



Structural MRI of the brain in neonates

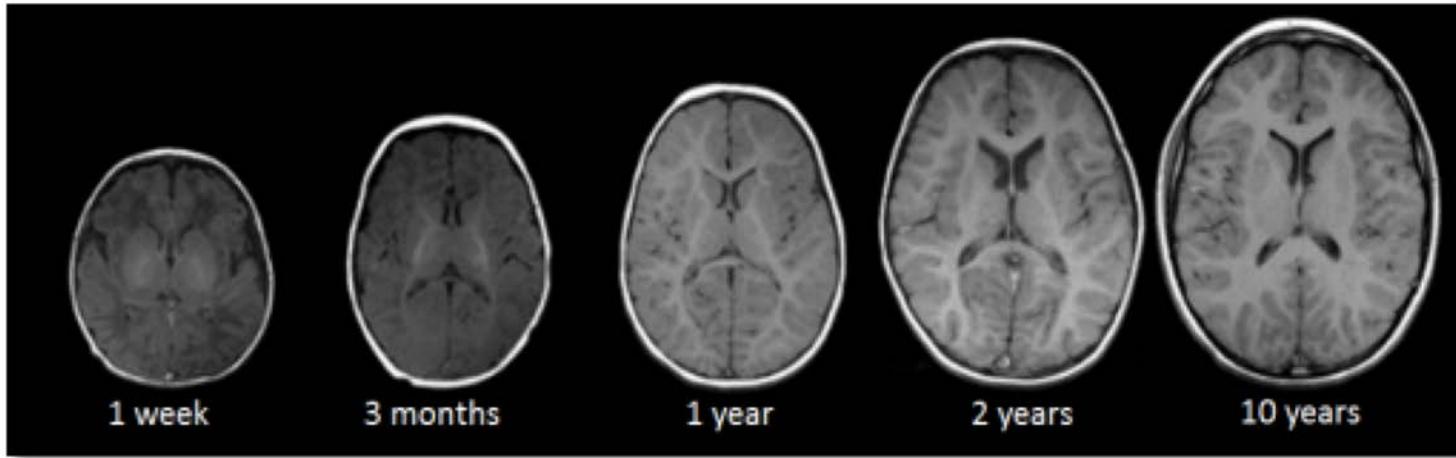


III SIZE MATTERS



<http://www.mrineonatalbrain.com/maryrutherfordimaging/downloads/presentations/Neonatal-MR-optimisation.pdf>

III NEONATAL POPULATION



Small head circumference (only $\frac{1}{4}$ of adult head)

Rapid maturation in first year of life.

Major changes in water content and myelination: adjusted protocols needed.

Non-cooperative → **high probability of motion artifacts**

Patient preparation, materials used and acquisition protocols need to be adapted specifically for neonates.

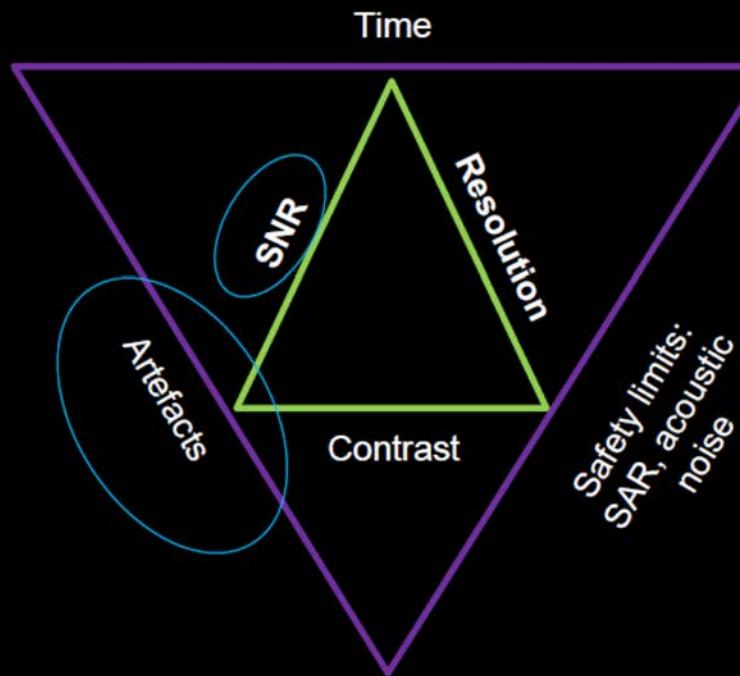
→ Using protocols for larger children and adults in neonates is a guarantee of poor images.

III PRIMARY CLINICAL INDICATIONS

- Severe prematurity → Born at less than 32 weeks gestational age
 - => PVL = periventricular leukomalacia (white matter disease of prematurity)
 - => Germinal matrix hemorrhage
- Asphyxia (= lack of oxygen, usually with difficult birth delivery)
 - → evaluation of brain damage
 - → Prognosis
 - → Clinical management (supportive, expectant, palliative)
- Dysmorphism and abnormal muscle tone (hypotonia, hypertonia)
 - → Malformations? Abnormal development of the brain
 - → Inborn errors of metabolism (gray and/or white matter disorders). Fault in DNA leads to accumulation of toxic metabolites.
 - → Management (e.g. removing nutritional substances that are not broken down correctly from diet)
 - → Prognosis (often poor)
 - → Genetic screening/counseling parents/family members (future pregnancies)
- Epilepsy / convulsions / infantile spasms
- Infection (eg. bacterial or tuberculous meningitis)

III IMAGE QUALITY OPTIMIZATION

Image quality optimisation process in MRI



III OPTIMIZATION OF IMAGE QUALITY ON 3T

Advantages at high field MRI

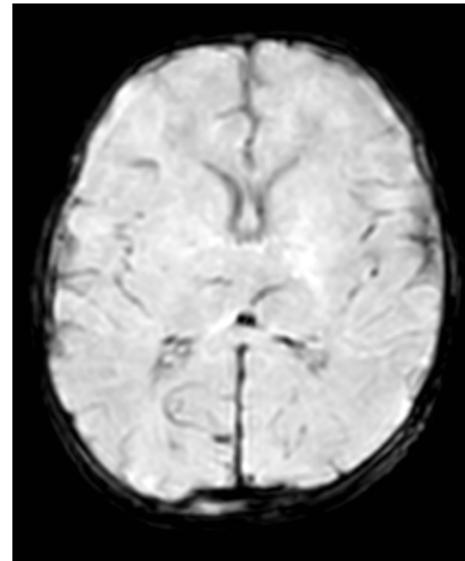
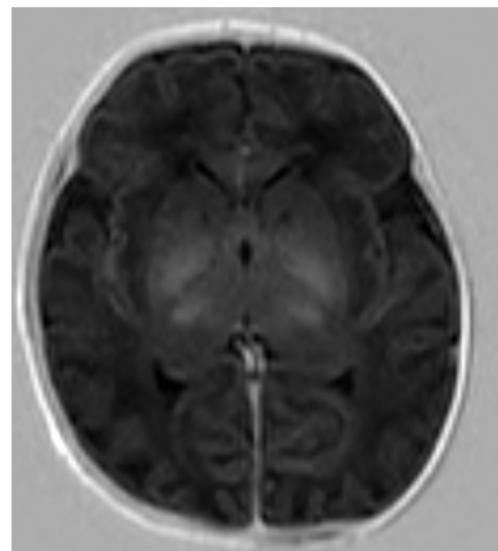
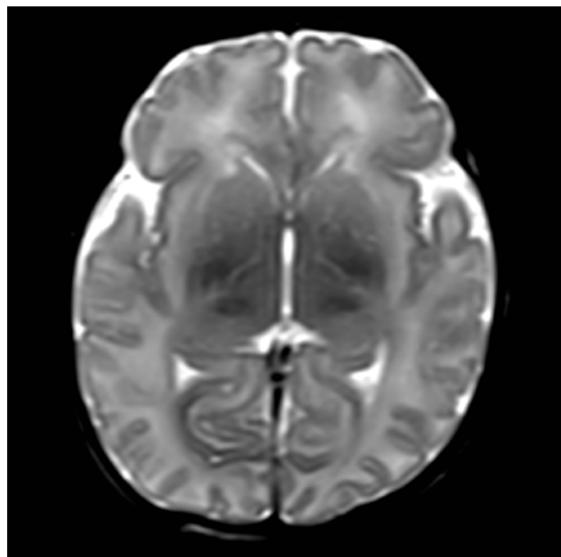
Higher signal to noise ratio

Spatial resolution

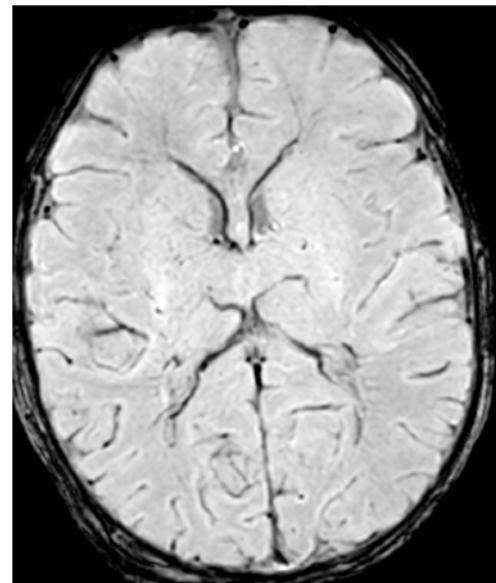
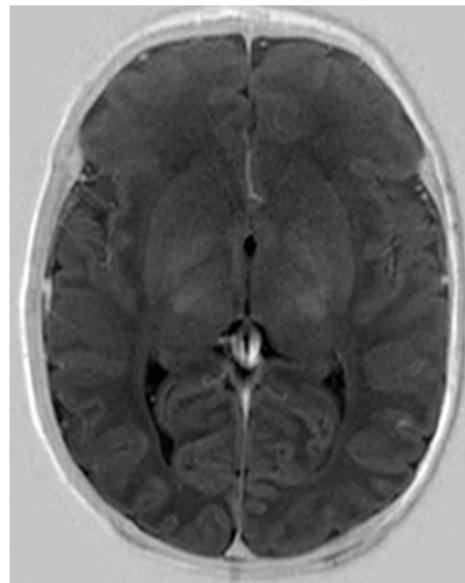
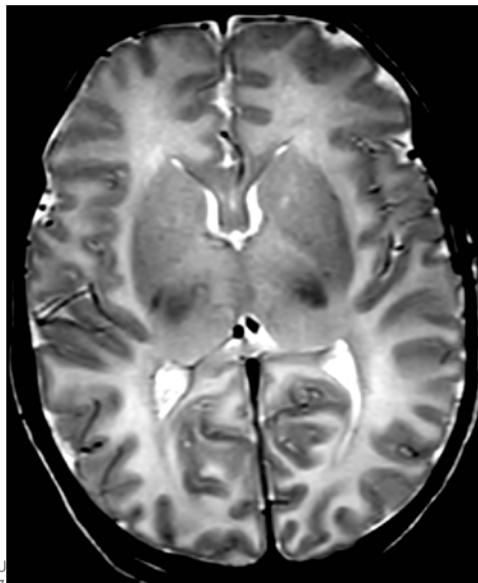
Temporal resolution

III IMAGE QUALITY OF NEONATAL MRI OF THE BRAIN

1,5T



3T



III INFANT PREPARATION

Sedation

- Chloral hydrate (syrup)
 - Most used worldwide
 - Readily available
 - Relatively wide margin of safety - oldest and best studied sedative
 - Best taken on an empty stomach
 - Children sleep after 15-30 minutes
 - administration orally (tastes bad), via nasogastric tube (preferred at UZ Brussel) or rectally.
 - Dosage and method of administration are decision of the pediatrician, not of the radiologist!
 - Collaboration with children's daycare hospital
 - MONITORING !
- Feed and sleep
 - Scanning in natural sleep after feeding
 - Works best with children < 6 weeks
 - Requires good coordination/timing
 - Greater chance of non-diagnostic images
- UZ Brussel :
 - Feed and sleep for infants up to 3 months of age
 - Sedation with chloral hydrate in children up to 12 months of age.
 - Older children under anesthesia (remote team anesthesia) – Propofol (Diprivan)



III INFANT PREPARATION

Ear protection – immobilization with vacuum mattress



- Mini Muffs; *Natus Medical, San Carlos, California*
- Headphone

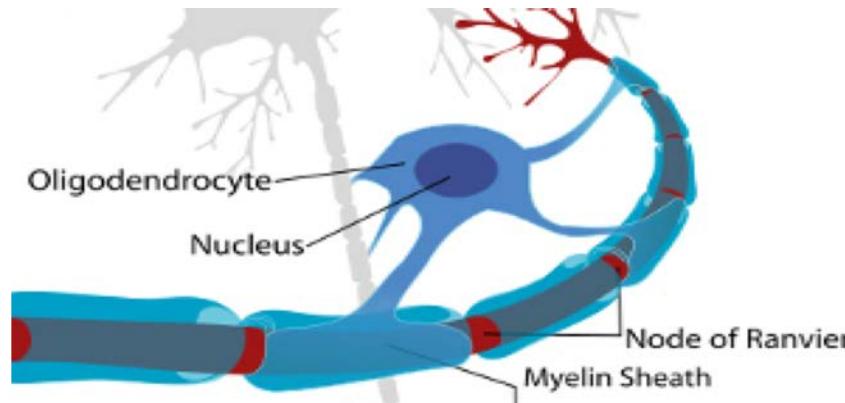
Foto : Mathur, A.M., Neil, J.J., McKinstry, R.C. et al. Pediatr Radiol (2008) 38: 260. <https://doi.org/10.1007/s00247-007-0705-9>

A sedated infant still asleep post MRI within the Med-Vac immobiliser.



