Prenatal Diagnosis of Central Nervous System (CNS) Pathologies: does Fetal MRI help in their management?









Indications: ACOG Recommendations

#### **Common Indications:**

elevated BMI

Oligo/ Anhydramnios

Scarring of the abdomen

Position of the fetus that allows only restricted

**US** assesment

## **Special Indications:**

Situations, where MRI allows a better estimation of the intrauterine situation than US alone

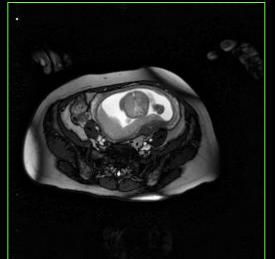
## Increased BMI + Anhydramnios

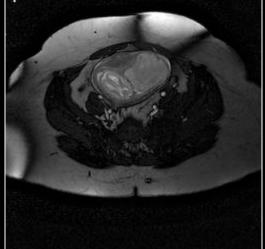




GW 19+1











#### Definitely indicated (>48%):

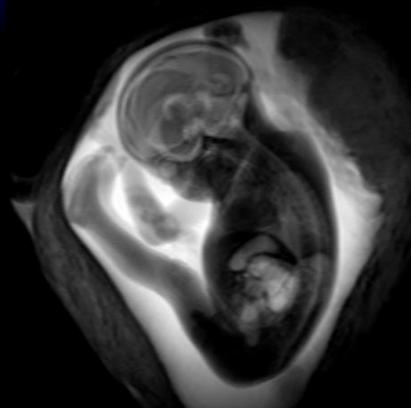
Callosal agenesis
Posterior fossa anomalies
Microcephaly

#### <u>Indicated (30-48%):</u>

Ventriculomegaly
Neural tube defects
Diaphragmatic Hernia

#### Low priority (10-30%):

Pulmonary anomalies, Multiple malformations Abdominal wall defects



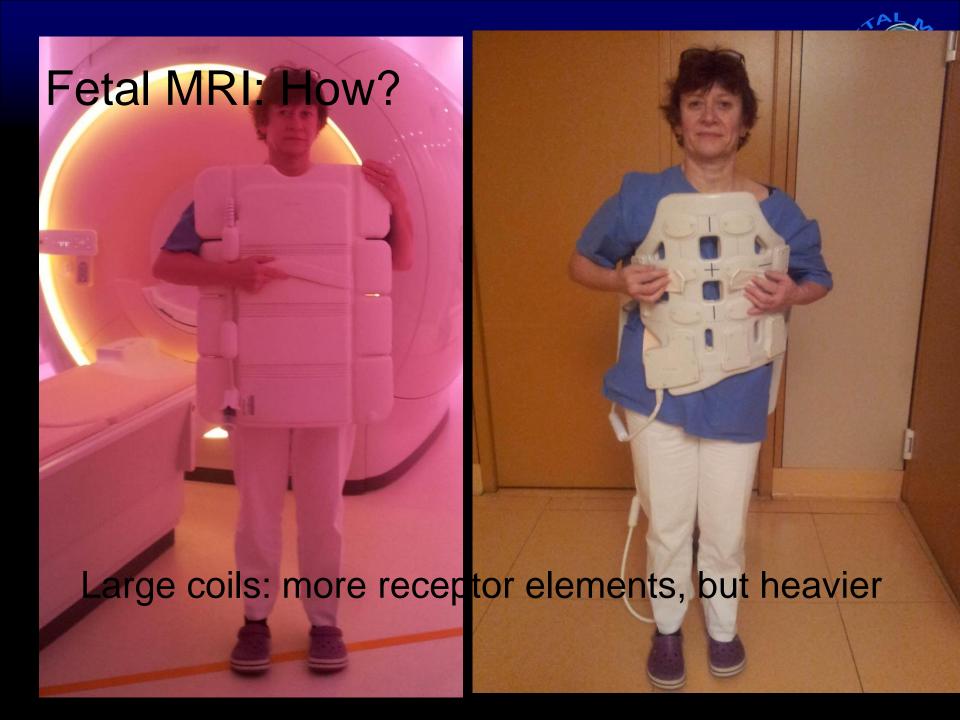
Very Low priority (0-10%): Congenital heart defects, Urinary tract, Twins, Cleft lip

