Development of the Fetal Cerebral Cortex in the Second Trimester: Assessment with 7T Postmortem MR Imaging

Z. Zhang, Z. Hou, X. Lin, G. Teng, H. Meng, F. Zang, F. Fang, and S. Liu

ABSTRACT

BACKGROUND AND PURPOSE: Few investigators have analyzed the fetal cerebral cortex with MR imaging of high magnetic strength. Our purpose was to document the sulcal development and obtain quantitative measurements of the fetal brain in the second trimester.

MATERIALS AND METHODS: The brains of 69 fetal specimens, with GA 12–22 weeks, were first scanned on a 7T MR imaging scanner. Then the sequential development of the different fissures and sulci was analyzed, and quantitative measurements of the cerebral cortex were obtained.

RESULTS: A new chronology of sulcal development during 12–22 weeks GA was summarized. Before 12 weeks, few sulci were present; by 16 weeks, many sulci were present. The 16th week could be considered the most intensive time point for sulcal emergence. Most sulci, except for the postcentral sulcus and intraparietal sulcus, were present by 22 weeks GA. Measurements of the fetal brains, each with different growth rates, linearly increased with GA, but no sexual dimorphisms or cerebral asymmetries were detected.

CONCLUSIONS: The second trimester is the most important phase, during which most sulci are present and can be clearly shown on 7T postmortem MR imaging. It is apparent that the specific time during which neuropathologic features of sulci appear, previously thought to be well understood, should be redefined. Quantitative data provide assistance in the precise understanding of the immature brain. The present results are valuable in anatomic education, research, and assessment of normal brain development in the uterus.

ABBREVIATIONS: GA = gestational age; US = ultrasonography

The developmental process of the fetus in vivo is divided into the first, second, and third trimesters, which are closely associated with the different developmental stages of the CNS. The first and second trimesters correspond to the neurulation, differentiation of cerebral vesicles, and neurogenesis.1-3 Disorders of migration are more likely to occur in the second trimester. Either undermigration or overmigration of neurons will lead to cortical abnormalities, such as gray matter heterotopia, agryria/pachygyria, heterotopia within the molecular layer (layer 1 of the cortex), and neuronal heterotopia.4 Therefore, it is important to study fetal brain development in the second trimester.

The sulci first appear as shallow fossa and then develop into a deeper and more curved pattern on the cerebral cortex.5,6 The timing of the appearance of these different types of sulci is so precise that neuropathologists consider sulcation to be a reliable estimation of GA and consequently a good marker of fetal brain maturation.6 Fetal sulcal development has been studied in neuropathology,5 with US7 and MR imaging in vitro8 and in vivo.6 However, few investigators have described the normal patterns of fetal sulcation with an MR imaging scanner of high magnetic strength.

The dimensions of the developing fetal brain change rapidly in early life, and morphometric normative data are crucial for the assessment of normal maturation and diagnosis of brain anomalies.9 Currently, prenatal morphometric studies have been focused as early as 17 weeks GA10, but only the linear biometric values were obtained on 2D images without 3D reconstruction of the fetal brain. In addition, the existing MR imaging postprocess-
The brains were not removed from the calvaria and were scanned by a 7T micro-MR with a maximal gradient of 360 mT (70/16 PharmaScan; Bruker BioSpin, Bremen, Germany). A rat body coil with an inner diameter of 60 mm was selected to scan all of the fetuses. For T1-weighted images, section thickness was 0.8 mm; section interval, 0.8 mm; TR, 384.4 ms; TE, 15.8 ms; matrix size, 256 × 256; number of excitations, 4; and field of view, 6 × 6 cm. For T2-weighted images, section thickness was 0.5 mm; section interval, 0.5 mm; TR, 17,000 ms; TE, 50 ms; matrix size, 256 × 256; number of excitations, 4; and field of view, 6 × 6 cm.

| GA     | Number |
|--------|--------|
| 12     | 3      |
| 13     | 3      |
| 14     | 3      |
| 15     | 4      |
| 16     | 5      |
| 17     | 7      |
| 18     | 4      |
| 19     | 6      |
| 20     | 16     |
| 21     | 12     |
| 22     | 6      |

### Materials and Methods

#### Subjects

A total of 85 fetal specimens of 12–22 weeks GA were obtained from medically indicated or spontaneous abortions, and fetal deaths from hospitals in the Shandong Province of China.