Therese Ibrahim et al. Arch Dis Child Fetal Neonatal Ed 2015;100:F465-F466

III MYELINATION

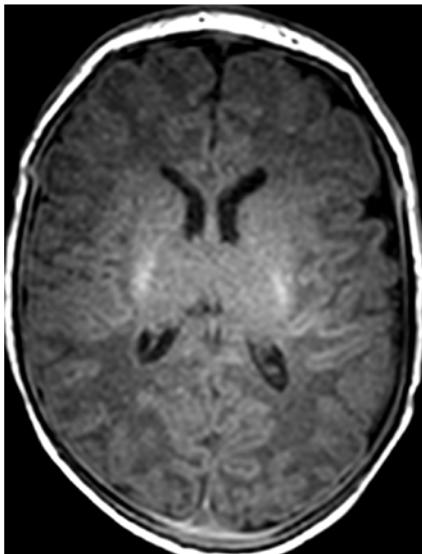


- Has a major impact on T1 and T2 tissue contrast
- Myelin shaft is a plasma membrane surrounding axons in the nervous system
- In adults 50-60% of dry weight of the brain
- Forms an insulator => Requires less energy for conduction: up to 5000 times less energy for the same speed at much smaller diameters (Ritchie, 1982)
- Important for normal brain functioning
- In newborns most of the white matter is not yet myelinated.
- After one year, the brain is already largely myelinated.

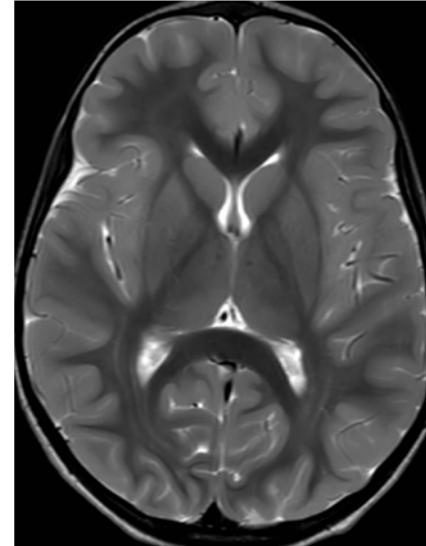
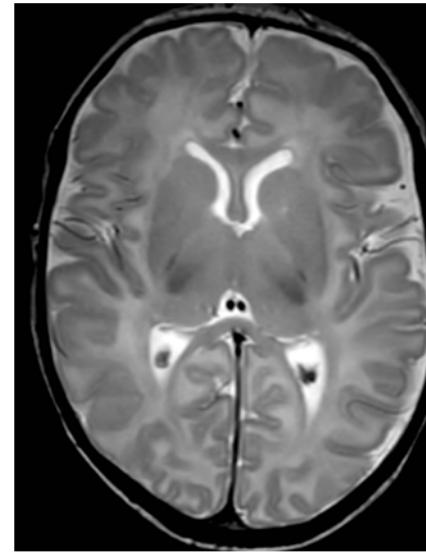
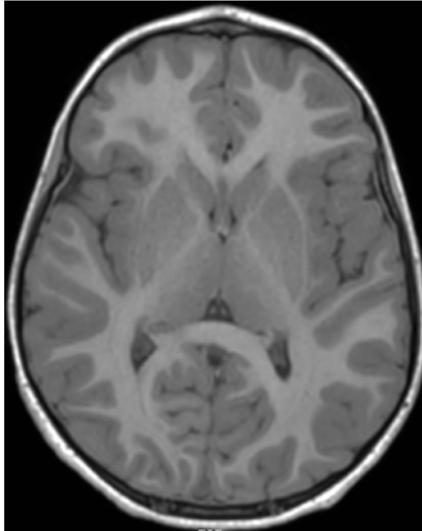
III MYELINATION AND TISSUE CONTRAST

Lack of myelination in neonates: Inverted gray-white contrast on T1 and T2

5 days



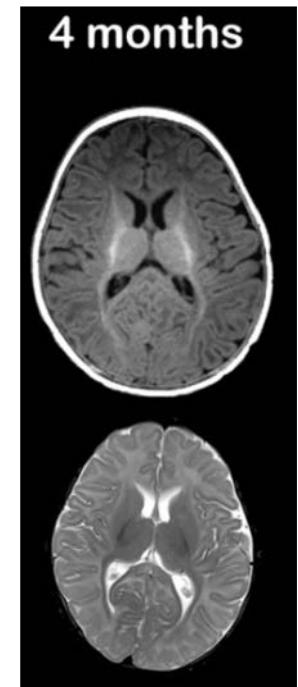
3 years



III EVALUATION OF MYELINATION

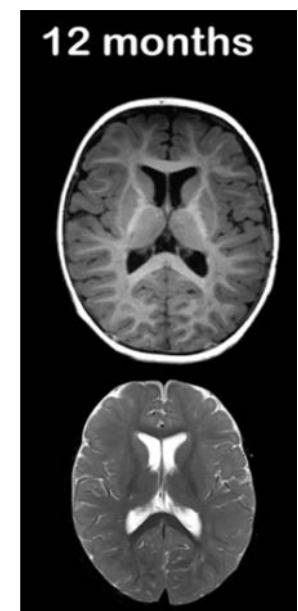
T1

- most sensitive sequence in children <1 year of age
- myelination represented by T1 hyperintensity



T2

- most sensitive sequence in children between the age of 1 and 2 demonstrating a gradual shift from hyper- to hypo-intense relative to grey matter
 - myelination represented by T1 hypointensity
-
- The only area to remain hyperintense after the age of 2 years, and often for quite some time, is the peritrigonal region ⁴ which is called terminal zones of myelination
 - (Radiopaedia)



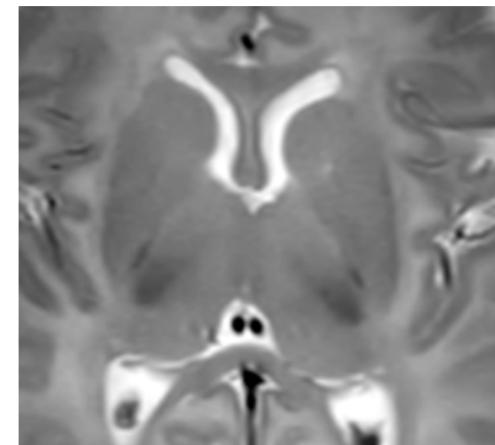
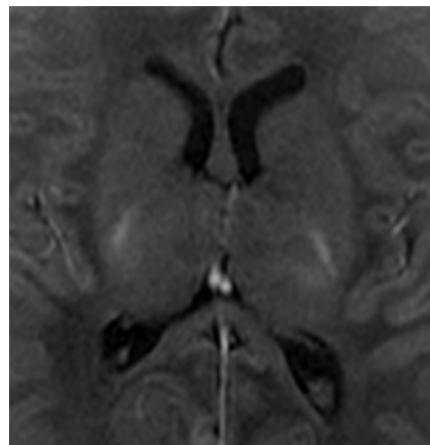
III MYELINATION

Progression of myelination is predictable and correlates strongly with developmental milestones.

1. Central to peripheral
2. Caudal to rostral
3. Dorsal to rostral
4. Sensory before motor

T1: Myeline = hyperintens

T2: Myeline = hypointens

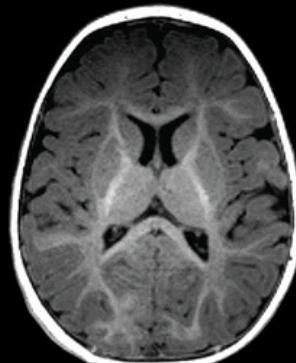
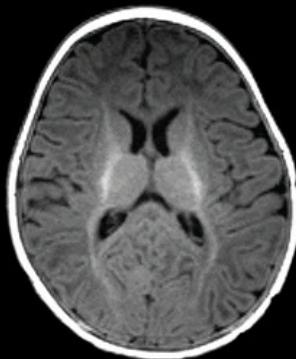


Genetic defects can make myelin formation more difficult or cause degradation of already formed myelin → Leukodystrophy

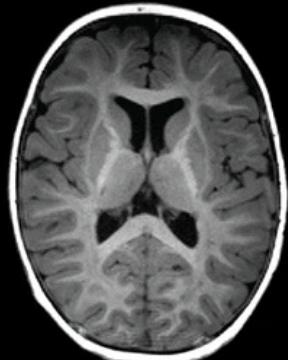
III NORMAL MYELINATION

4 months 6.8 months 12 months 30 months 72 months

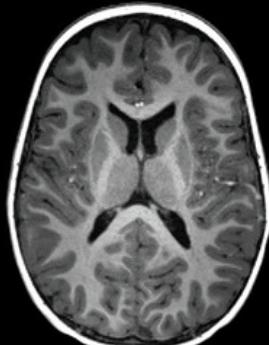
T1



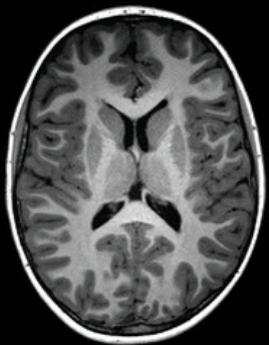
12 months



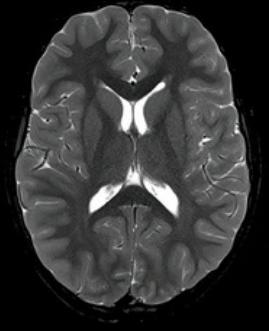
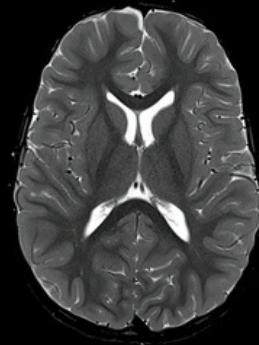
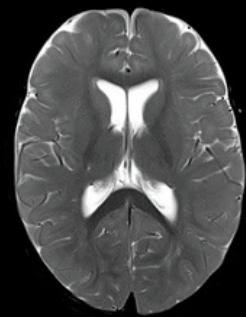
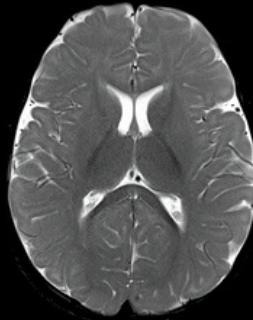
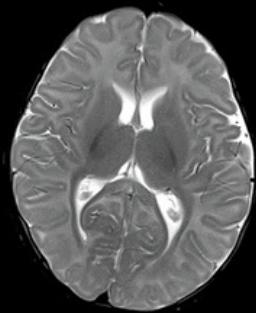
30 months