## Fetal MRI: How?

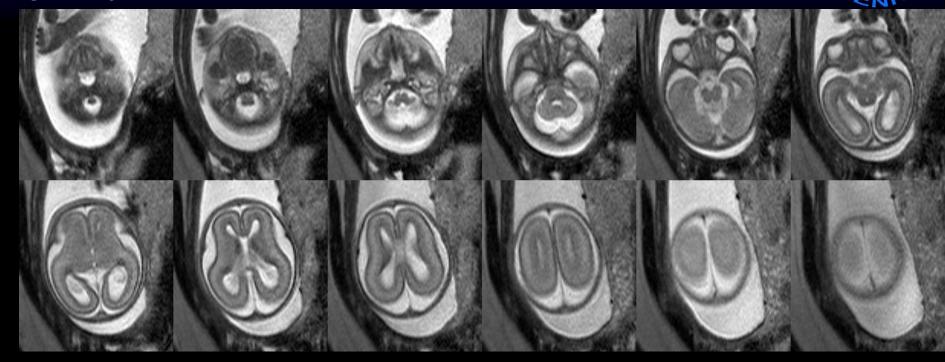


Modern 1.5 or 3T, big bores – less claustrophobia





## Fetal MRI: How? GW 20+4

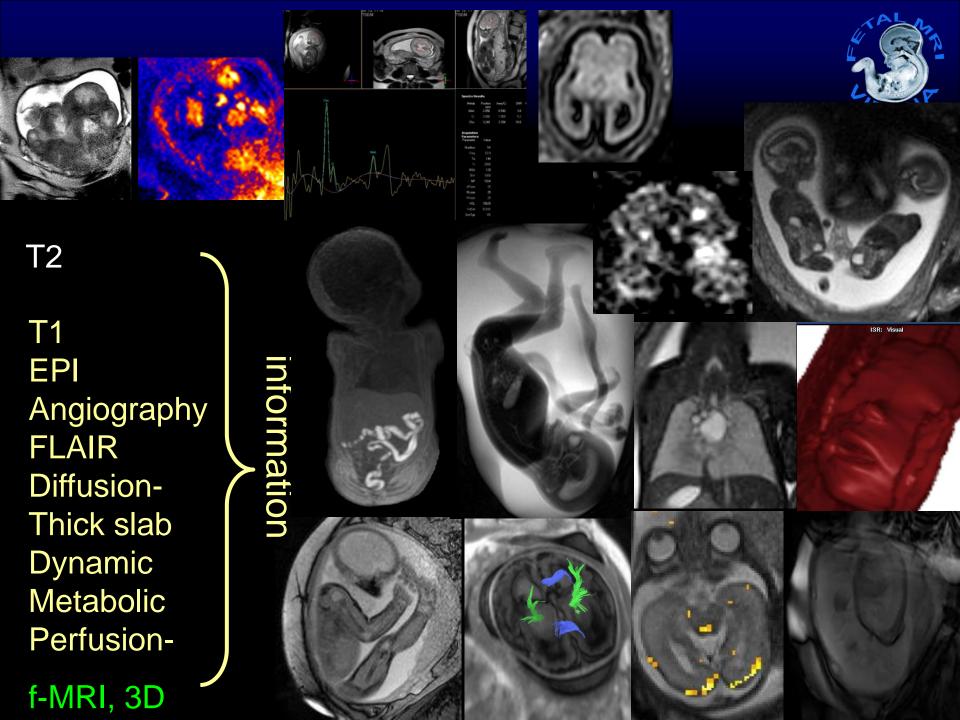


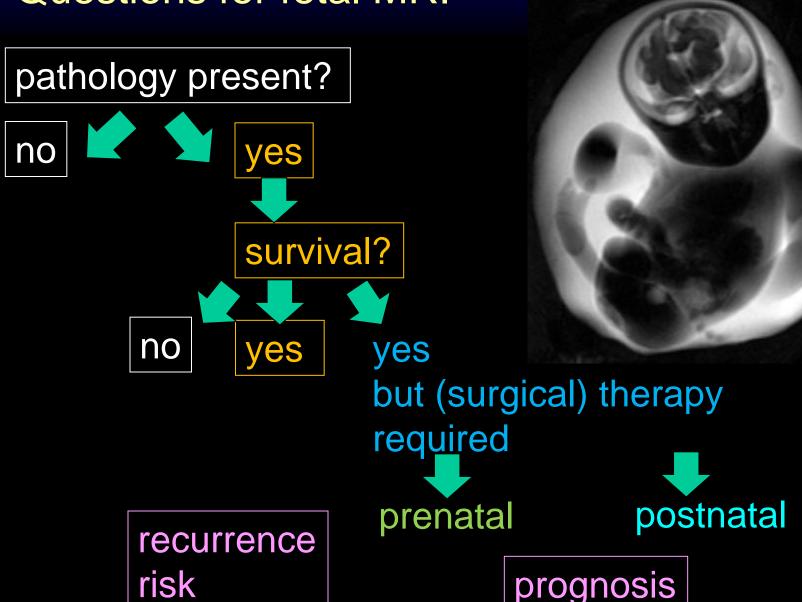
## Sequence

= Choice of parameters influencing signals, resolution...