The inclusion criteria were the same as those in our previous research: 1) maternal pregnancy records showing an absence of documented fetal chromosomal abnormalities, stressful intrauterine conditions, maternal genetic disease in the families, or a history of seizures in the case of eclampsia; 2) results of US examination for the fetus during pregnancy and results of postmortem MR imaging examinations for the specimen indicating an anatomically normal and developmentally appropriate fetal CNS; and 3) further validated detailed autopsy combined with neuropathologic examinations also describing no detectable CNS malformations.

Finally, 69 specimens with an appropriately developed fetal brain were reserved for the study (GA dispositions and numbers of the chosen specimens are listed in Table 1). This study was conducted after approval of the Ethical Committee at Shandong University.

#### Image Acquisition

The brain structures were assigned and annotated based on a histology atlas of second-trimester fetal brains. The T2-weighted images were segmented and reconstructed by Amira 4.1 software (www.amira.com). The segmentation on the cortical surface was performed at the borderline between the cortical plate and the marginal zone (Fig 1A, C). The 3D reconstruction models were automatically obtained after segmentation. We simultaneously segmented all images twice using 2 anatomists to obtain a mean.

After segmentation, a polygonal surface model was created. The Amira software was used to generate a triangular surface grid for the brain embedded in a voxel dataset. Then the surface area for each triangular grid was summed to obtain the surface area of the whole brain. A brain mask for the 3D volume was created by segmentation. All of the voxels in the mask were accumulated to obtain brain volume, and a linear interpolation method was used to get the voxels in the gap between 2 sections.

Brain length, width, and height were also calculated. Brain height was defined as the vertical distance between 2 horizontal planes, containing the most superior and inferior points of the brain, respectively. Brain length and width were defined in a similar fashion (Fig 1D).

### Statistical Analysis

Our criterion to define a sulcus is the same as that used by Garel et al. A clear indentation at the surface of the brain was considered to be the earliest indication of a sulcus. One sulcus was considered to be present if it was observed in more than 75% of cases, detectable if observed in 25% to 75% of cases, and absent if observed in less than 25% of cases. A paired t test was used to detect significant differences in sex and hemispheres. All of the statistical work was done with SPSS version 17.0 (SPSS, Chicago, Illinois).

### Results

Of the 69 specimens, 31 were male and 38 were female. All were from the Chinese Han ethnic group with an average age of 18.348 ± 2.869 weeks GA. The GA span of male specimens was 13–22 weeks (mean ± SD, 18.129 ± 2.592 weeks GA), and the GA span of female specimens was 12–22 weeks (mean ± SD, 18.526 ± 3.100 weeks GA). There were no significant sex differences in GA (P = .571). There was no statistically significant interobserver variation (P = .810) at preliminary analysis.

### Sulci

The chronology of sulcal development during 12–22 weeks GA is listed in Table 2. At 12 weeks GA, few sulci except the interhemispheric fissure and the lateral sulcus were present (Fig 2A, -E). By 16 weeks GA, many sulci were present, such as the central sulcus, the superior frontal sulcus, calcarine fissure, and parieto-occipital sulcus (Fig 2B, -F, and -J), and the 16th week GA could be considered to be the most intensive point of sulcal emergence. At 20 weeks GA, the present sulci became a little deeper and more obvious on the cortical surface (Fig 2C, -G, and -L). The collateral sulcus and the posterior part of the superior temporal sulcus

---

**Table 1: GA dispositions and numbers of chosen specimens (n = 69)**

| GA     | Number |
|--------|--------|
| 12     | 3      |
| 13     | 3      |
| 14     | 3      |
| 15     | 4      |
| 16     | 5      |
| 17     | 7      |
| 18     | 4      |
| 19     | 6      |
| 20     | 16     |
| 21     | 12     |
| 22     | 6      |
emerged as well. Most sulci, except the postcentral sulcus and intraparietal sulcus, were not present until 22 weeks GA (Fig 2 D, H). At this point, all of the visible sulci were straight, immature, and had not developed the secondary branches (Fig 2). In addition, most appeared deeper and more distinguishable at the right hemisphere, especially the superior frontal sulcus and the central sulcus (Fig 2 B, D).

The lateral sulcus, which was the first to be distinguished, was the most obvious of all of the sulci. During 12–22 weeks GA, the lateral sulcus became deeper and wider, with the opercula insulae approaching each other. However, the opercula insulae did not fold until 22 weeks GA (Fig 2 E–H).

The calcarine fissure and the parieto-occipital sulcus were the most prominent in the medial cerebral surface. The emergence time of the calcarine fissure was a little earlier than that of the parieto-occipital sulcus, and they gradually became separate by 20 weeks GA (Fig 2 I–L).

