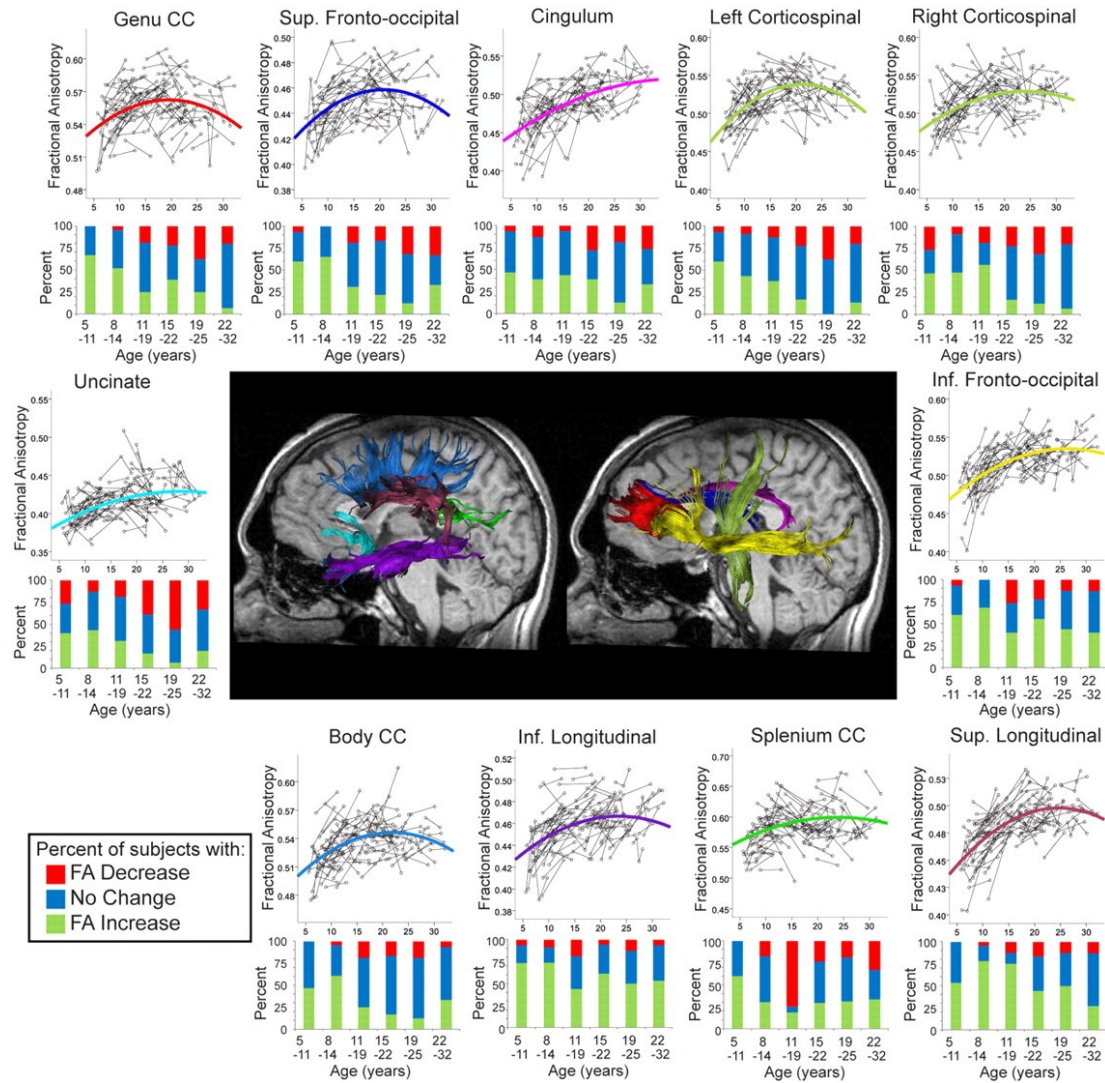


Brain myelination across development. Top panels: Images show age-related increase in brain size and white matter intensity acquired longitudinally from 1 child.

Bottom panels: Age-related differences in the organization of corpus callosum white matter (higher values = greater organization of fiber tracts)

Gilmore JH, Lin W, Gerig G. Fetal and neonatal brain development. *Am J Psychiatry*. 2006;163:2046.

Longitudinal age-related changes of fractional anisotropy.

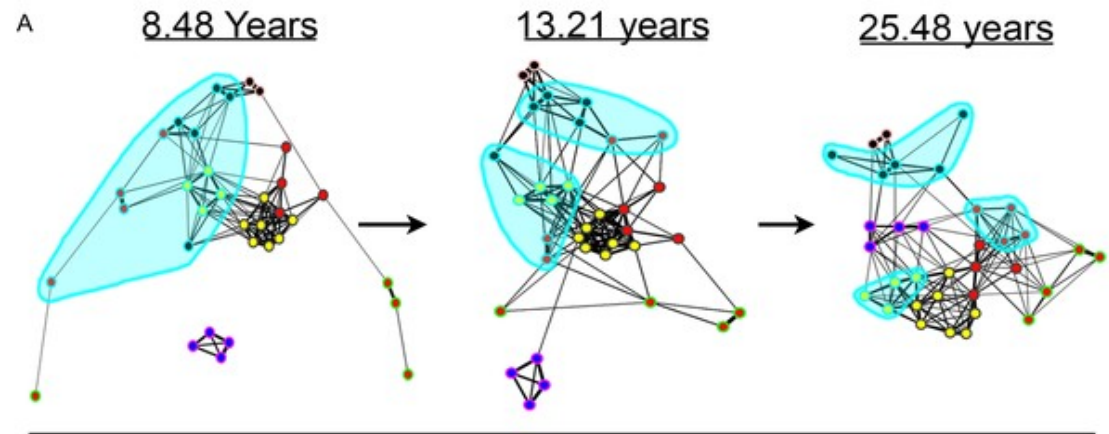


Lebel C, and Beaulieu C. *J. Neurosci.* 2011;31:10937-10947

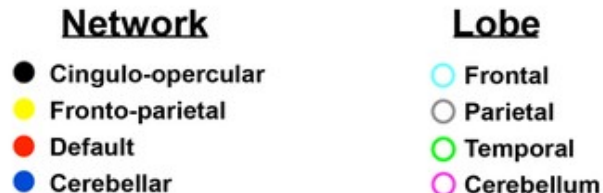
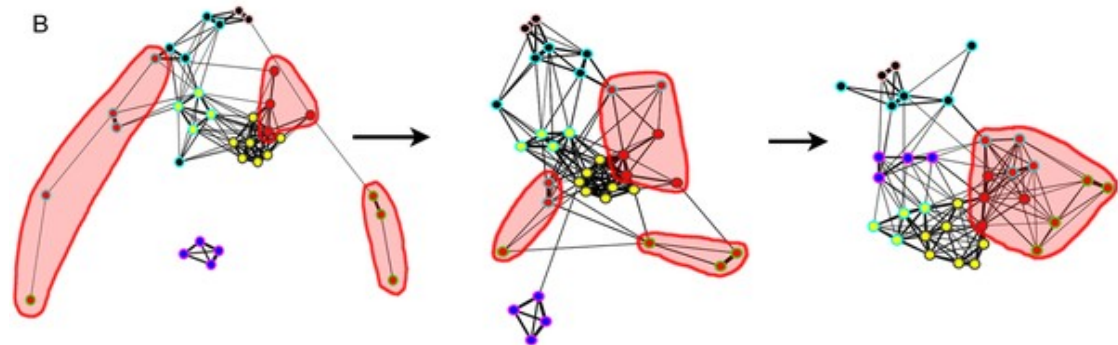
n=103

Figure 2. Over age the graph architecture matures from a “local” organization to a “distributed” organization.

A:
Anatomically-clustered regions segregate with age.