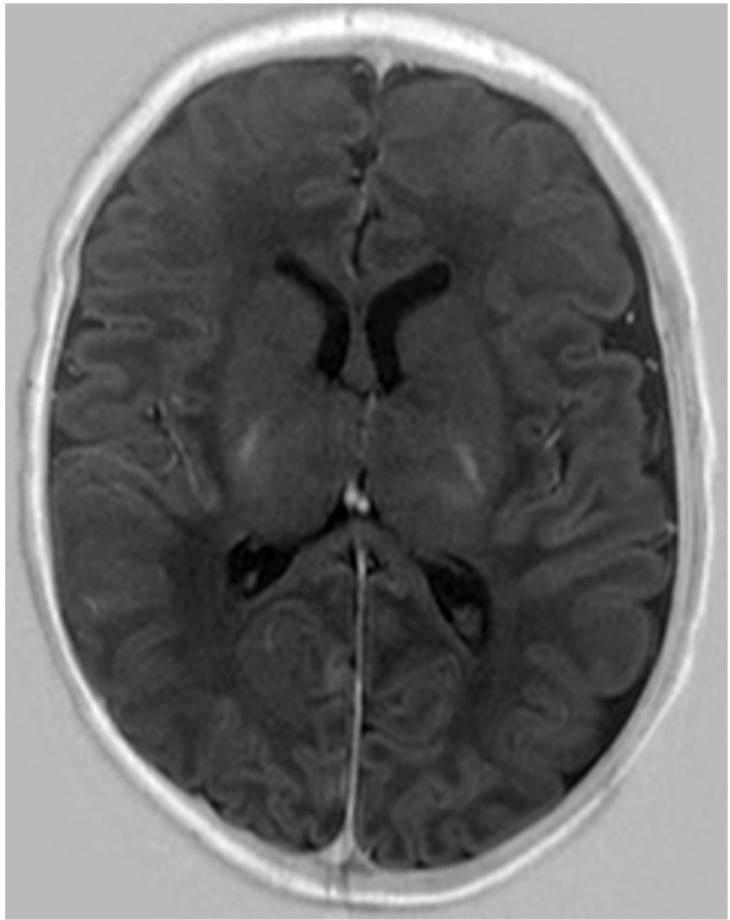
72 months



T2

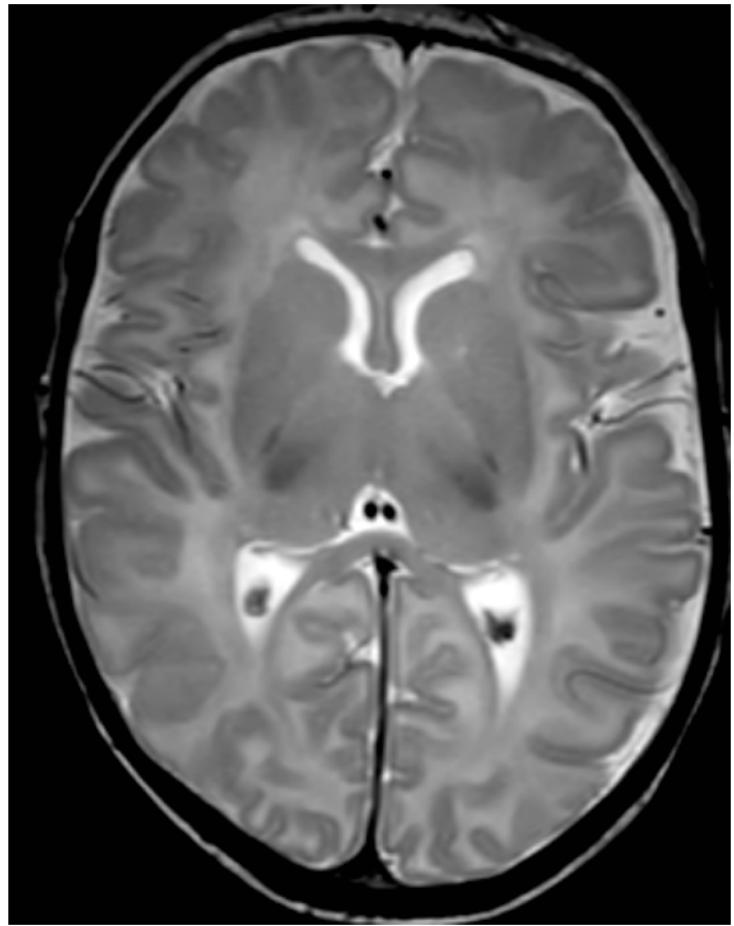


III BASIC SEQUENCES



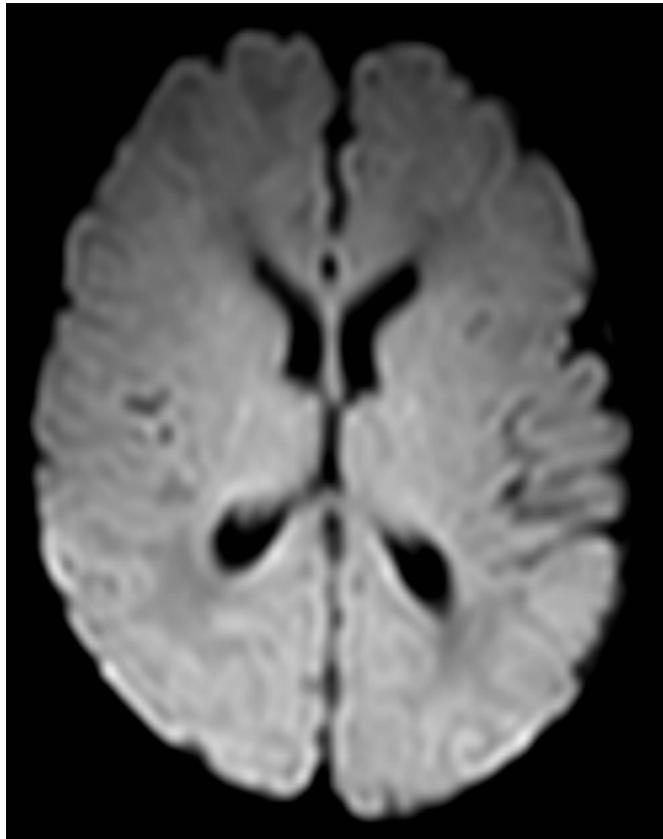
T1 TSE (3 mm)

TR 4130
TE 120

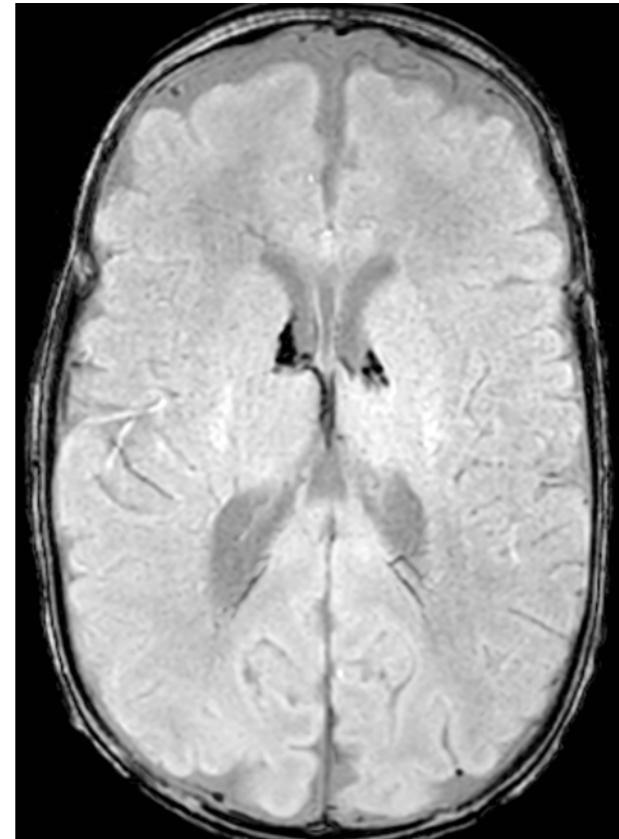


T2 TSE (3.5 mm)

III BASIC SEQUENCES

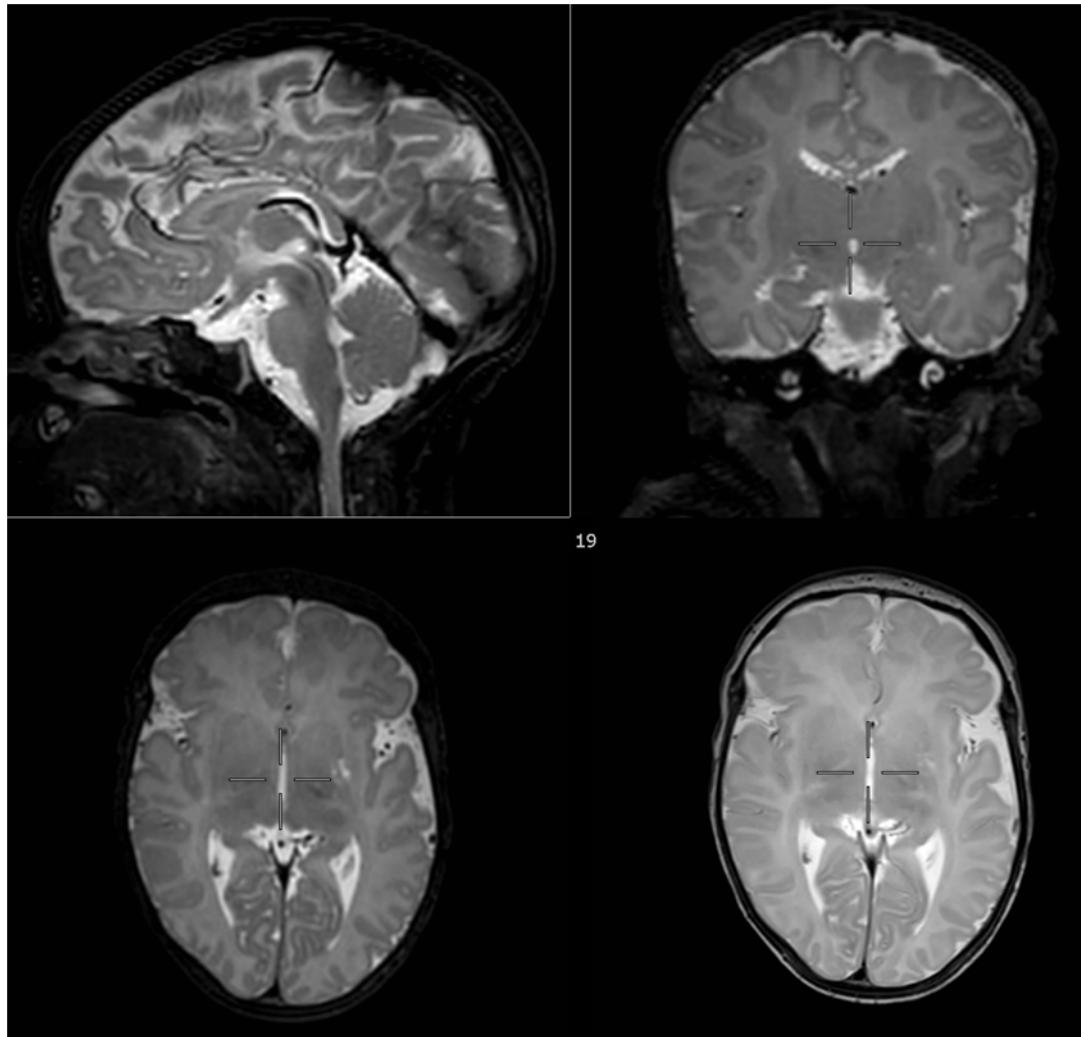


DWI b₀-b1000 (4mm)



SWI (3D 2 mm)