The appearance of all of the other sulci, such as the central sulcus, the posterior part of the superior temporal sulcus, and the superior frontal sulcus, did not change much and remained shallow and tiny before 22 weeks GA (Fig 2).

**3D Visualization Model of the Fetal Brain**

The 3D visualization model demonstrates well the morphologic changes of the cortical surface (Fig 3). Most sulci, delineated as shallow fossae on the cerebral surface, could be observed (Fig 3), such as the lateral sulcus, central sulcus, and superior frontal sulcus. The length, depth, and location of the sulci could also be clearly demonstrated (Fig 3). The sulci were discontinuous and appeared as a cluster of separated shallow fossae, which merged together to form a single immature sulcus after 16 weeks GA (Fig 3 B–D), such as the central sulcus and inferior temporal sulcus. The lateral sulcus was described as a wide fossa on the cortical surface (Fig 3). It became deeper as GA increased, and the operculum insulae did not fold until 22 weeks GA (Fig 3 D).

**Quantitative Assessment of the Fetal Brain**

From the slope of the lines, it could be concluded that the total cerebrum was developing at a different speed. The brain surface area and volume increased the fastest (Fig 4), followed by its length (Fig 5 A). The height increased the slowest (Fig 5 C). There were no sexual dimorphisms or cerebral asymmetries in the measurements (P > .05).

**DISCUSSION**

**Sulcation in the Second Trimester**

Development of the fetal cerebral cortex is defined by changes of the sulci, because they are indicators of brain maturation. Exploring the sequential development of sulci can greatly enrich our knowledge of prenatal radiology and will supply certain guidance for assessing normal cortical development in vivo. Our results show that most of the sulci can be observed much earlier on 7T postmortem MR imaging than on in vivo fetal MR imaging. An example of this is the central sulcus, which is commonly observed on fetal MR imaging at 24–25 weeks GA, but can be clearly observed on 7T postmortem MR imaging at 16 weeks GA. Thus, we...

---

**Table 2: Chronology of sulcal development during 12–22 weeks GA**

| Sulci                              | Observed 25%–75% | Present ≥ 75% | Sulci                              | Observed 25%–75% | Present ≥ 75% |
|------------------------------------|------------------|--------------|------------------------------------|------------------|--------------|
| Medial Cerebral Surface             |                  |              | Lateral Cerebral Surface            |                  |              |
| Interhemispheric fissure           | –<sup>b</sup>    | 12           | Superior frontal sulcus            | 12               | 15           |
| Callosal sulcus                    | 12               | 14           | Inferior frontal sulcus            | 15               | 16           |
| Cingular sulcus                    | 12               | 14           | Posterior part of superior temporal sulcus | 16               | 18           |
| Calcarine fissure                  | 13               | 15           | Inferior temporal sulcus           | 15               | 16           |
| Parieto-occipital sulcus           | 15               | 16           |                                   |                  |              |
| Ventral Cerebral Surface           |                  |              | Vertex                             |                  |              |
| Hippocampic fissure                | 12               | 14           | Precentral sulcus                  | 16               | 18           |
| Orbital sulcus                     | 15               | 16           | Central sulcus                     | 15               | 16           |
| Collateral sulcus                  | 16               | 18           |                                   |                  |              |
| Occipitotemporal sulcus            | 20               | 22           |                                   |                  |              |
| Olfactory sulcus                   | 15               | 16           |                                   |                  |              |

<sup>a</sup> Sulci were present on 7T postmortem MR imaging scans.

<sup>b</sup> No cases were studied.
believe that MR imaging with higher magnetic strength can delineate small sulci earlier.

There were inconsistencies between our results and those of previous research. For example, we found the central sulcus at 16 weeks GA, but Chi et al5 found it at 20 weeks GA on neuropathologic examinations. We found that the superior frontal sulcus and inferior temporal sulcus had already been present at 16 weeks GA in almost all of the specimens, but Chi et al5 described the neuro-

pathologic appearance of the these 2 sulci at 25 weeks and 30 weeks GA, respectively. These inconsistencies need to be studied further. Genetic, environmental, nutritional, and other differences exist between our generation and past generations, as well as between white and Asian races,17 which may affect the developmental speed of the fetal brain. The emergence time of sulci, described in great detail in anatomic and embryologic books and in published studies, should be re-examined.
There are many possible reasons for the inconsistencies between our results and those described in past research, such as fluid loss of the fetal brain caused by fetal death or formalin fixation, method of describing the presence of a sulcus (being observed in more than 75% or 50% cases), method of examination (US or MR imaging of different magnetic strength), different races (Asian or Western), and other unknown reasons. It is thought that all of these variables cannot lead to such an appreciable difference between our chronology of sulcal development and that of Chi et al.