Continuous series of images In 20 sec









pathology present?

Frequent question: isolated ventriculomegaly on Ultrasound

		Ventricle in mm
	Normal	<10mm
Classification	Borderline	8.5mm – 10mm
	Mild VM	10mm – 15mm
	Severe VM	>15mm

Pagani G, Thilaganathan B, Prefumo F. Neurodevelopmental outcome in isolated mild fetal ventriculomegaly: systematic review and meta-analysis. UOG. 2014;44(3):254-60.



pathology present?

Premature gyri



GW 24 healthy GW 23+5



Cobble stone Liss encephaly

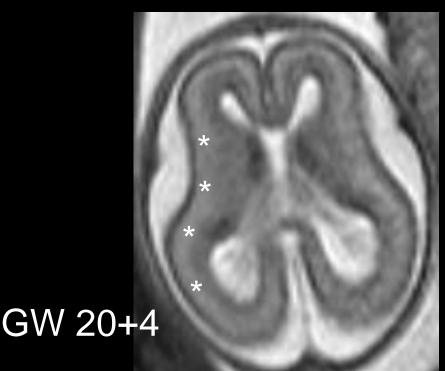
GW 29+3



pathology present?

no

MRI does not only show the surface of the brain but also the developing parenchyma



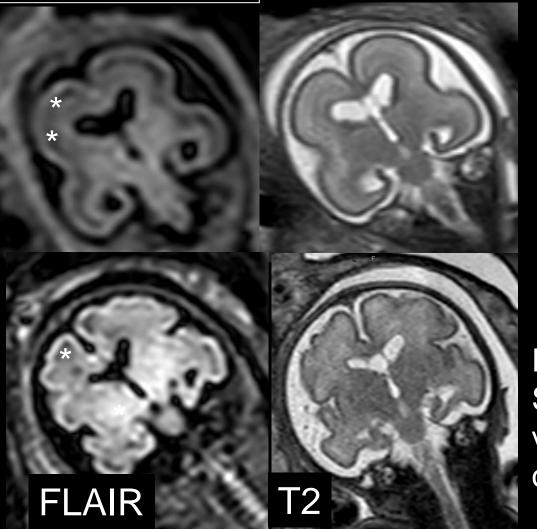
Most important structure: subplate (\*): integrity crucial for normal cortical devlopment

Kostovic I: The Anatomical Record 267;1-6 (2002)



## pathology present?

no



**GW 23** 

**GW** 29

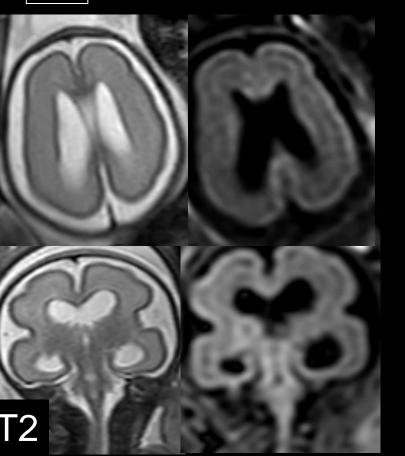
From GW 24 onwards Subplate (\*) better visible on FLAIR than on T2



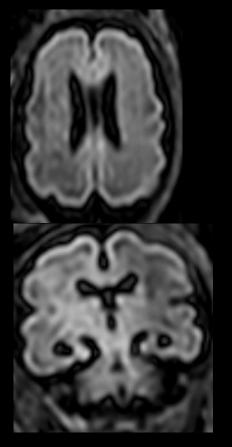
## pathology present?

no

**GW 26** 



**GW 31** 

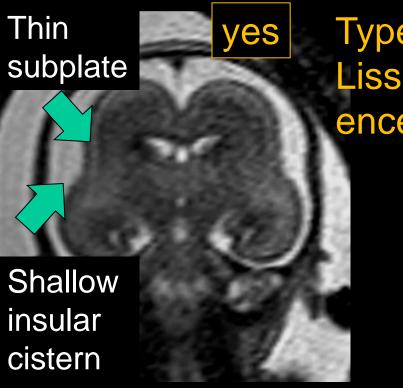


Borderline Ventriculomegaly delineation of subplate normal!