III BASIC SEQUENCES



3D T2 TSE

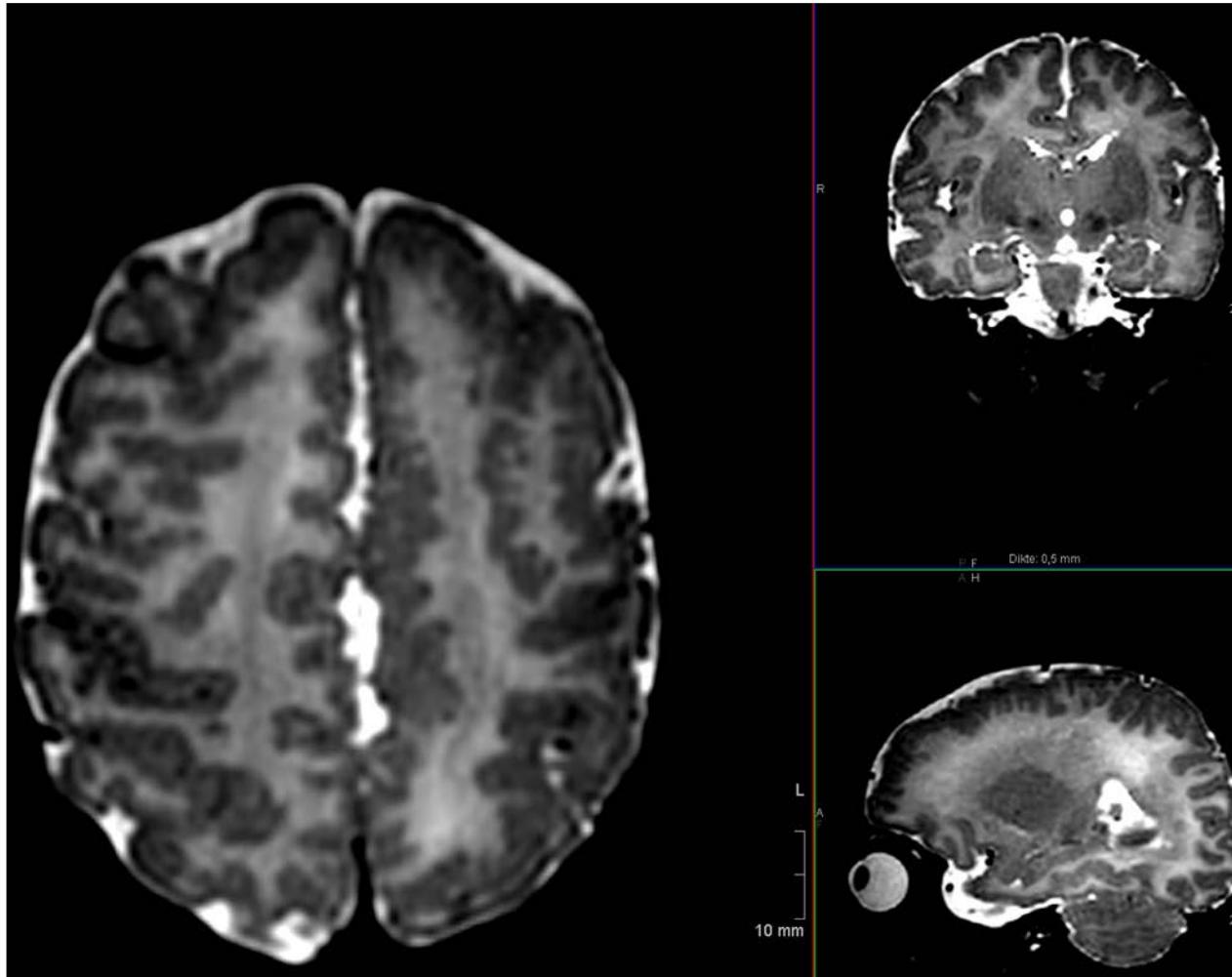
Multiplanar reconstructions

- Less contrast in white matter
- Lower in plane resolution
- Better delineation of the cortex
 - detection of subtle cortical malformations
- Flow void through aqueduct

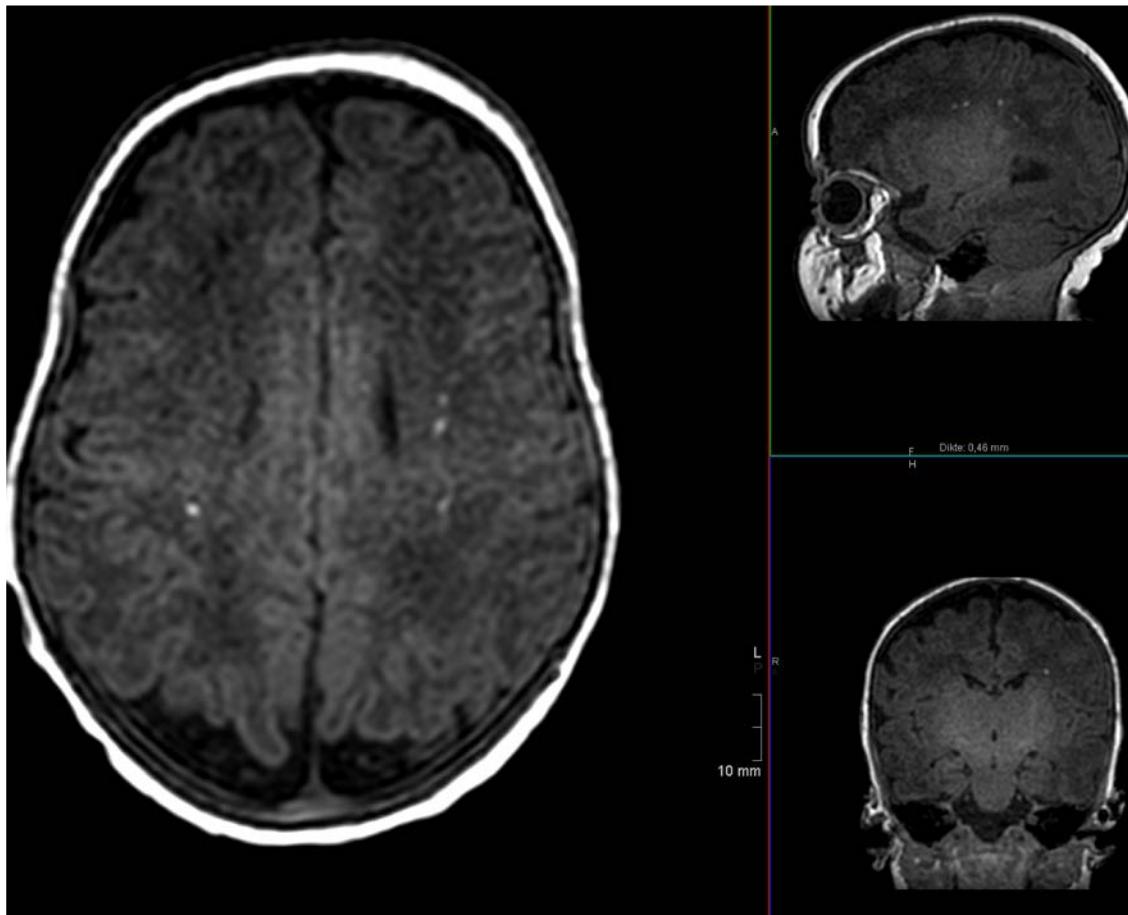
TR:2500
TE:330
FOV 190

III BASIC SEQUENCES

3D T2 + MPR: polymicrogyria (PMG)



III BASIC SEQUENCES



3D T1 TFE

Multiplanar reconstruction

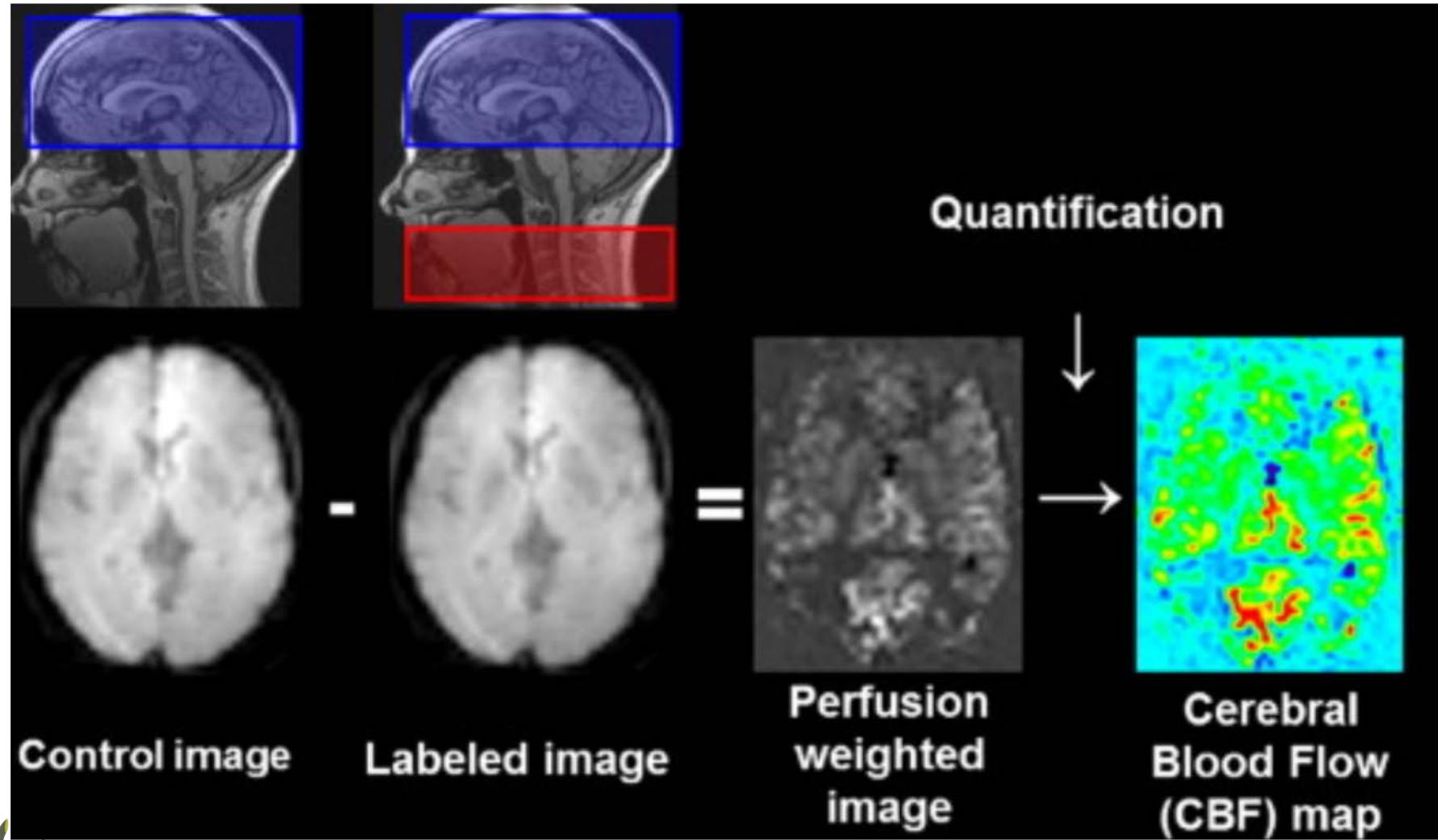
- Detection of small white matter lesions
- location in motor tract => risk of impaired motor development

TR:6100

TE:2750

FOV 180

III ASL (ARTERIAL SPIN LABELING)