In our study, the sulci are described as a cluster of separated shallow fossae at 16 weeks GA, termed “sulcal roots” in previous research. It can be concluded that sulcal development starts from several separated fossae on the cerebral surface, these fossae then join together, and a single complete sulcus is formed. The 16th week of GA may be considered as the original point for formation of the sulcal roots.

Quantitative Measurements of the Fetal Brain in the Second Trimester

Quantitative analysis in our study may provide valuable assistance in modeling normal fetal brain development. Currently, quantitative data on the fetal brain mainly come from research on neonates or premature infants, and very little is known about the range of normal values in the first and second trimesters. Therefore, our measurements, obtained from a large cohort, may be a valuable reference to determine whether the cortex is developing normally during the midtrimester. Although the fetuses in our study have undergone formalin fixation, which slightly reduces the volume and increases the width of the sulci, previous studies have confirmed the clinical application of these measurements.

Our measurements of the surface area and volume of the brain at 13–21 weeks GA were approximately 10 cm² and 5 cm³ greater than those of Huang et al., obtained with 11.7T and 4.7T post-mortem MR imaging, respectively. However, our results might be more accurate because they were obtained with 7T MR imaging from a greater amount of specimens, and all of the specimens were scanned in situ, which may have avoided deformations caused by human factors or gravity when the brain is removed from the cranial cavity.

Brain length in our research was roughly 0.5 cm smaller than the fronto-occipital diameter of 20–22 weeks GA, described by Parazzini et al., and was roughly 1 cm smaller than the fronto-occipital diameter of 17–22 weeks GA, described by Moreira et al. For brain width, our research demonstrates a width of approximately 0.5 cm greater than the cerebral biparietal diameter described by Parazzini et al. and Moreira et al. These inconsistencies are mainly the result of different definitions of brain measurements found in our research and in theirs. Our measurements may be more convincing because they were obtained from 3D visualization models based on MR imaging of high magnetic strength instead of 2D in vivo fetal MR imaging.

There have been 2 opposing views in US research about whether sexual dimorphisms and cerebral asymmetries are present.
ent in fetal brains in the second half of gestation, but MR imaging studies have discovered these features in premature and neonatal brains. Our results, however, indicate that no significant sexual dimorphisms and cerebral asymmetries are present during the second trimester. Because the total number of subjects and their distribution in each GA group are relatively small, it can only be suspected that brain sexual differentiation starts in the third trimester, when the brain has finished the initial developmental phase and will step into an accelerated and complicated developmental period.

**Superiority of 7T Postmortem MR Imaging in Demonstration of the Fetal Brain in the Second Trimester**

The low-image contrast of in vivo fetal MR imaging precludes the application of postprocessing tools dedicated to the adult brain. However, postmortem MR imaging obtained with high magnetic strength is of high quality and is sufficient for segmentation, reconstruction, and quantitative analysis. The consistency in demonstrating the cortical surface between the gross anatomy and the 3D visualization model has been proved. Manual segmentation, which may be both tedious and time-consuming for large imaging, is much more precise than automatic segmentation, especially for the immature fetal brain. It is thought that these 3D visualization models, with dynamic demonstration, can exactly delineate developmental changes of the fetal brain during the second trimester, and supply quantitative data about the cortical surface area and volume that is hard to obtain by US, in vivo MR imaging, or histology studies. Our results may benefit the comprehension of immature structures in the second trimester.

**CONCLUSIONS**

The second trimester is the most important phase, during which most sulci are present and can be clearly shown on 7T postmortem MR imaging. Perhaps the specific time during which neuro-pathologic features of sulci emerge, previously thought to be well understood, should be redefined. Quantitative data assist greatly in the precise understanding of the immature brain. Measurements of fetal brains with different growth rates increase linearly with GA. Fetal brain sexual dimorphisms and asymmetries may arise and develop during the third trimester. Our results are valuable in anatomic education, research, and assessment of normal brain development in the uterus.

**ACKNOWLEDGMENTS**

We thank Hamed Haghazaz, Sam Hobel, and Xuntao Yin for linguistic advice during the revision.