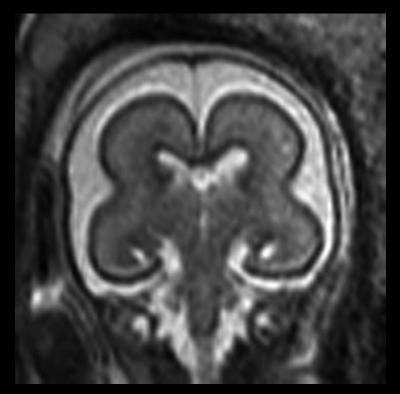


## pathology present?

**GW 22** 



Type I Liss encephaly



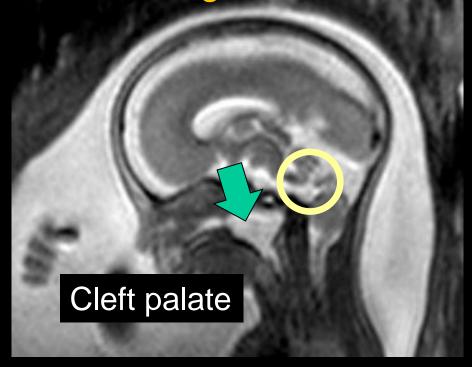
GW 20 normal



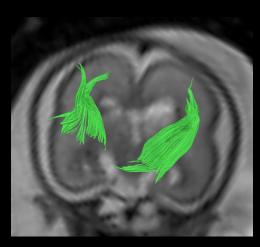
survival?

no

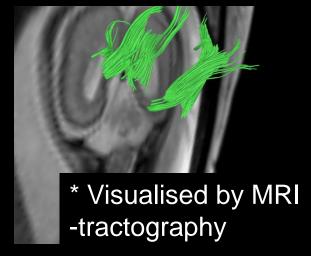
Brainstem segmentation disorder



GW 22 Brainstem interrupted



Corticospinal tract absent Infratentorially\*

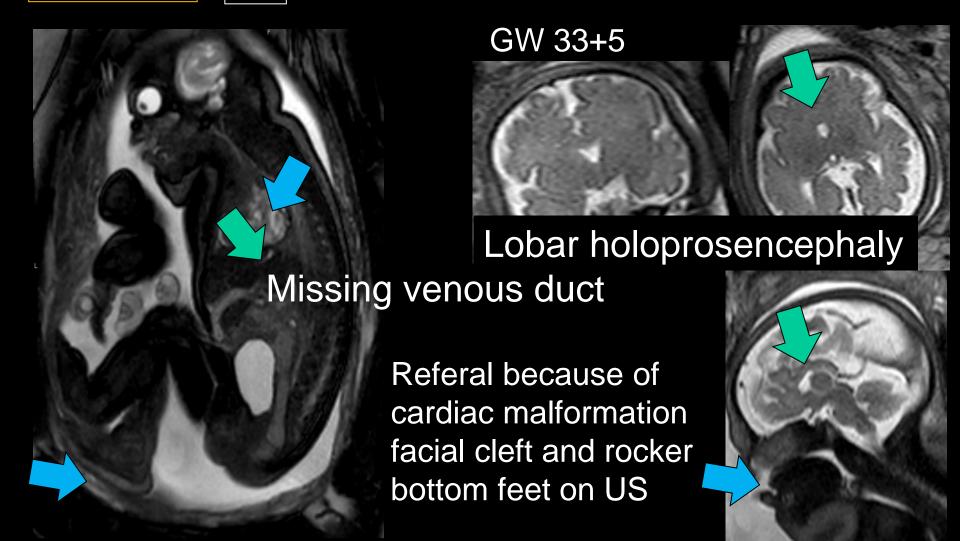




survival?

no

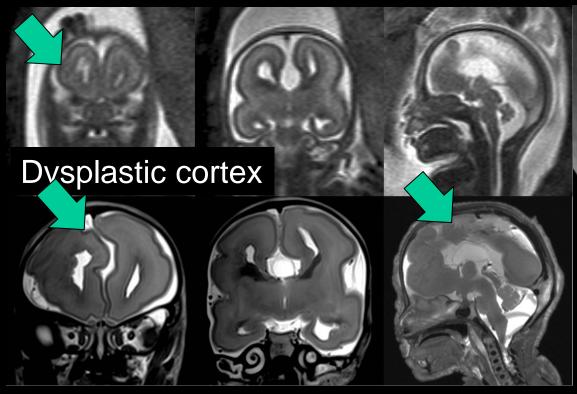
Trisomy 13



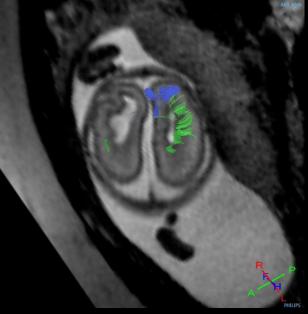


survival?

Yes. Females only...but....







