ABSTRACT
Visualization of the fetal face with three- and four-dimensional (3D/4D) ultrasound is a unique experience for both, the parents-to-be and the operator. While the future parents are primarily interested in seeing the surface of the fetal face and facial movements, the operator uses the different display modes for a precise fetal malformation check. The multiplanar mode with a simultaneous display of the three perpendicular planes allows an accurate demonstration of a normal or an abnormal fetal profile. Even if an image of the fetal face is acquired in an oblique position, the stored volume can be rotated by the rotational controls in all three directions, until the face is seen precisely in the median plane. The different surface modes enable the operator to detect abnormal protuberant structures or surface defects, while the transparent mode (maximum mode) reveals ossification defects. During a targeted ultrasound examination of the fetal face, five different regions have to be assessed: the forehead, orbits and eyes, nose, mouth, and chin. With 3D ultrasound the following fetal anomalies can be detected: anomalies including the forehead (anencephaly, protuberant forehead, abnormal metopic suture, frontal encephalocele/meningocele), anomalies of the orbits and the eyes (orbital hypoplasia/microphthalmia, hypertelorism, hypotelorism, cataract, nasolacrimal cyst/dacryocystocele, cyclopia/proboscis), abnormalities of the nose (flat nose, absent nasal bone), abnormalities of the mouth (cleft lip/cleft palate, epignathus, macroglossia, chin anomaly (retrognathia/micrognathia). 3D ultrasonography allows a comprehensive evaluation of the fetal face with different display modes. In contrast to 2D ultrasound, 3D ultrasound enables a detailed demonstration of the soft tissue of the fetal face and thus contributes to a better understanding of the malformation by both the physician and the future parents.

Keywords: Congenital anomalies, Fetal face, Three-/Four-dimensional ultrasonography.

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INTRODUCTION
Visualization of the fetal face with 3D/4D ultrasound is a unique experience for both, the parents-to-be and the operator. While the future parents are primarily interested in seeing the surface of the fetal face and facial movements, the operator uses the different display modes for a precise fetal malformation check. The multiplanar mode with a simultaneous display of the three perpendicular planes allows an accurate demonstration of a normal or an abnormal fetal profile. Even if an image of the fetal face is acquired in an oblique position, the stored volume can be rotated by the rotational controls in all three directions, until the face is seen precisely in the median plane. The different surface modes enable the operator to detect abnormal protuberant structures or surface defects, while the transparent mode (maximum mode) reveals ossification defects.

During a targeted ultrasound examination of the fetal face, five different regions have to be assessed: the forehead, orbits and eyes, nose, mouth, and chin.

ANOMALIES INCLUDING THE FOREHEAD
Anencephaly
Anencephaly is the most common and severe anomaly of the central nervous system with an incidence rate of 1:1000 births. The surface demonstration of the fetal head shows an absent superior vault, the missing cerebrum and large bulging eyes (“frog eyes”) (Fig. 1). The respective ultrasound appearance of the face has also been described as “Mickey Mouse face”. Anencephaly can be detected as early as 9–10 week’s gestation.3,4

Fig. 1: Surface view (HDlive) of an anencephalus at 30 weeks’ gestation
Anomalies of the Fetal Face

Protuberant Forehead

The term frontal bossing refers to the development of an unusually pronounced forehead caused by an enlargement of the frontal bone. In several cases, it has been observed in conjunction with abnormal enlargement of other facial bones. A protuberant forehead may be observed in different fetal abnormalities, such as achondroplasia, Crouzon syndrome, Fragile X syndrome, Hurler syndrome, Marfan syndrome, Pfeiffer syndrome, Rubinstein-Taybi syndrome, and Russell-Silver syndrome. Moderate frontal bossing may also be seen in some normal fetuses.

Sonographically, frontal bossing is best seen in a side view (Fig. 2), but it can also be unambiguously demonstrated in a frontal surface view.

ABNORMAL METOPIC SUTURE

The metopic suture can be demonstrated in the transparent mode from a front view (Fig. 3).

Wide Metopic Suture

Wide V-, Y- and U-shaped metopic sutures have been observed in fetuses with facial defects involving the orbits, nasal bones, lip, the palate, and mandible, as well as in fetuses with cerebellar abnormalities (Chaoui). A wide V-shaped metopic suture can also be observed in fetuses with chromosomal defects (Fig. 3A) or in those with osteogenesis imperfecta type II.