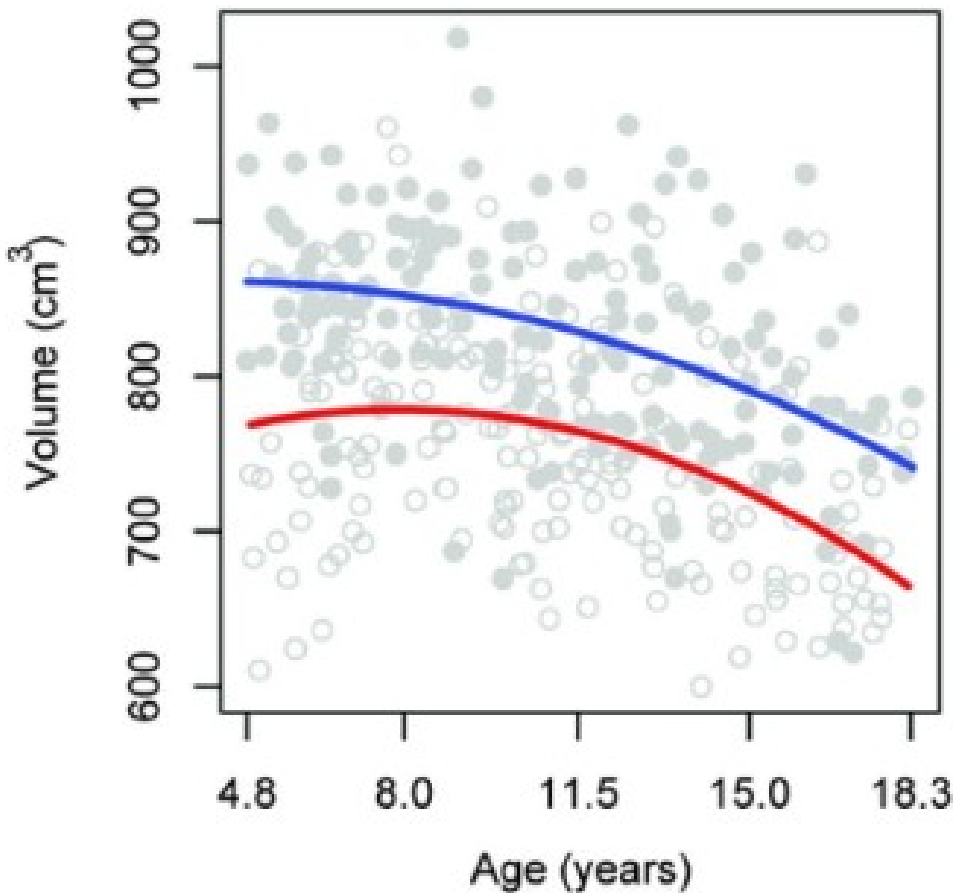


B. Functionally clustered networks become more integrated with age.

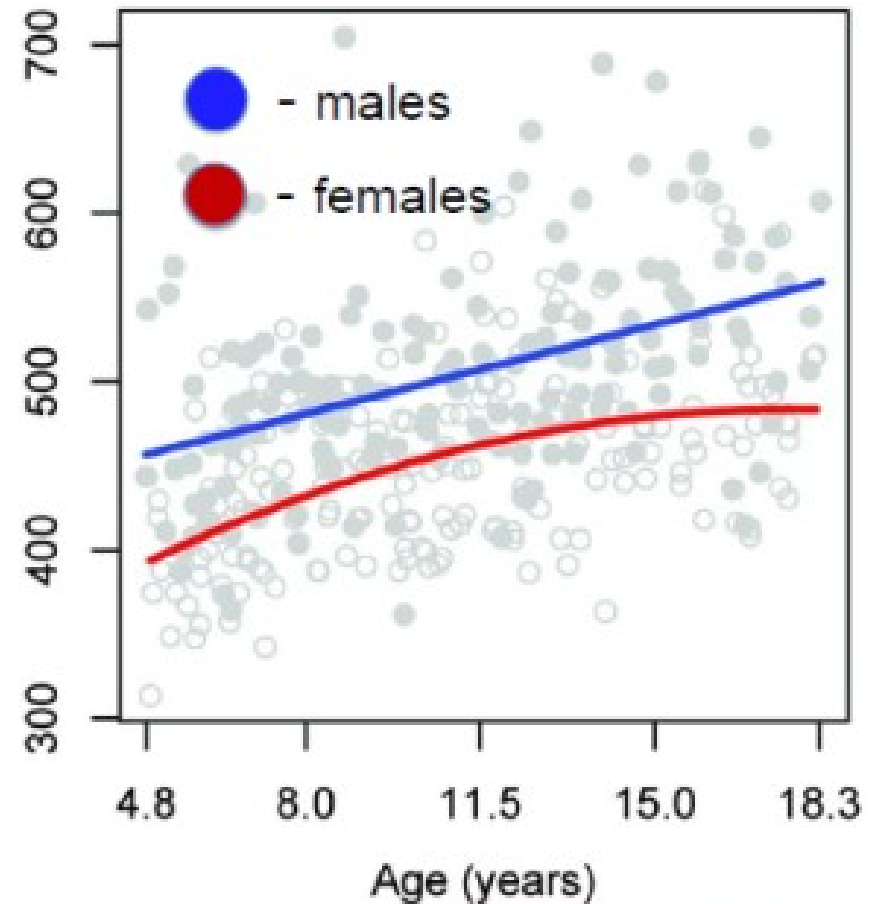


Changes Occur Within and Between Circuits

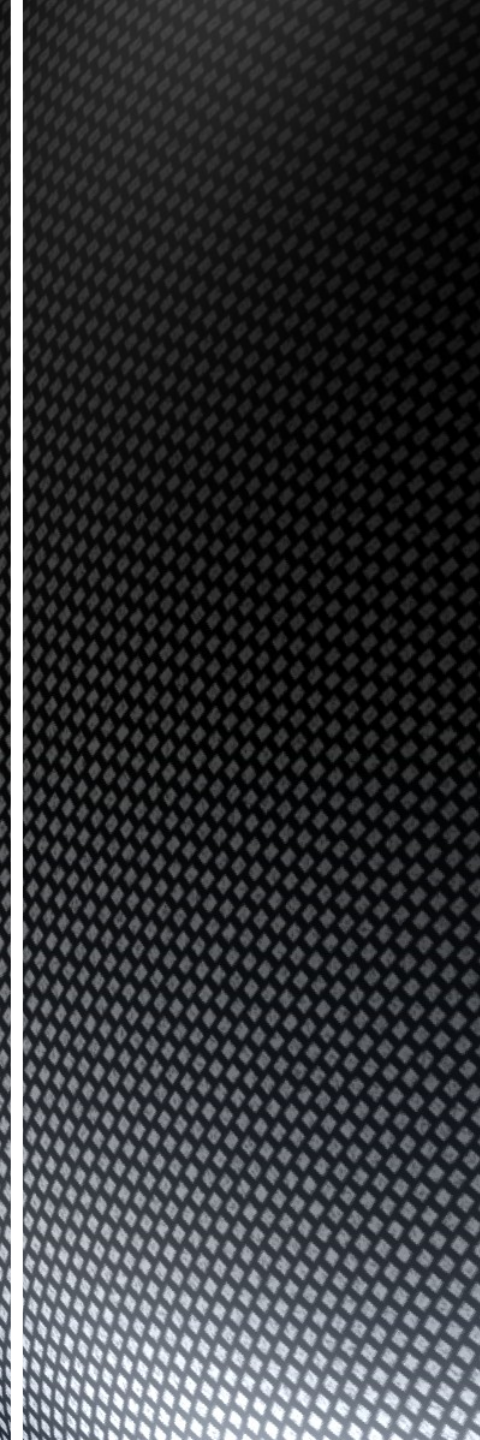
Gray Matter

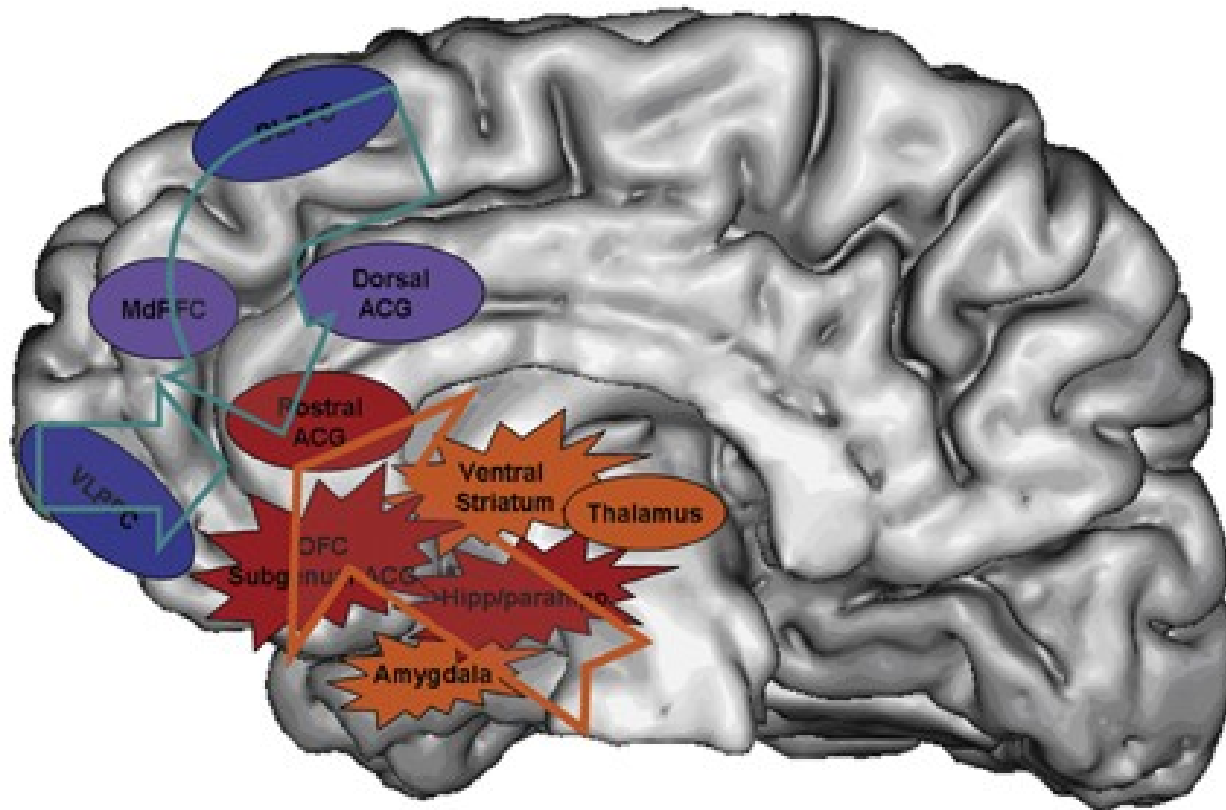


White Matter



What does development look like in neural regions that support emotional experience and regulation?

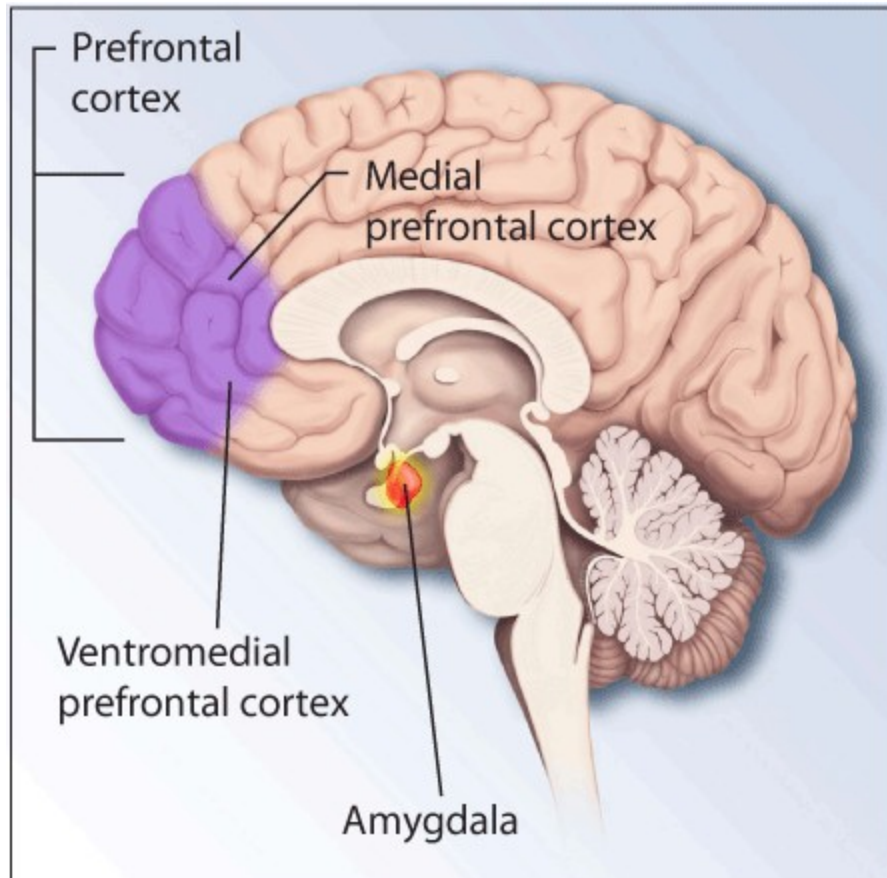




Orienting/Emotion Identification
 Automatic Emotion Regulation
 Voluntary Emotion Regulation
 Regions Implicated in Both Automatic
 and Voluntary Emotion Regulation

Phillips, M. L., Ladouceur, C. D., & Drevets, W. C. (2008). Neural systems underlying voluntary and automatic emotion regulation: Toward a neural model of bipolar disorder. *Molecular Psychiatry*, 13(9), doi:10.1038/mp.2008.82