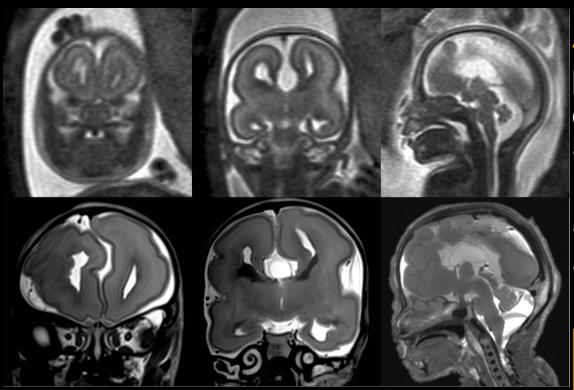
Corpus callosum agenesis with Probst Bundles\*





survival?

Yes. Females only...but....



Aicardi syndrome presumably X-linked dominant,

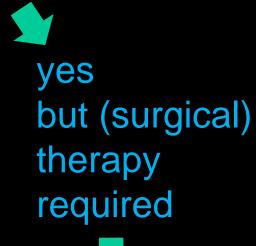
- agenesis of the corpus callosum,
- chorioretinal lacunae
- infantile spasms, with lethality in males.

Seizures severe neurological impairment....

Fernández-Ramos JA et al. Rev Neurol. 2013 Dec 1; (11):481-8.

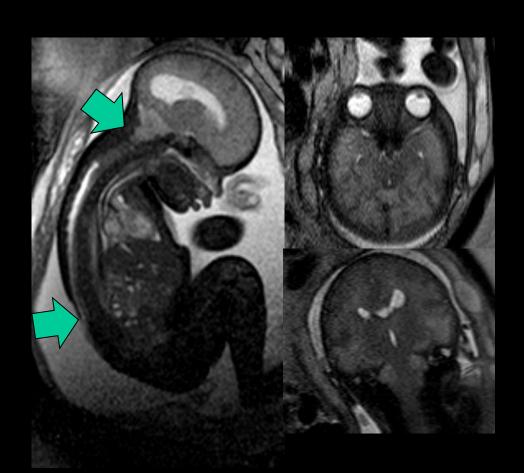


#### survival?



prenatal

Fetuses with open dystaphism improve after prenatal closure of the cele



GW 32, Chiari II malformation



## survival?

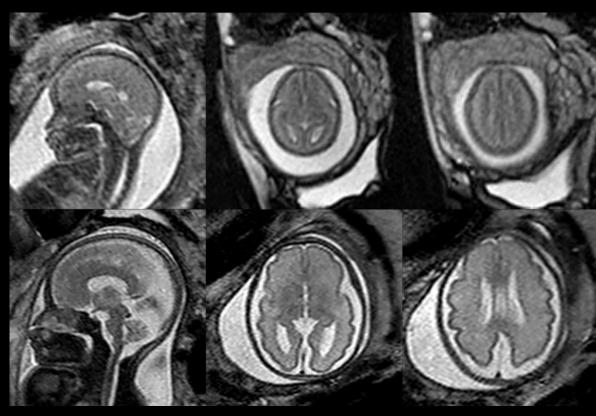


yes but (surgical) therapy required

prenatal

Fetuses with open dysraphism improve after prenatal closure of the cele

GW 23 before



GW 28 after

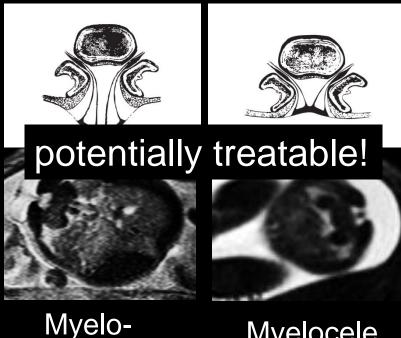
Courtesy Gregor Kasprian Vienna/ Houston

## Questions for fetal MRI "Spina bifida"



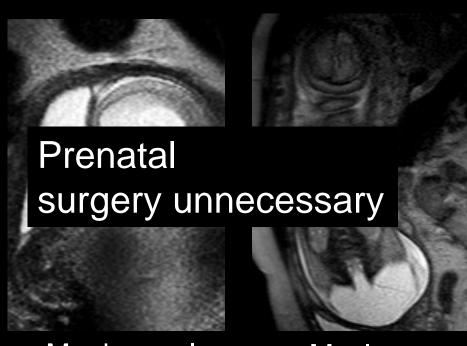
open neural tube defects <sup>1</sup>

closed neural tube defects 1



meningocele

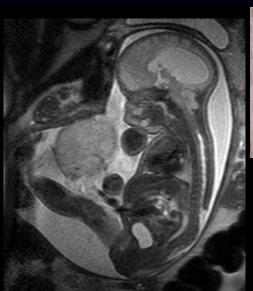




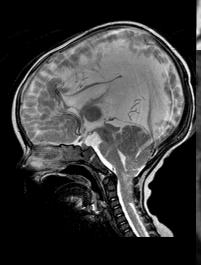
Meningocele Myelocystocele

1: Tortori-Donati P, Rossi AMD, Biancheri R. Pediatric neuroradiology. Berlin; [Great Britain]: Springer 2005.