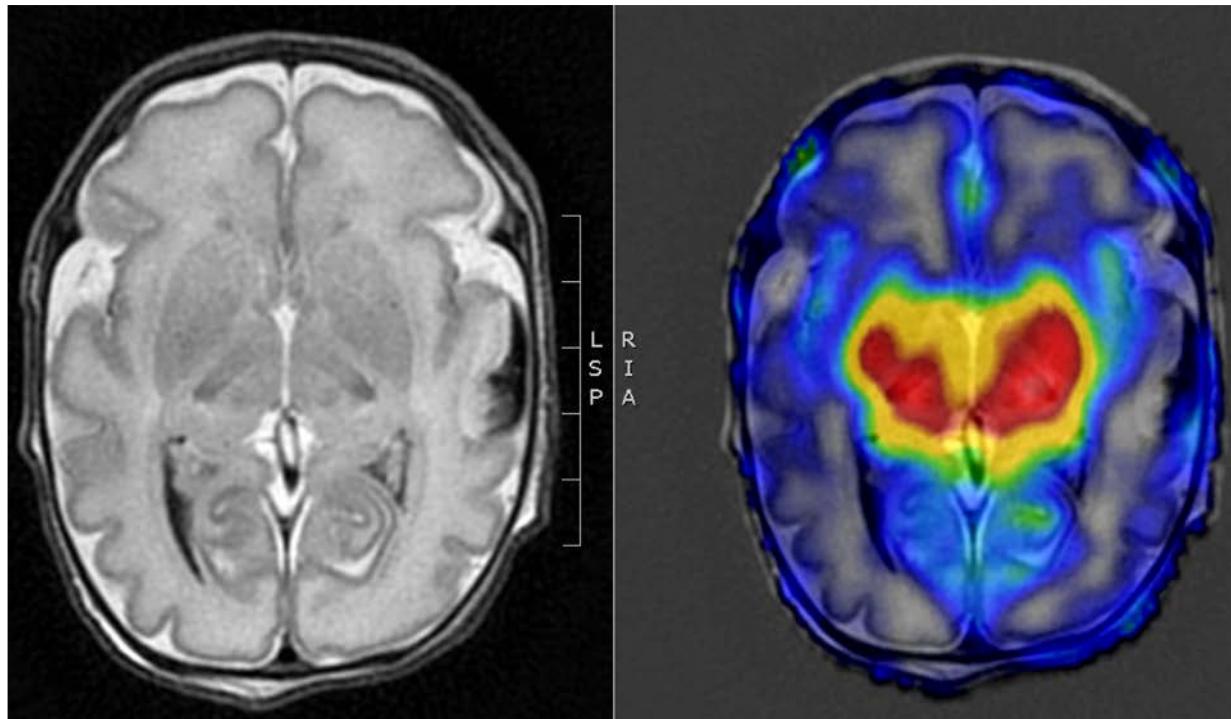




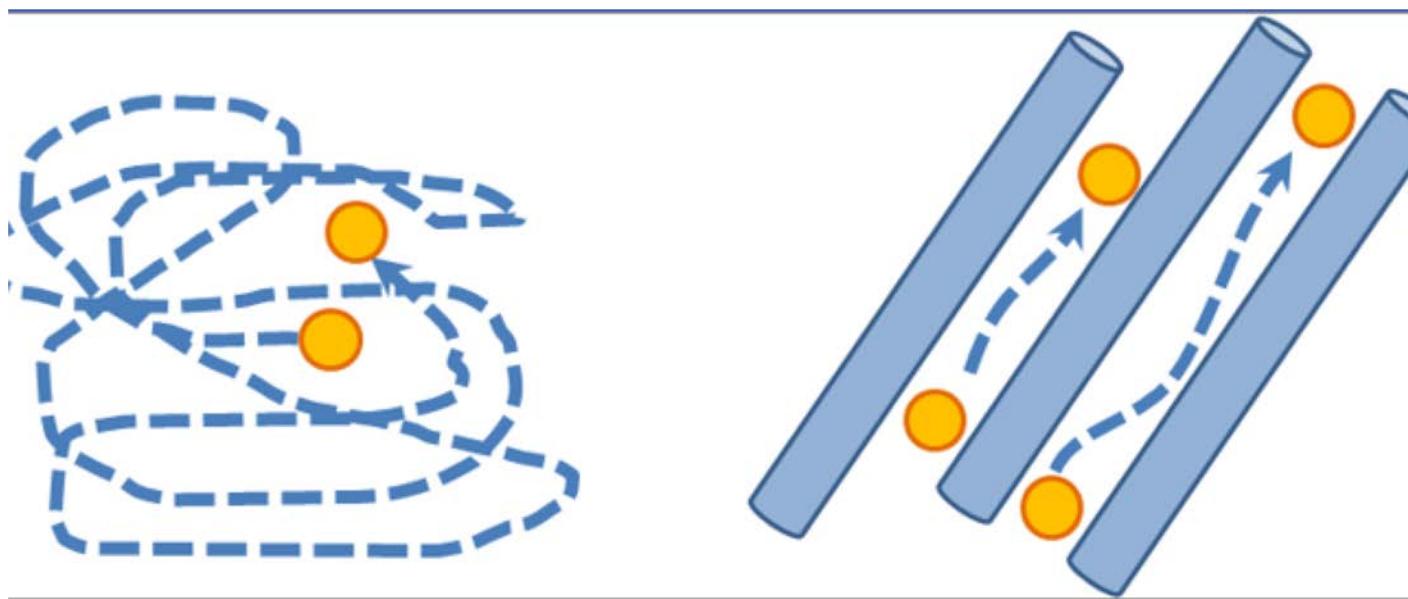
ASL

HIE (hypoxic ischemic encephalopathy)

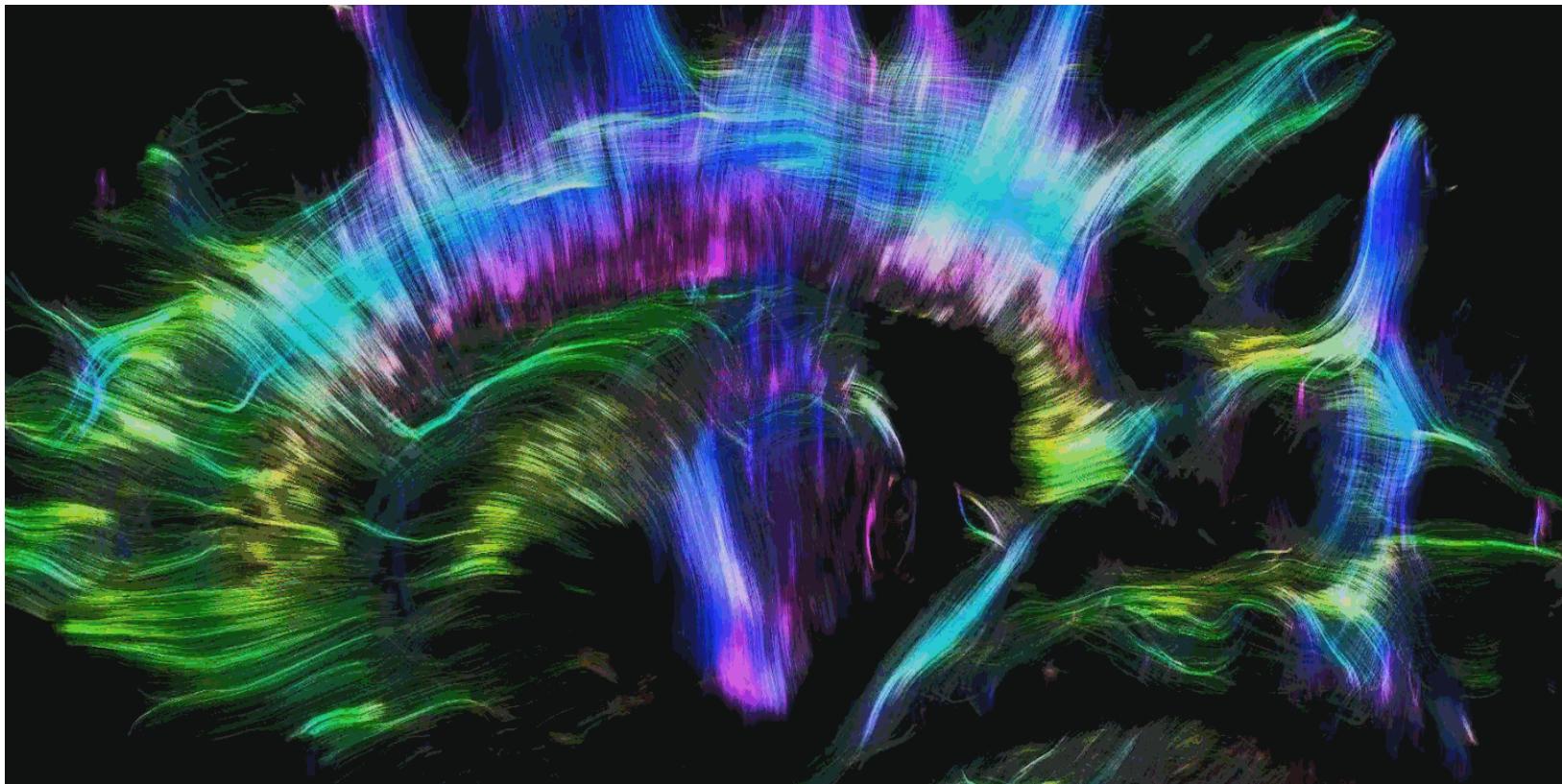
- asphyxia (due to problems before or during delivery)
- **Cardiac arrest**
- Severe oxygen deficit during longer time => **brain damage**
- Hyperperfusion (reperfusion) in basal ganglia + lactate (spectroscopy) → **worse outcome**



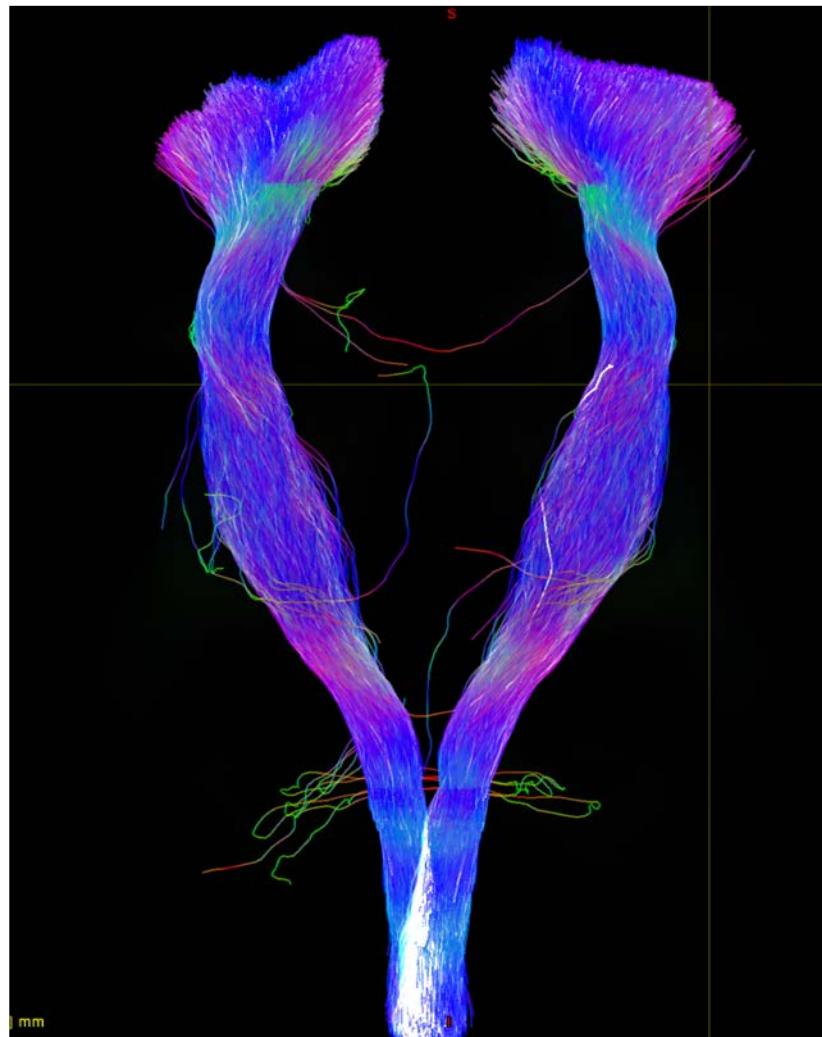
III DIFFUSION + TRACTOGRAPHY – BROWNIAN MOTION