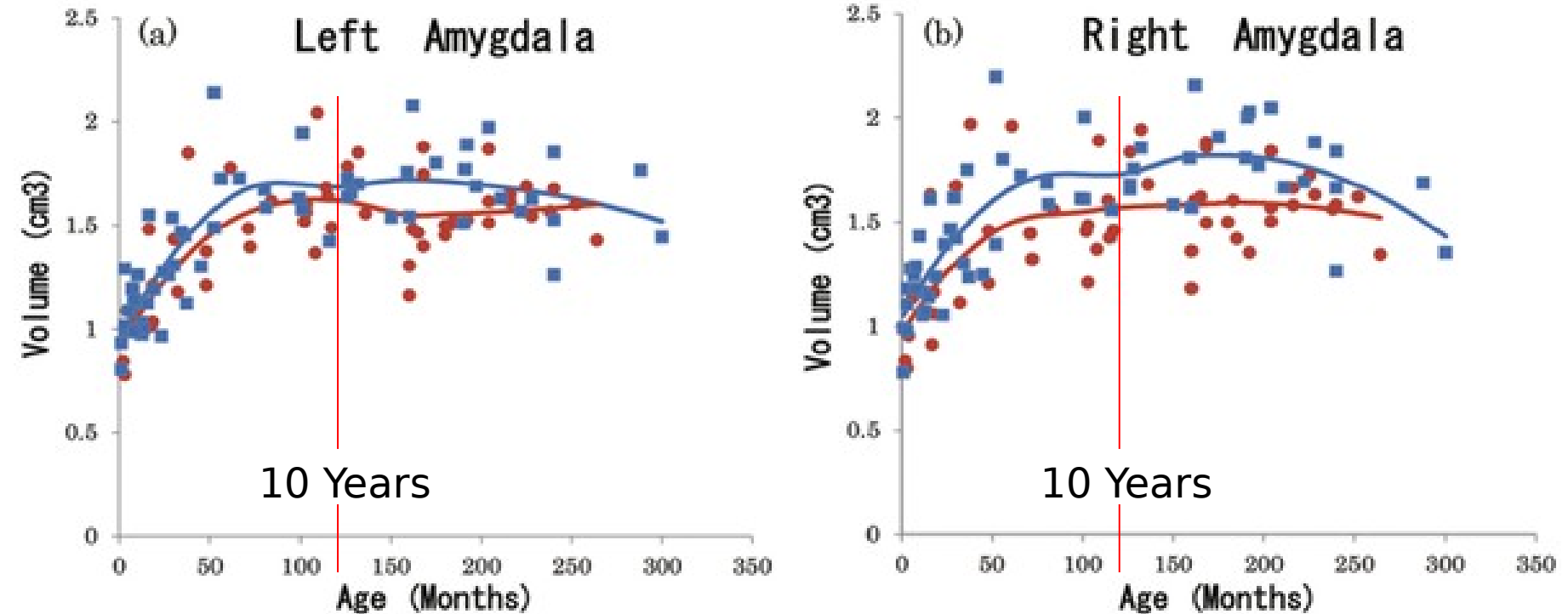
Voluntary
Regulation



Emotion

Emotions and Emotion
Regulation

Amygdala Volume Peaks Between Ages 9-11 Years

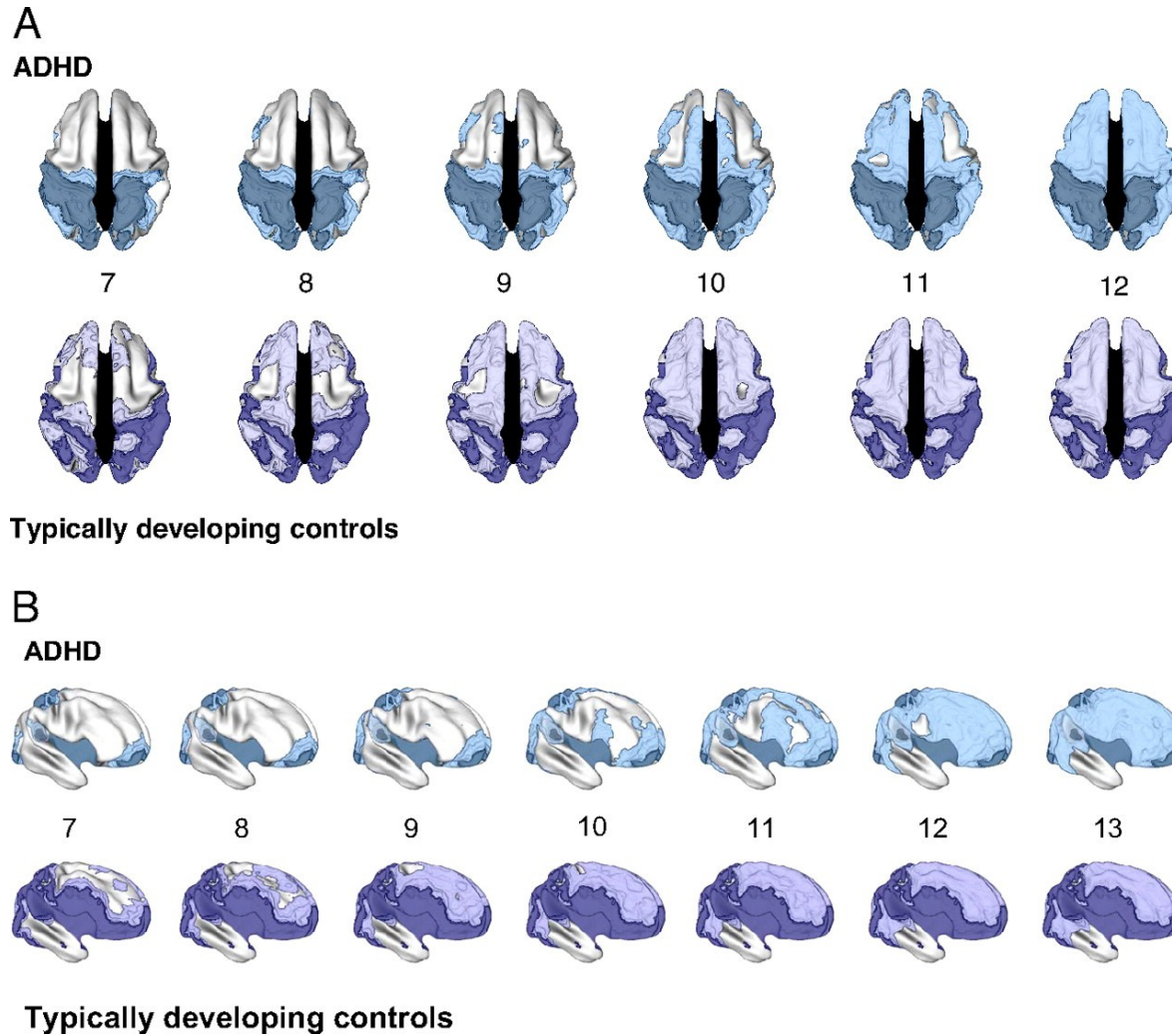


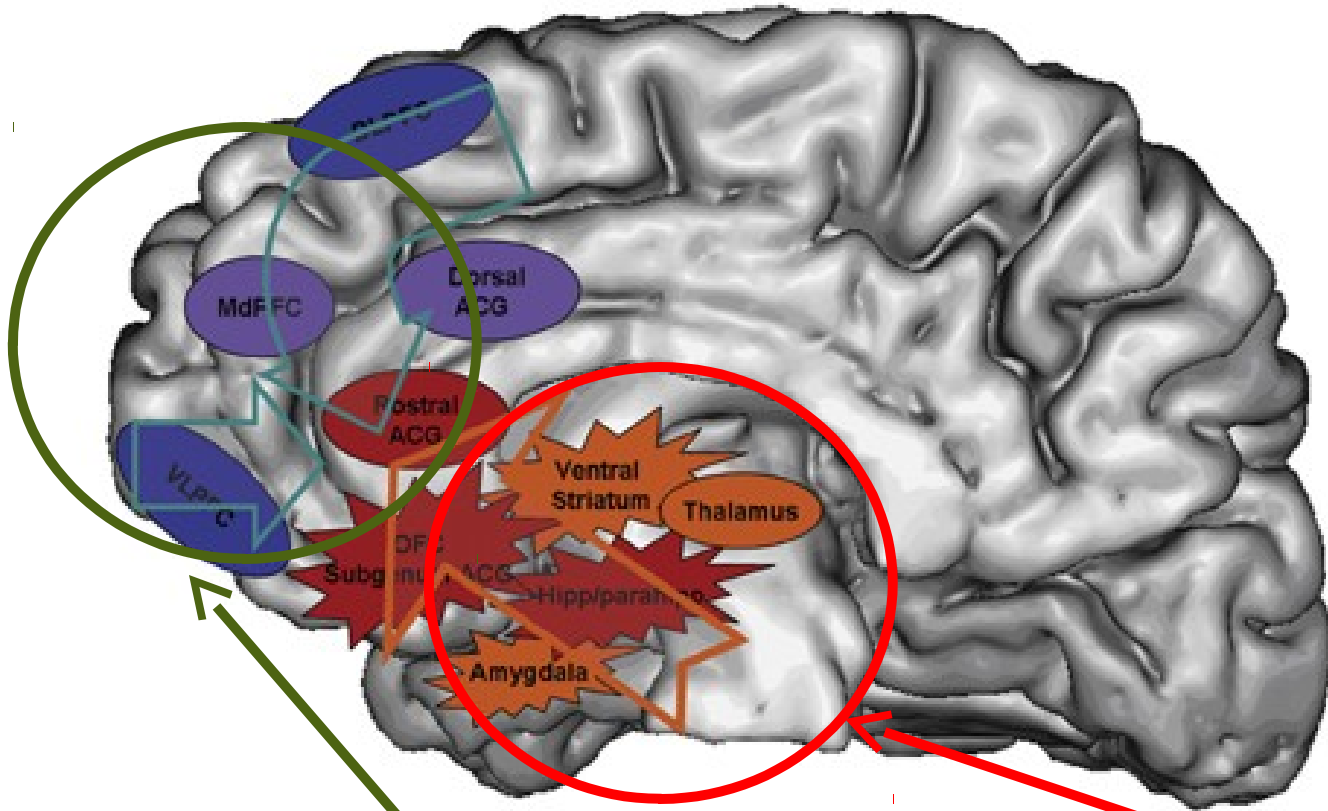
Males: blue line; Females: red line

Uematsu A, Matsui M, Tanaka C, Takahashi T, et al. (2012) Developmental Trajectories of Amygdala and Hippocampus from Infancy to Early Adulthood in Healthy Individuals. *PLoS ONE* 7(10): e46970.
doi:10.1371/journal.pone.0046970

<http://www.plosone.org/article/info:doi/10.1371/journal.pone.0046970>

The age of attaining peak cortical thickness in children with ADHD compared with typically developing children.