GW 29+4





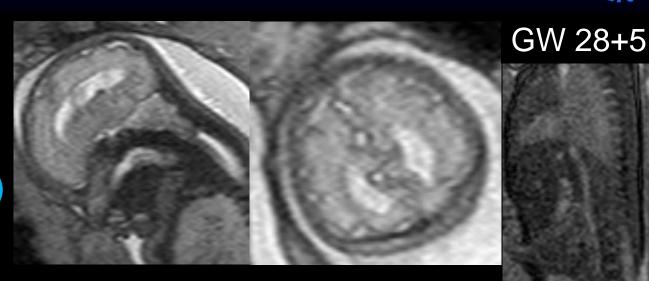
#### survival?

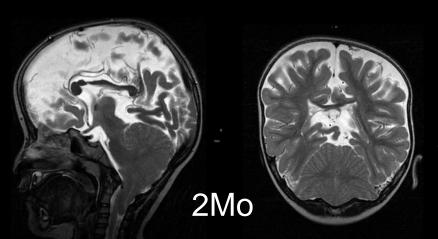


yes but (surgical) therapy required



Postnatal shunt with less impact on vermian herniation





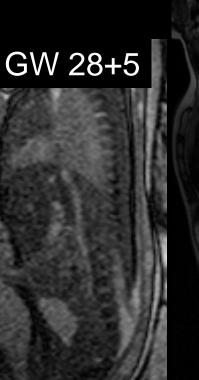
Danzer E et al. Dev Med Child Neurol. 2012;54(1):8-14.

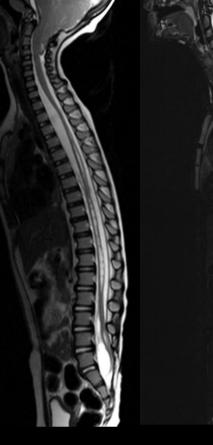


## survival?



yes
but (surgical)
therapy
required







Late spinal complications

postnatal

3a

12a



recurrence risk

# 2 month old with seizures

Partial Callosal Agenesis?

Callosal Dysgenesis?

Schizencephalic clefts?





recurrence risk

Fetal Thrombotic Vasculopathy

GW 26

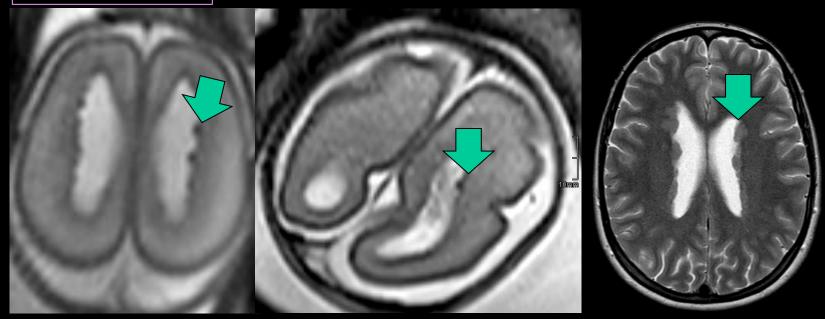
**Demised Co-Twin** 

Sato Y, Benirschke K. Pediatrics. 2006;117(1):113-7.



recurrence risk

Subependymal heterotopia



1<sup>st</sup> Fetus GW23+0 2<sup>nd</sup> Fetus GW21+3

mother

Filamin A gene mutation!



prognosis | Spinal level in open defects?







Hipflection: Kneeextgension: L1/2 L3

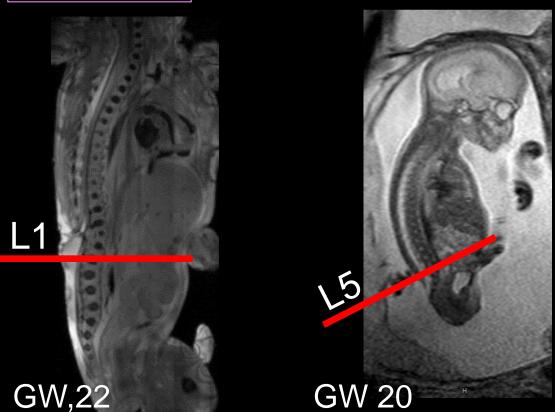
Kneeflection: L4 Foot dorsiflection: L5

Foot plantarflection: sacral

Lindseth RE. (1976) Treatment of the lower extremity in children paralyzed by myelomeningocele (birth to 18 months). AAOSIC Lectures 25: 76–82.