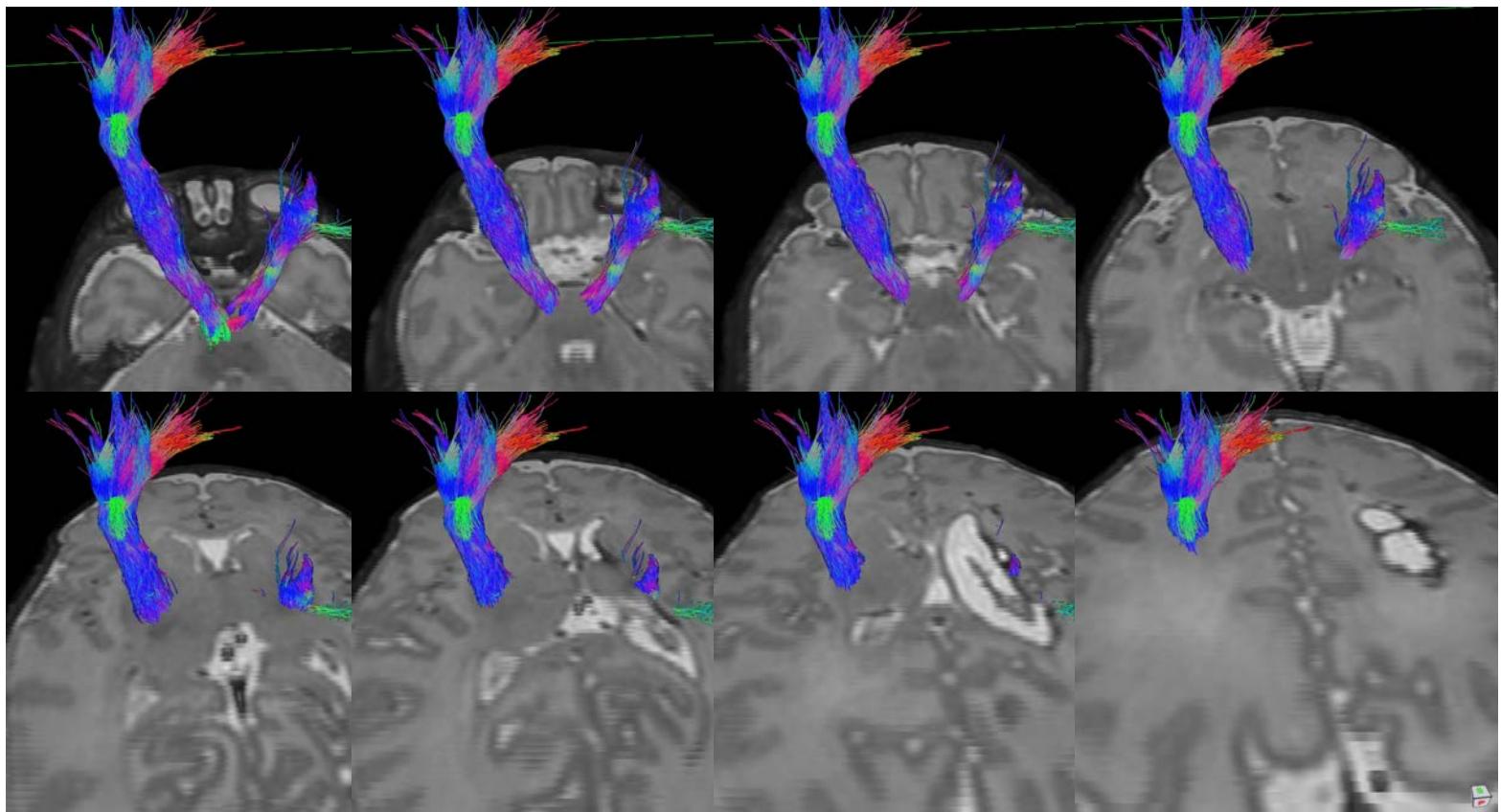


III NEONATALE DIFFUSIE TRACTOGRAFIE



III CORTICOSPINAL TRACT



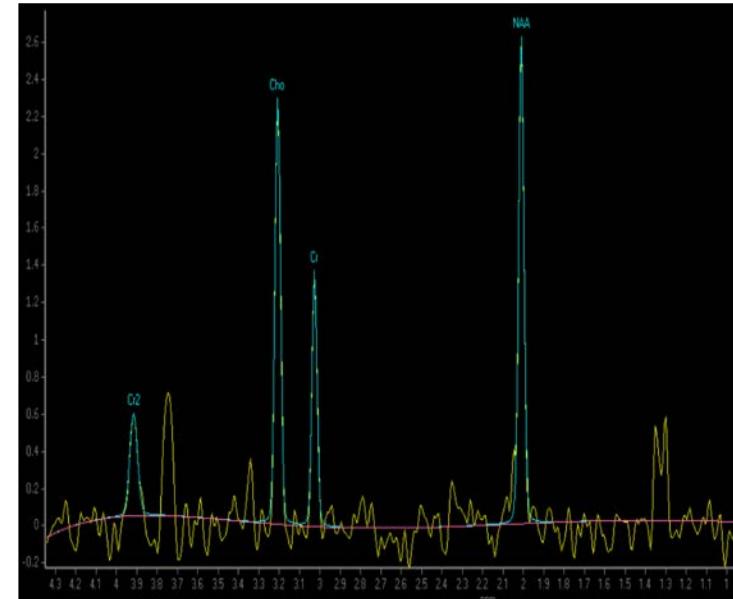
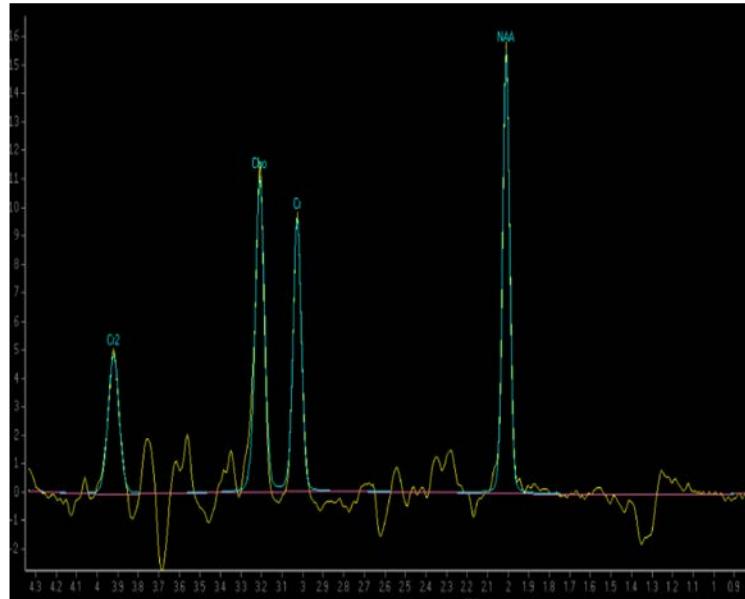


III MR SPECTROSCOPY

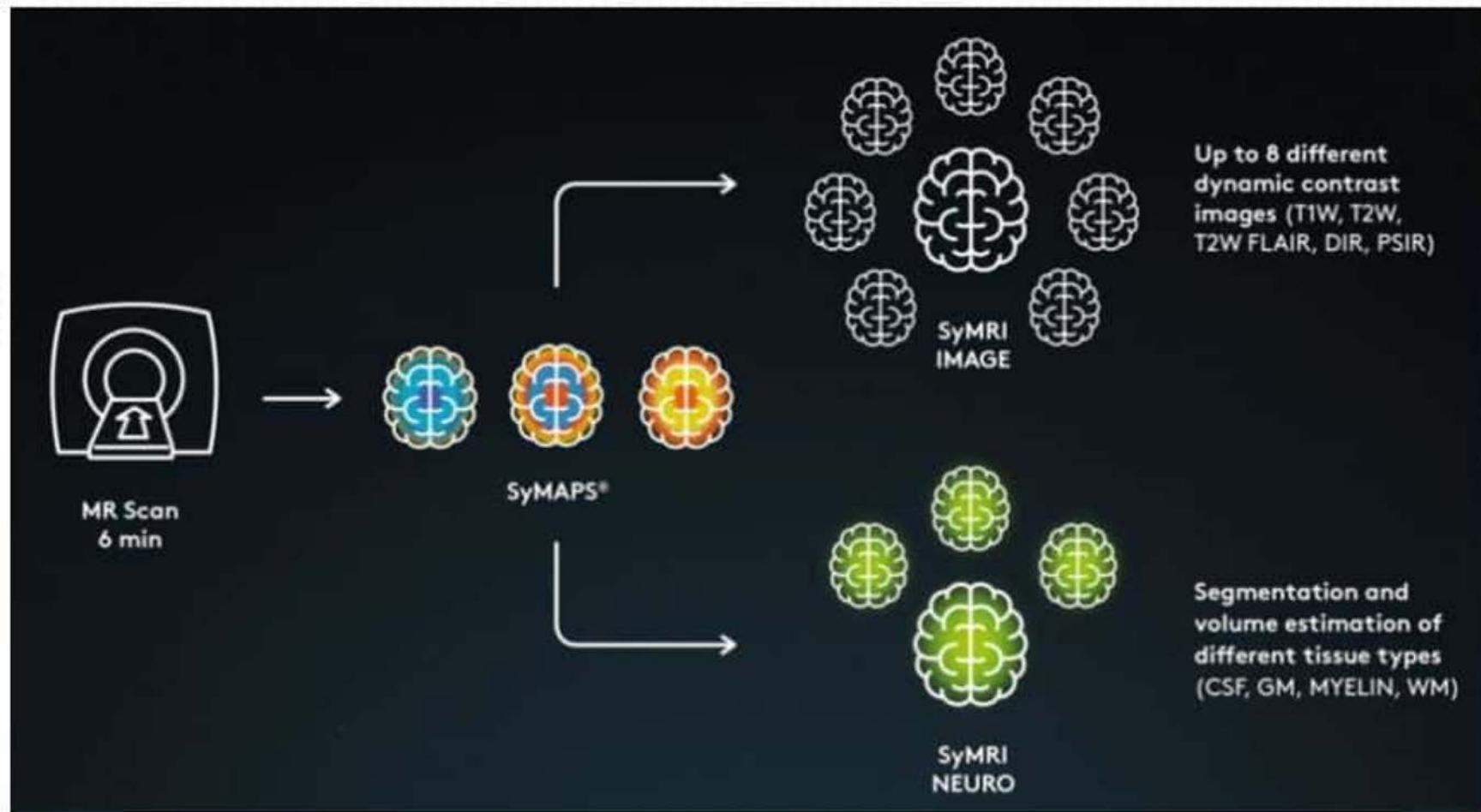
Hypoxia – mitochondrial white matter diseased : lactate ++

Canavan Leukodystrophy : NA +++++

Non-ketotische hyperglycémie : Glycine (short echo time)



III SYNTHETIC MRI



How SyMRI Works

III ADVANTAGES OF SYNTHETIC MRI

- **Quantitative measurements**, physical tissue properties => Easier to compare between different scanners/centers/references.
- **Parameters** : TR, TE, TI can be adjusted after the scan => synthetic T1, T2, STIR, PSIR, Proton density images.
- **Synthetic MRI Volumetry**
 - **Brain volume** at term age in prematurity is correlated with outcome later in life (cognitive, motor)
 - **Brain volumetry** in neonates currently only in specialized research centers
 - Techniques for volumetry in adults (freesurfer, SPM, FSL, Icometrx,...) do not work in neonates because of totally different contrast
 - >**Volume measurement** of the brain in routine with SyMRI, in less than 1 minute!
 - >**Based on known tissue specific T1 and T2 relaxation times**
 - >**No atlas needed**