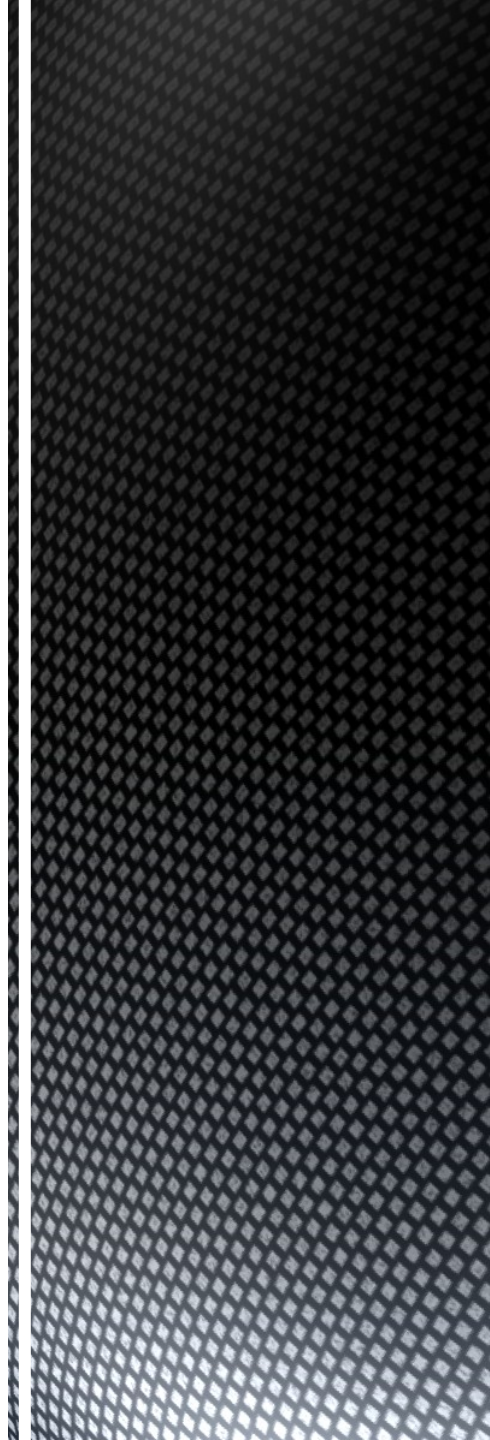
Orienting/Emotion Identification
 Automatic Emotion Regulation
 Voluntary Emotion Regulation
 Regions Implicated in Both Automatic
 and Voluntary Emotion Regulation

Bottom line:

Subcortical regions (orange, some red): develop early

Cortical regions (blue, purple): developmental trajectory extends into adulthood

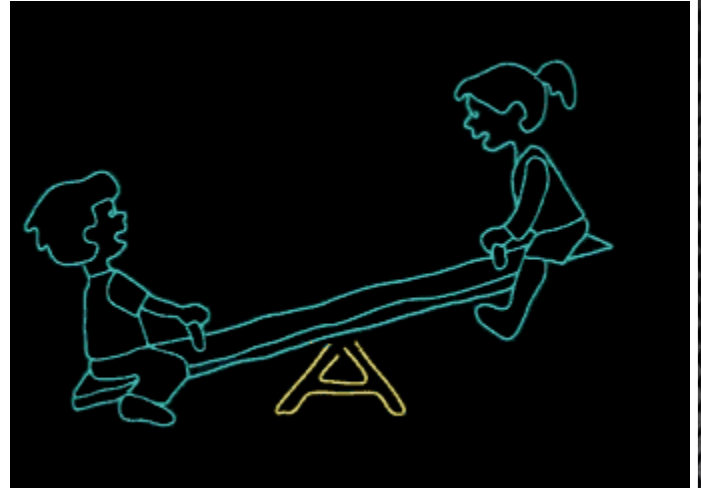
How can the dynamic interplay among intrapersonal and interpersonal characteristics lead to deviations from typical developmental trajectories?