prognosis | Spinal level in open defects?





assessments correlated ± 1 Level of anatomical defect

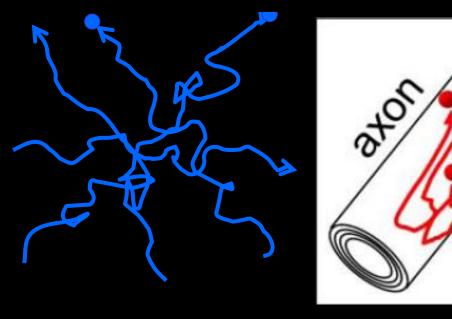


## prognosis

Use of diffusion- tensor imaging



Diffusion weighted imaging measures degree and directionality of water motion in tissue



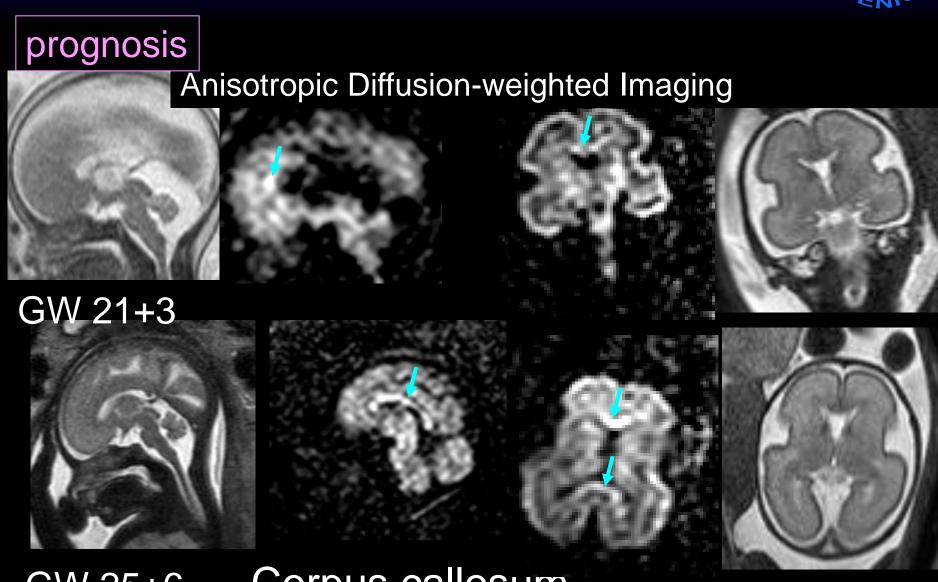
Stochastic motion Brownian motion

Isotropy

Anisotropy







GW 25+6

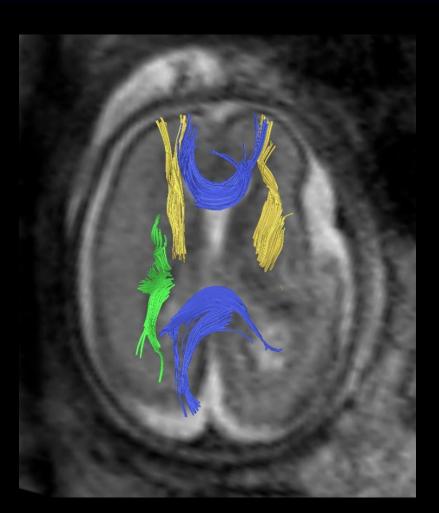
Corpus callosum





prognosis



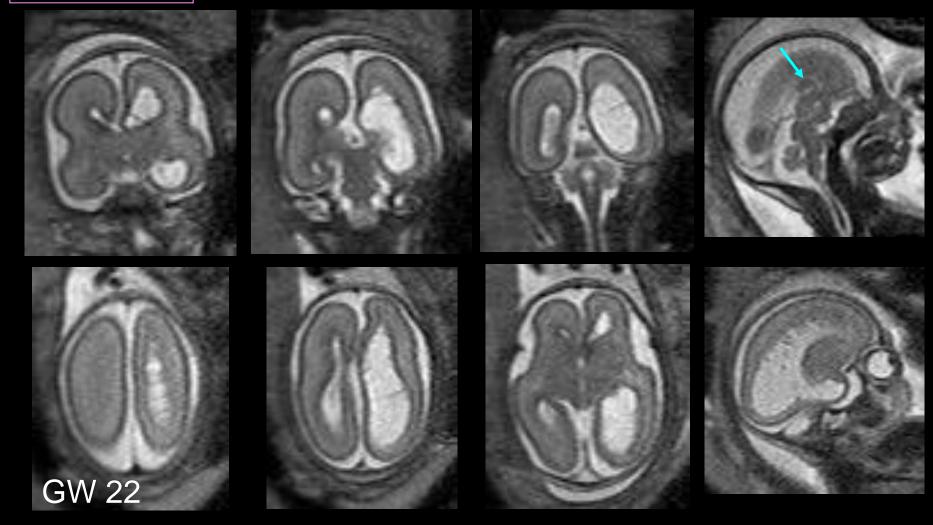


Connectivity at GW 23

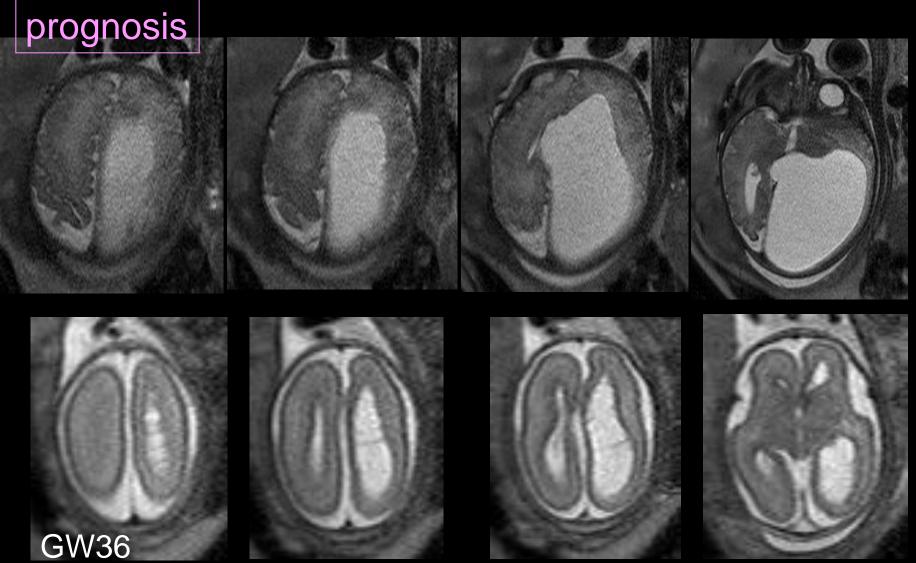


## prognosis

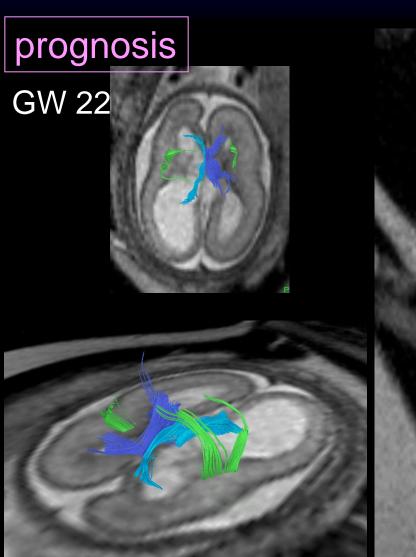