III SYNTHETIC MRI - MYELINE

C. Andica et al. / Journal of Neuroradiology 46 (2019) 268–275

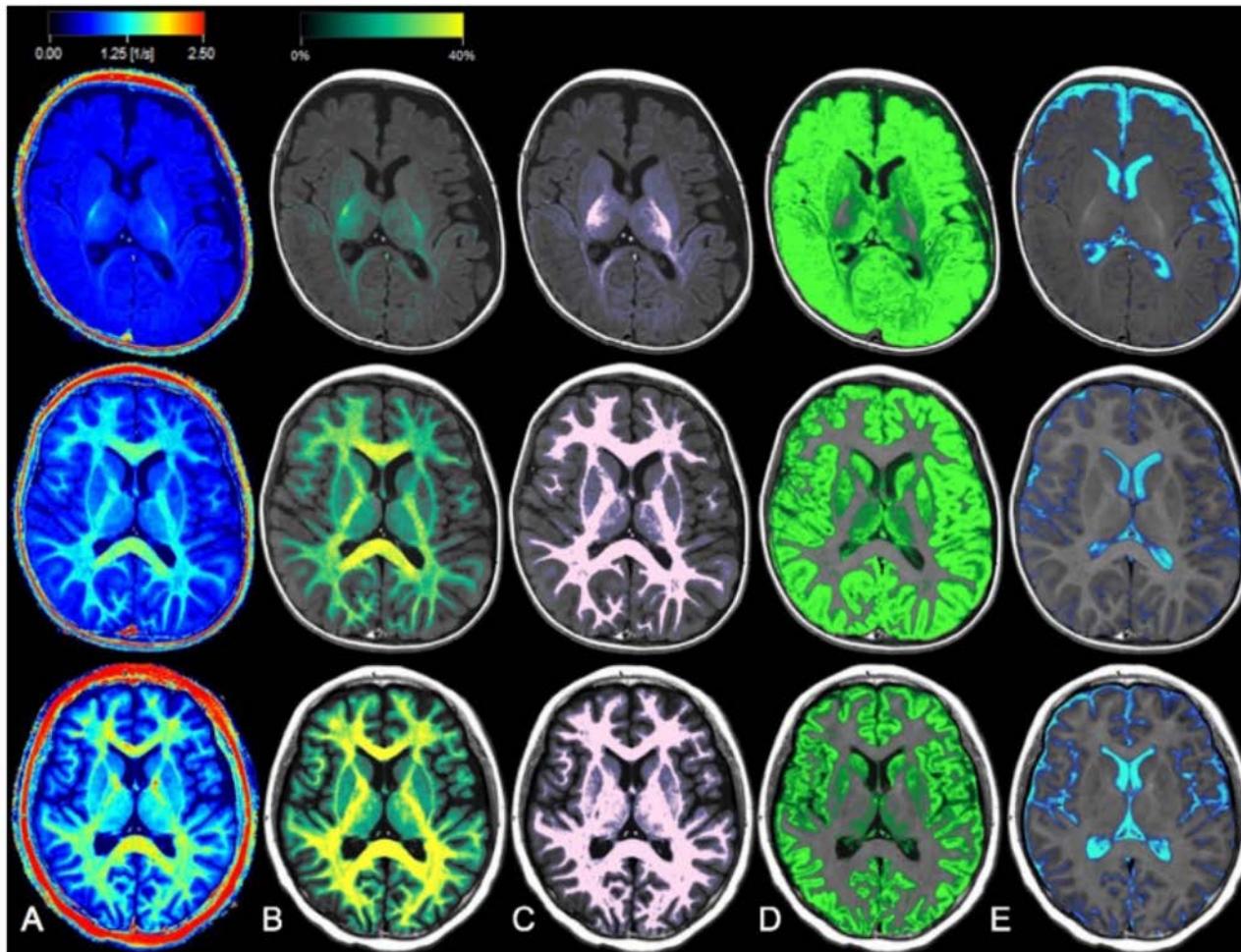
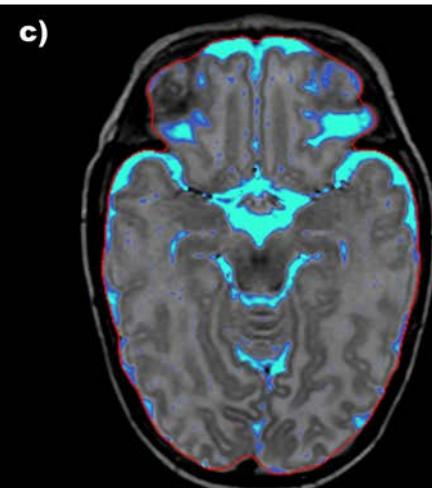
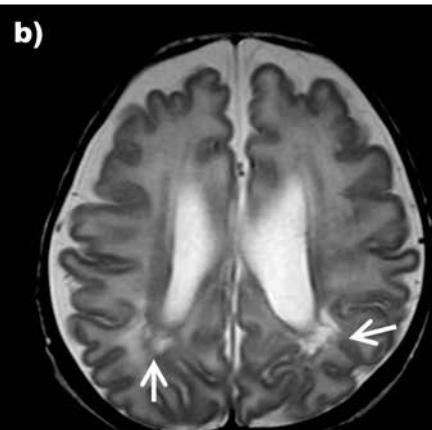
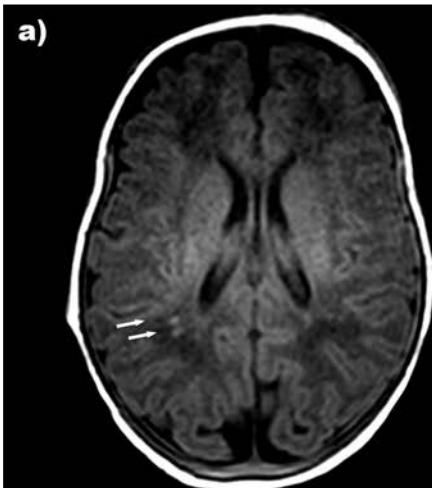


Fig. 1. The brains of a **3-month-old** male infant (first row), a **2-year-old** boy (second row), and a **13-year-old** girl (third row). SyMRI quantitative maps [only R1 map (shown on a scale 0–2.5 s⁻¹) is shown (A)], myelin partial volume map (shown on a scale 0–40%) (B), WM segmentation (C), GM segmentation (D), and CSF segmentation (E) can be used to evaluate development of the brain. However, in young children, WM and GM segmentation is suboptimal because the unmyelinated WM is assigned as GM, which is a limitation of the currently implemented algorithm

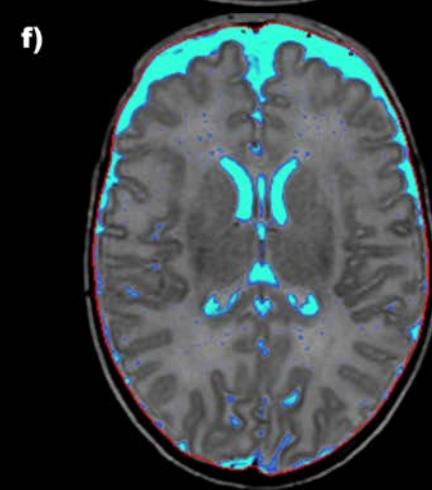
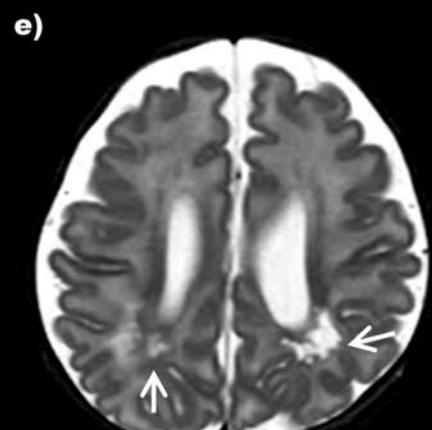
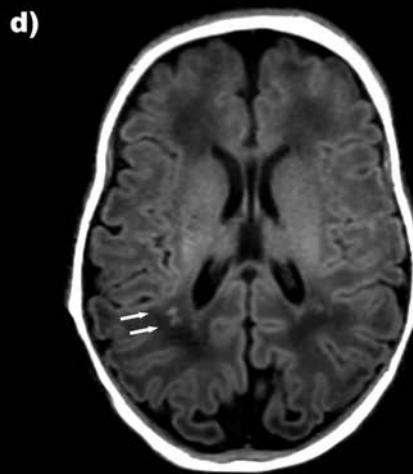
III SYNTHETIC MRI