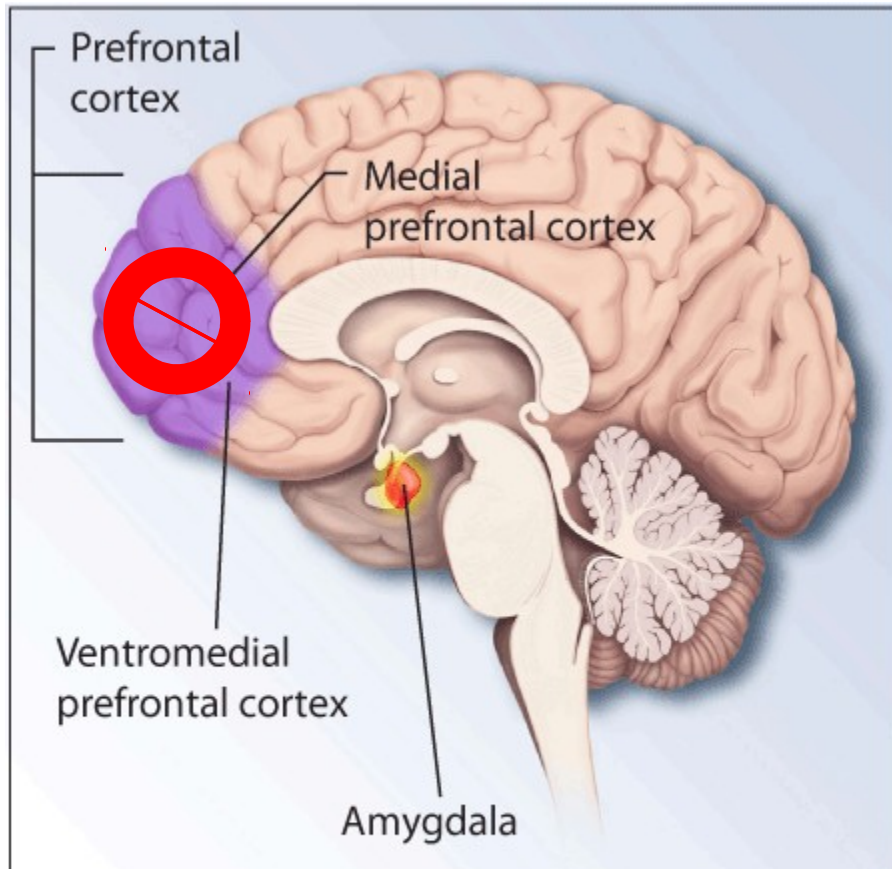


Adaptive Emotion Regulation

**Regulatory
Control**

Emotion



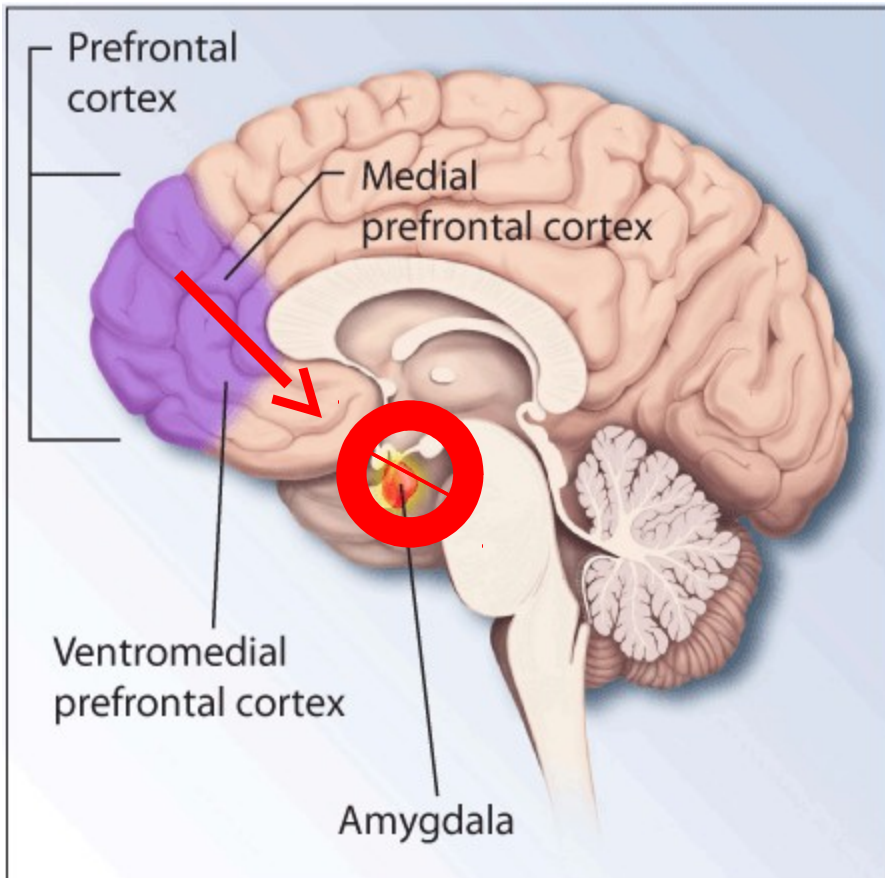


Regulatory Control

Emotion



Under-regulation



Regulatory Control

Emotion



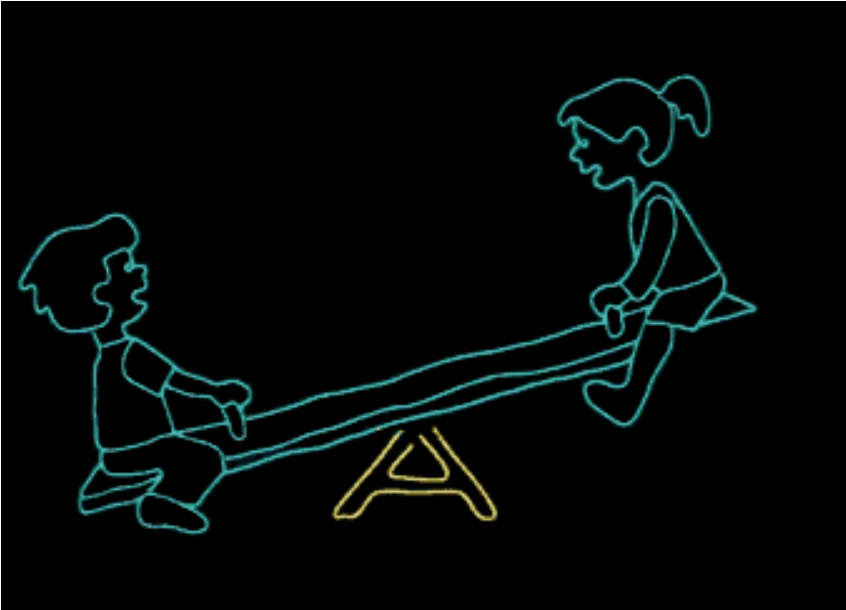
Detachment/Over-regulation

Regulatory
Control

Emotion

Regulatory
Control

Emotion



What determines whether one develops adaptive or maladaptive emotion regulation capacities?

● Intraindividual



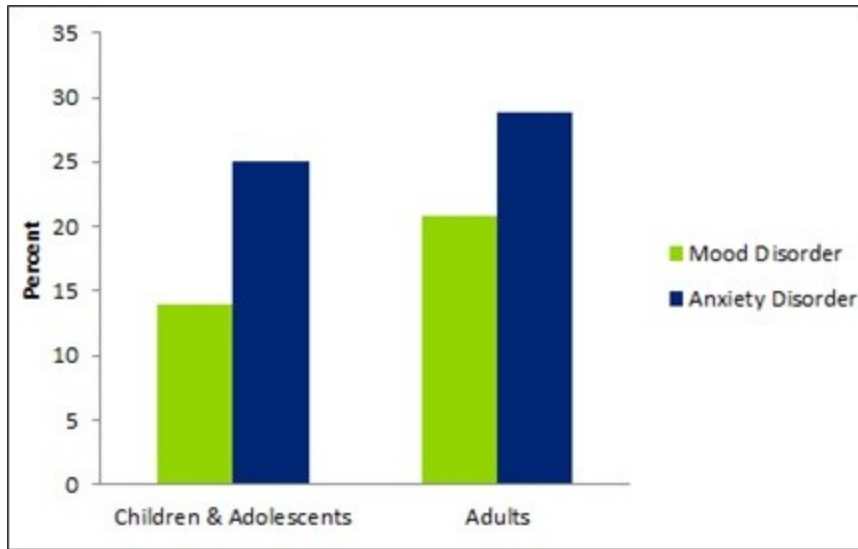
Neurobiological
Expressions

● Interindividua l moderators:

Prenatal and family



**Factors that
Interact to
Determine
Success at
Emotion
Regulation**



Source: NIMH Statistics, "Any Mood Disorder in Children"

Table 1 Gender distribution among nondepressed/nonanxious, subthreshold-depressed/anxious and depressed/anxious groups

	Levels of anxiety		Levels of depression	
	Boys %	Girls %	Boys %	Girls %
No anxiety/depression	50.35	49.65	50.35	49.65
Subthreshold- anxiety/ depression	38.24	61.76	38.24	61.76