**REFERENCES**

1. Encha-Razavi F, Sonigo P. *Features of the developing brain.* *Childs Nerv Syst* 2003;19:426–28
2. Prayer D, Kasprian G, Krampl E, et al. *MRI of normal fetal brain development.* *Eur J Radiol* 2006;57:199–216
3. Glenn OA. *MR imaging of the fetal brain.* *Pediatr Radiol* 2010;40:68–81
4. Fogliarini C, Chaumoitre K, Chapon F, et al. *Assessment of cortical maturation with prenatal MRI: part II: abnormalities of cortical maturation.* *Eur Radiol* 2005;15:1781–89
5. Chi JG, Dooling EC, Gilles FH. *Gyral development of the human brain.* *Ann Neurol* 1977;1:86–93
6. Garel C, Chantrel E, Brisse H, et al. *Fetal cerebral cortex: normal gestational landmarks identified using prenatal MR imaging.* *AJNR Am J Neuroradiol* 2001;22:184–89
7. Monteagudo A, Timor-Tritsch IE. *Development of fetal gyri, sulci and fissures: a transvaginal sonography study.* *Ultrasound Obstet Gynecol* 1997;9:222–28
8. Hansen PE, Ballesteros MC, Soila K, et al. *MR imaging of the developing human brain. Part 1. Prenatal development.* *Radiographics* 1993;13:21–36
9. Parazzini C, Righini A, Rustico M, et al. *Prenatal magnetic resonance imaging: brain normal linear biometric values below 24 gestational weeks.* *Neuroradiology* 2008;50:877–83
10. Moreira NC, Teixeira J, Themudo R, et al. *Measurements of the normal fetal brain at gestation weeks 17 to 23: a MRI study.* *Neuroradiology* 2011;53:43–48
11. Habas PA, Kim K, Rousseau F, et al. *Atlas-based segmentation of developing tissues in the human brain with quantitative validation in young fetuses.* *Hum Brain Mapp* 2010;31:1348–58
12. Zhang Z, Liu S, Lin X, et al. *Development of fetal cerebral cortex: assessment of the folding conditions with post-mortem magnetic resonance imaging.* *Int J Dev Neurosci* 2010;28:537–43
13. Zhang Z, Liu S, Lin X, et al. *Development of laminar organization of the fetal cerebrum at 3.0T and 7.0T: a postmortem MRI study.* *Neuroradiology* 2011;53:177–84
14. Zhang Z, Liu S, Lin X, et al. *Development of fetal brain of 20 weeks gestational age: Assessment with post-mortem magnetic resonance imaging.* *Eur J Radiol* 2011;80:e432–39
15. Meng H, Zhang Z, Geng H, et al. *Development of the subcortical brain structures in the second trimester: assessment with 7.0-T MRI.* *Neuroradiology* 2012;54:1153–59
16. Bayer SA, Altman J. *The human brain during the second trimester.* In: *Atlas of Human Central Nervous System Development*, Indianapolis: CRC Press; 2005
17. Tang Y, Hojatkashani C, Dinov ID, et al. *The construction of a Chinese MRI brain atlas: a morphometric comparison study between Chinese and Caucasian cohorts.* *NeuroImage* 2010;51:33–41
18. Régis J, Mangin JF, Ochialli T, et al. “Sulcal root” generic model: a hypothesis to overcome the variability of the human cortex folding patterns. *Neurol Med Chir (Tokyo)* 2005;45:1–17
19. Dubois J, Benders M, Borradori-Tolsa C, et al. *Primary cortical folding in the human newborn: an early marker of later functional development.* *Brain* 2008;131:2028–41
20. Dubois J, Benders M, Cachia A, et al. *Mapping the early cortical folding process in the preterm newborn brain.* *Cereb Cortex* 2008;18:1444–54
21. Torkildsen A. *The gross anatomy of the lateral ventricles.* *J Anat* 1934;68:480–91
22. Huang H, Xue R, Zhang J, et al. *Anatomical characterization of human fetal brain development with diffusion tensor magnetic resonance imaging.* *J Neurosci* 2009;29:4263–73
23. Tiele B, Alberti C, Adamsbaum C, et al. *Cerebral biometry in fetal magnetic resonance imaging: new reference data.* *Ultrasound Obstet Gynecol* 2009;33:173–81
24. Kivilevitch Z, Achiron R, Zalel Y. *Fetal brain asymmetry: in utero sonographic study of normal fetuses.* *Am J Obstet Gynecol* 2010;202:359.e1–8
25. Gilmore JH, Lin W, Prastawa MW, et al. *Regional gray matter growth, sexual dimorphism, and cerebral asymmetry in the neonatal brain.* *J Neurosci* 2007;27:1255–60