Volumetry - SyMRI

Conv

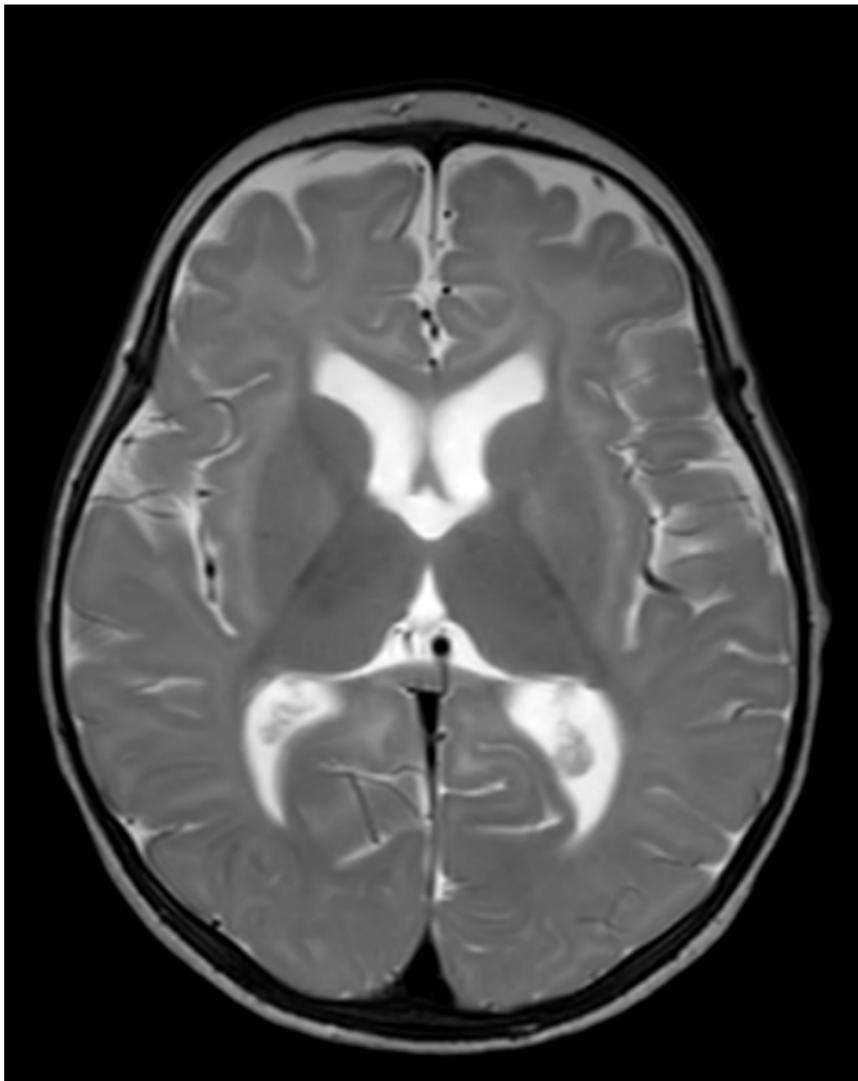


Syntetisch

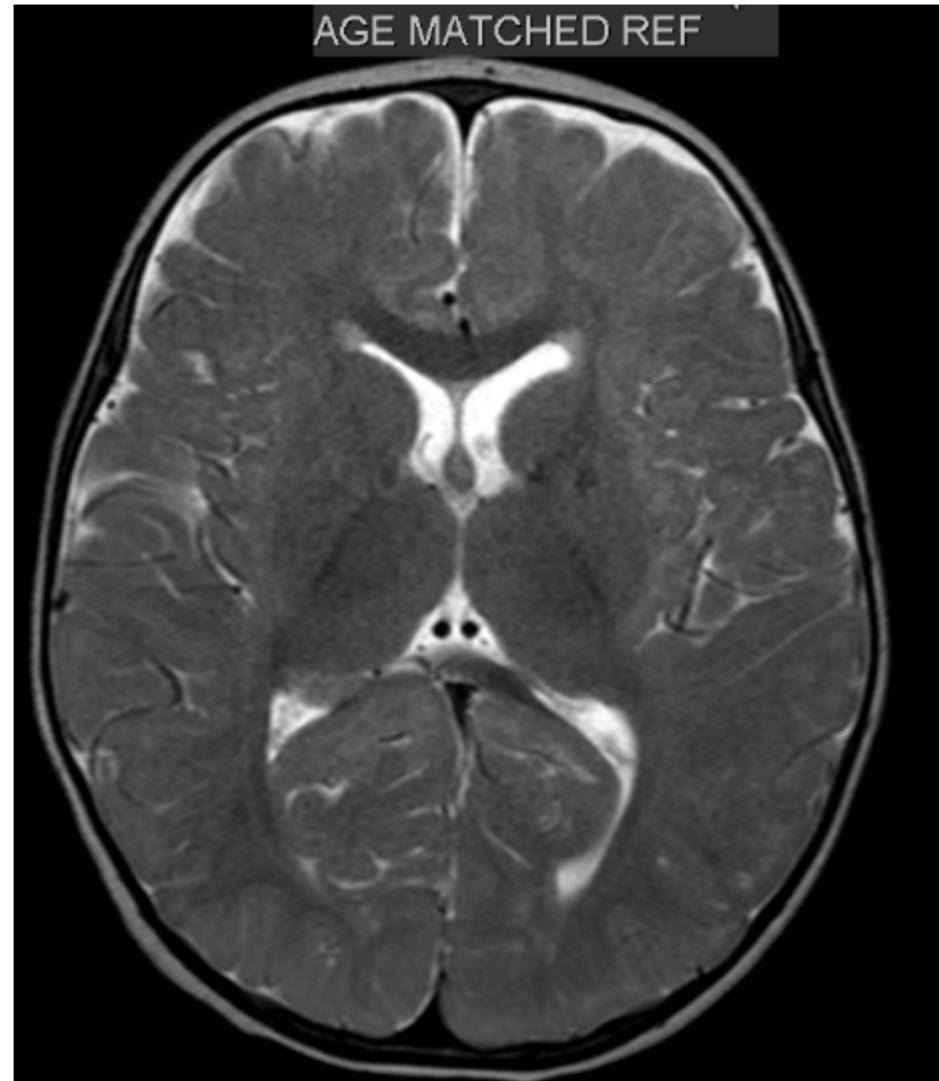




NORMAL OR NOT? COMPARATIVE IMAGES!!!

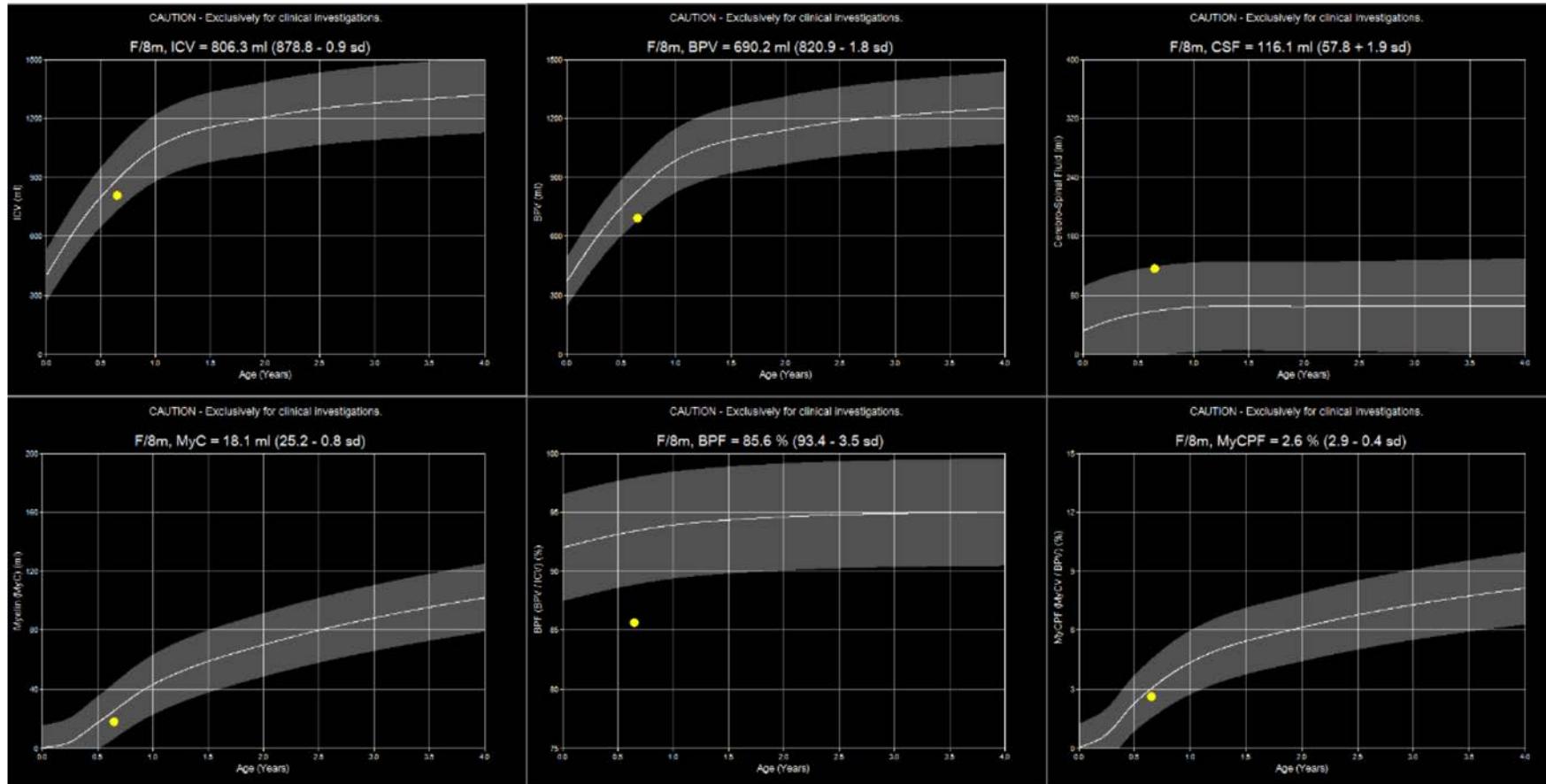


Girl, 8 months old



AGE MATCHED REF

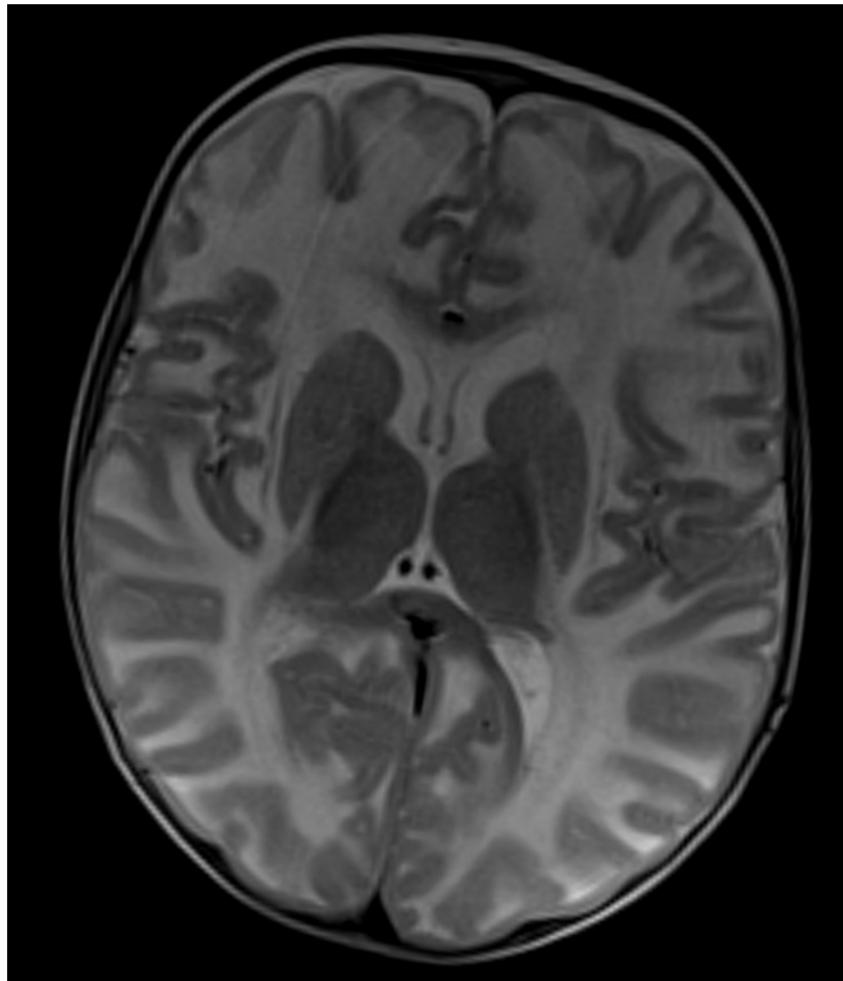
III NORMAL OR NOT? AGE MATCHED REFERENCES



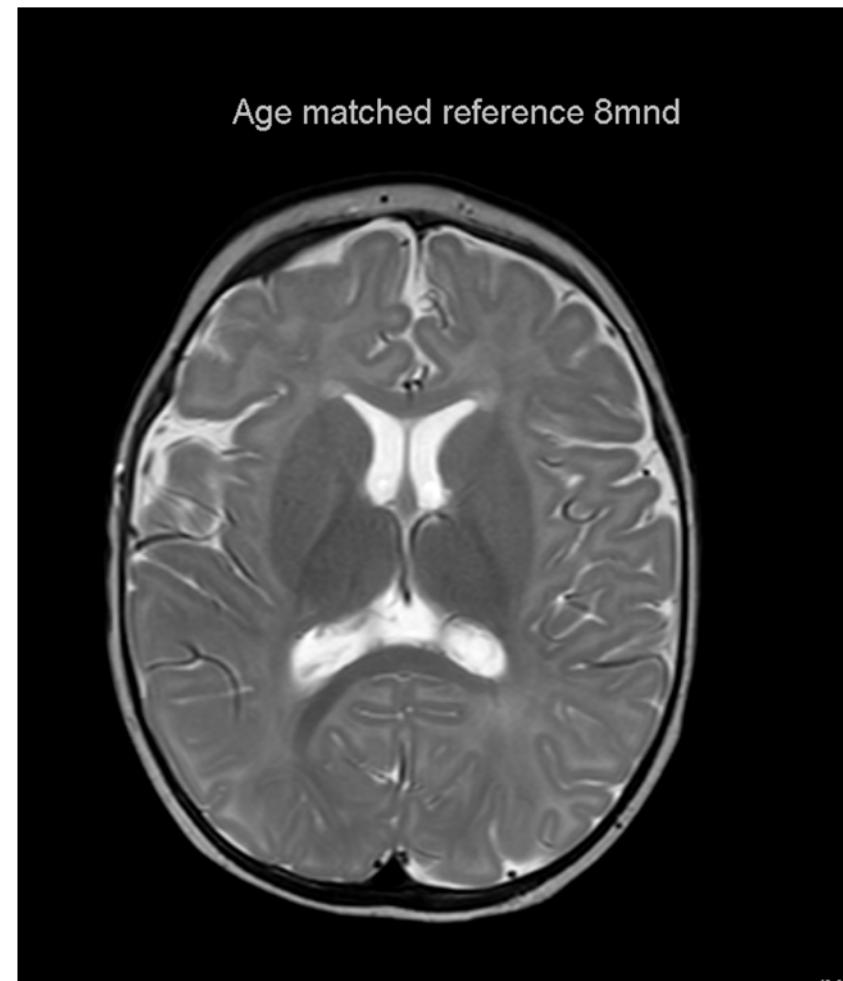
Synthetic MRI – Reference curves



NORMAL OR NOT? COMPARATIVE IMAGES!!!



Boy, 8 months old , GLIACAM mutation (megalencephalic leukoencephalopathy with cysts)



Age matched reference 8mnd

III CONCLUSION

- MRI in neonates and young children is a challenge!
- Custom protocols with adjusted FOV, TE and TR are needed.
- Good sedation is important for image quality
 - ➔ Chloralhydrate syrup or “feed and sleep”
 - ➔ Good collaboration needed with pediatricians/day hospitalists/technologists/radiologists
- Myelination proceeds according to fixed pattern and has a major influence on tissue contrast
- Synthetic MRI supplement/alternative to conventional images
 - ➔ Quantitative, brain volumetry, synthetic image reconstructi
- Importance of reference images and values



THANK YOU



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