Full anxiety/depression	24.07	75.93	24.07	75.93

N = 12,395.

Anxiety and Related Internalizing Conditions as a Model

Balázs, J., et al. (2013). Adolescent subthreshold-depression and anxiety: Psychopathology, functional impairment and increased suicide risk. *Journal Of Child Psychology And Psychiatry*, 54(6), 670-677. doi:10.1111/j.1469-7610.2012.02516.x

- Intraindividual moderators
 - Prenatal environment
 - Family environment
 - Parent mood/anxiety disorders
 - Family interaction patterns
 - School/peer environment

- Interactions

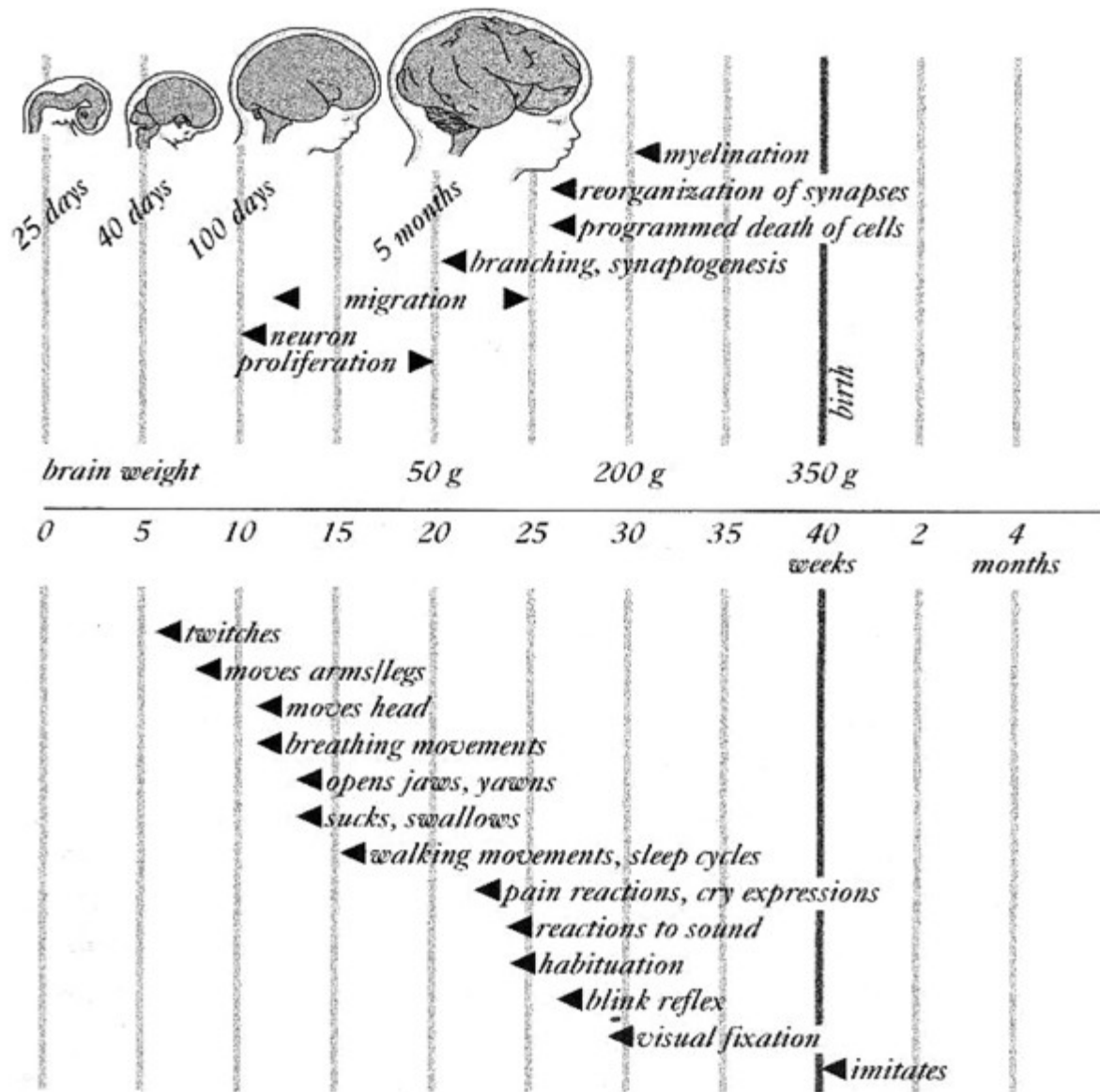
- McGrath article

Genetic Profile

- Maternal physical health
- Maternal mental health

Prenatal
Environment

Outcomes associated with fetal risk exposure during pregnancy



The development of the brain displayed in a schematic form. *Source:* The Newborn Brain. Edited by Lagercrantz H, Hanson M, Evrard P, Rodeck C. Cambridge University

Table 3

Logistic regression analysis of anxiety status.