## Callosal Hypogenesis

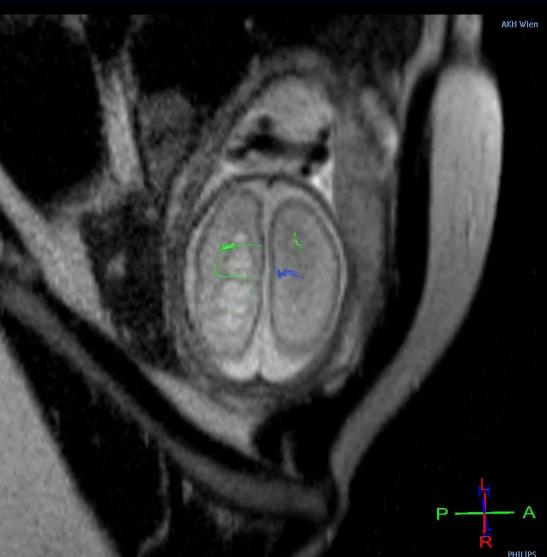












Questions for fetal MRI prognosis 5a **GW 22** 

prognosis

## At 5 Years

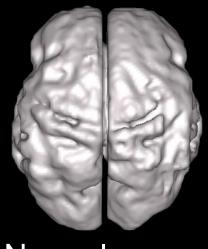
Normal intelligence
Symmetrical use of arms
Epilepsy with rare seizures,
EEG focus left
hemisphere





## Take home:

Fetal MRI can help with more accurate prenatal diagnosis and prognosis and thus support the managment of a complicated pregnancy



Normal



CC agnesis



CC agenesis with malformations



#### 27th World Congress on Ultrasound in Obstetrics and Gynecology

16 – 19 September 2017, Vienna, Austria