Variable	B (SE B)	exp B
Constant	.81 (.16)**	1.11
Age	-.16 (.14)	.85
Sex	.04 (.32)	1.04
Mother's anxiety score	.01 (.01)	1.01
Prenatal risk factors	.57 (.22)**	1.76
Visuospatial deviation	.14 (.05)**	1.15
Prenatal risk factors × visuospatial deviation	.10 (.06)*	1.11

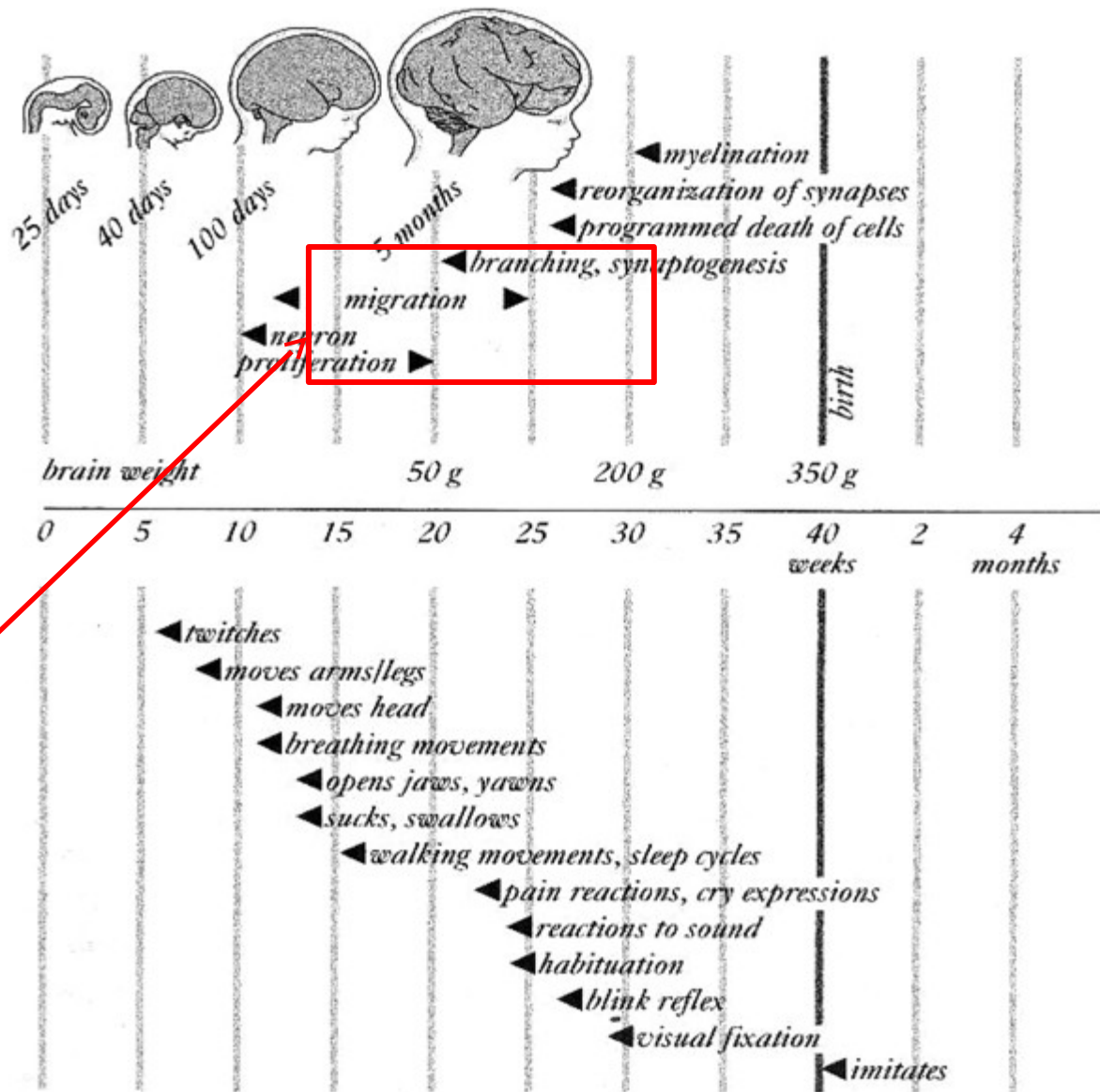
Note: $R^2 = 13.50$ (Nagelkerke). Model $\chi^2(21.28) = p < .01$.* $p < .05$.** $p < .01$.**Table 4**

B and exp B (proportionate change in odds for being high-anxious) 0, 1, 2, and 3 prenatal risk factors (linked to level of visuospatial deviation).

	0 prenatal risk factor × visuospatial deviation	1 prenatal risk factor × visuospatial deviation	2 prenatal risk factor × visuospatial deviation	3 prenatal risk factor × visuospatial deviation
B	.06	.17	.27	.66
exp B	1.06	1.18	1.31	1.92

Cumulative prenatal risk exposure significantly predicted anxiety in 8-12 year olds

Simon, E., Bögels, S., Stoel, R., & De Schutter, S. (2009). Risk factors occurring during pregnancy and birth in relation to brain functioning and child's anxiety. *Journal Of Anxiety Disorders, 23*(8), 1024-1030. doi:10.1016/j.janxdis.2009.07.002



Outcomes associated with maternal anxiety during pregnancy

Research focused on 19th-31st weeks of gestation

The development of the brain displayed in a schematic form. *Source: The Newborn Brain. Edited by Lagercrantz H, Hanson M, Evrard P, Rodeck C. Cambridge University*

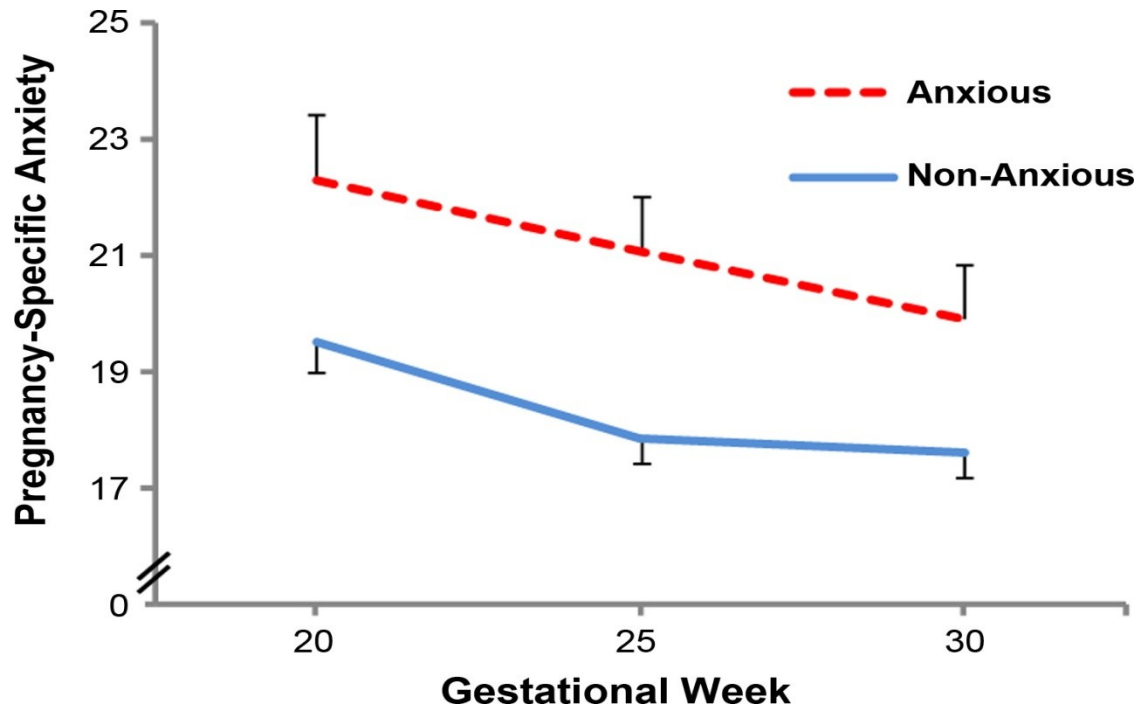
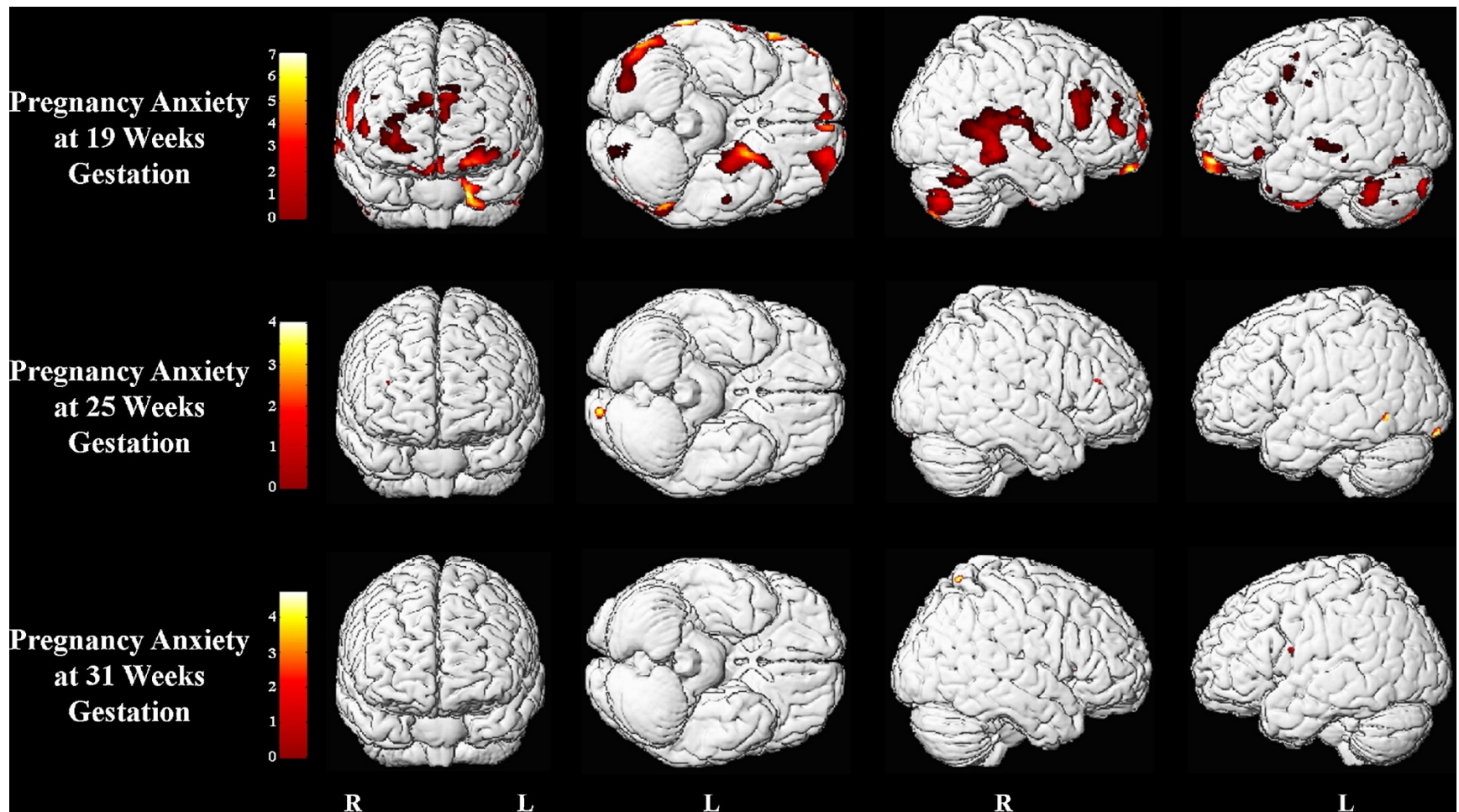


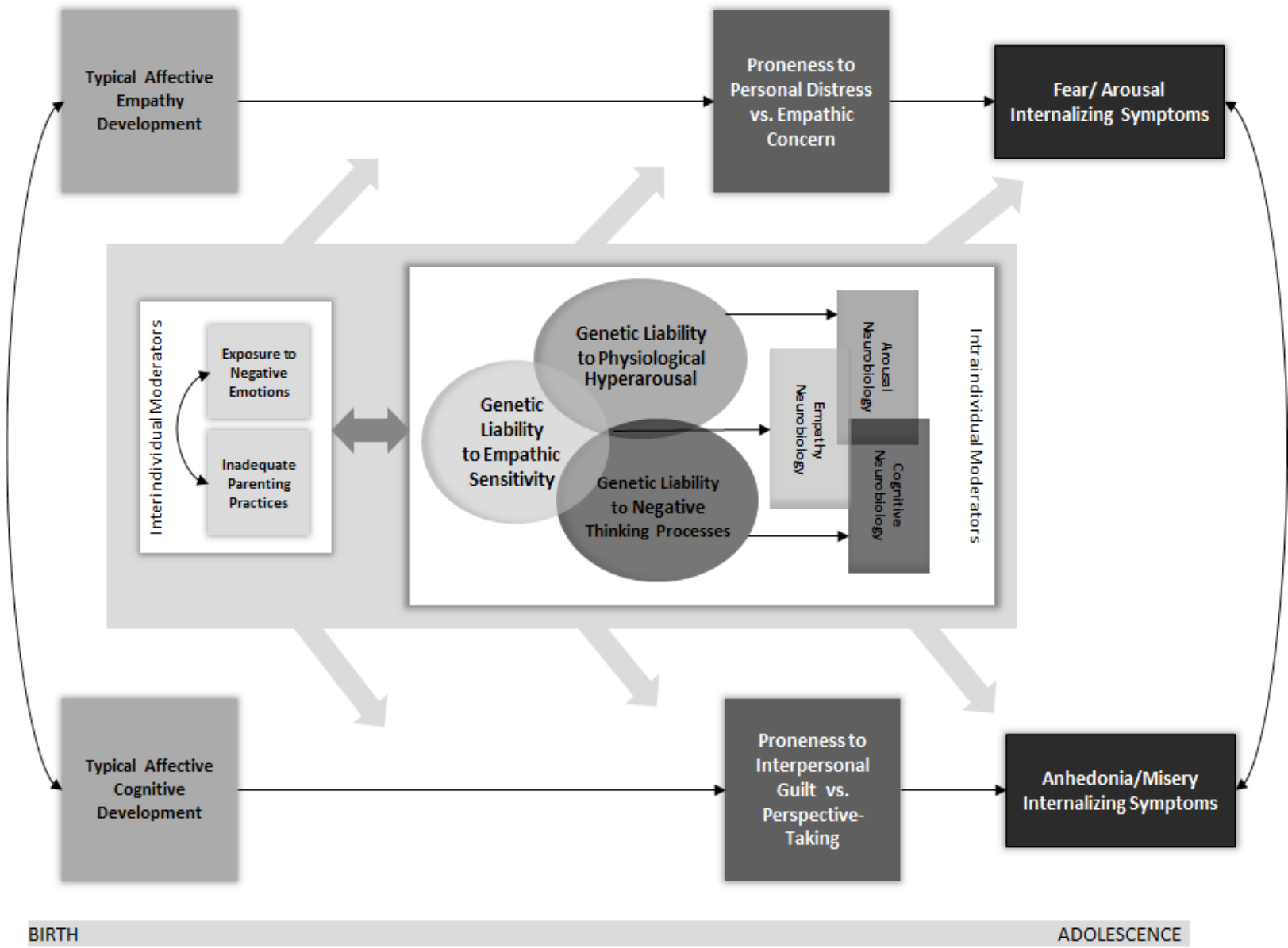
Figure 2 Pregnancy-specific anxiety predicts anxiety classification during preadolescence. Children who are in the normal range for anxiety were exposed to lower pregnancy-specific anxiety. In contrast, children who were exposed to elevated pregnancy-specific anxiety during gestation are significantly more likely to be rated in the anxious/borderline anxious range.

N=178 mothers & their 6-9 year old children



Areas of reduced gray matter volume in association with pregnancy anxiety at 19, 25 and 31 weeks gestation. Voxels with $p < 0.001$ (uncorrected) are displayed.

Buss et al. (2010). High pregnancy anxiety during mid-gestation is associated with decreased gray matter density in 6-9-year-old children. *Psychoneuroendocrinology*, 35(1), 141 - 153. [dx.doi.org/10.1016/j.psyneuen.2009.07.010](https://doi.org/10.1016/j.psyneuen.2009.07.010)



Tone & Tully (2014). In press. *Development & Psychopathology*