Narrow Metopic Suture

Premature closure of the suture (craniosynostosis) or the presence of an additional bone between the frontal bones may be a sign for holoprosencephaly and abnormalities of the corpus callosum. Premature closure of the metopic suture is also found in Apert syndrome (Fig. 3B). Fetal neurosonography is mandatory in all fetuses with craniosynostosis.

Frontal Encephalocele/Meningocele

Frontoethmoidal encephalocele represents a frontal skull defect with herniated brain substance. This is in contrast to frontal meningocele where the herniation contains only leptomeninges with cerebrospinal fluid. The incidence of frontoethmoidal encephalocele is 1:40,000 live births. The presence of a frontal lesion has been observed in 15% of all encephaloceles.

A frontoethmoidal encephalocele is identified sonographically as a cystic solid mass between the orbits. In the presence of a frontal meningocele, only a hernial sac filled with liquid is observed between the orbits (Fig. 4A). In both findings, a skull defect is visualized behind the protrusion (Fig. 4B).

ANOMALIES OF THE ORBITS AND THE EYES

Orbital Hypoplasia/Microphthalmia

In orbital hypoplasia, the orbital size ranges below the 5th percentile and the eye may be abnormally small (Fig. 5). The most extreme situation is encountered in anophthalmia where the eye is completely missing from the orbit. Microphthalmia may occur as an isolated abnormality, but is also seen in chromosomal defects (trisomy 13, triploidy), holoprosencephaly and various other syndromes.

Hypertelorism

Hypertelorism is defined as an abnormally increased interorbital distance (inner and outer orbital distances) above the 95th percentile. A number of different fetal

Fig. 2: Achondroplasia (37 weeks’ gestation). The profile (lateral surface view) of the fetal face shows pronounced frontal bossing.

Figs 3A and B: Transparent demonstration of the skull from a front view. (A) Wide V-shaped metopic suture in a fetus with trisomy 22 (20 weeks’ gestation); (B) Premature closure of the metopic suture in Apert syndrome (37 weeks’ gestation).
anomalies associated with hypertelorism has been described: chromosome anomalies, various syndromes (e.g., Apert syndrome, Crouzon syndrome), ethmoidal encephalocele and facial haemangioma. Ultrasonic measurements of the inner and the outer orbital distances can be performed in both the coronal (Fig. 6A) and axial demonstration of the orbits.

The high risk for associated malformations requires both, a targeted ultrasound examination and karyotyping.

Hypotelorism

Hypotelorism is defined as a decreased interorbital distance below the 5th percentile. There is a high risk for associated malformations (chromosome anomalies, holoprosencephaly, microcephaly, and Meckel syndrome. As in hypertelorism, measurements of the inner and the outer orbital distances can be performed in the coronal (Fig. 6B) or axial demonstration of the orbits.

Cataract

A cataract is an opacification of the crystalline lens in the eye. Cataracts represent a rare fetal condition that may be caused by intrauterine infections (cytomegaly, rubella, varicella), and chromosome abnormalities (trisomies 13, 18, 21), or it may form a part of a syndrome.

A beginning cataract appears sonographically as a hyperechoic dot in the fetal lens, and a complete cataract is visualized as a solid hyperechoic disc in the lens (Fig. 7).
Anomalies of the Fetal Face

Nasolacrimal Duct Cyst/Dacryocystocele

The nasolacrimal duct cyst is a unilateral or bilateral benign cyst of the nasolacrimal duct. Embryogenesis is attributed to the failure of the valve of Hasner at the distal end of the nasolacrimal duct. In dacryocystocele, there is an obstruction of the lacrimal drainage system both above (Rosenmüller valve) and below the sac (valve of Hasner).16,17

A lacrimal duct cyst/dacryocystocele appears sonographically as a cystic mass in the inferomedial canthus (Fig. 8). Dacryocystoceles are typically not identifiable until 30 weeks' gestation.18 In view of the fact that they may be a part of various syndromes, the investigator should carefully examine the fetus for other associated anomalies.19

Cyclopia/Proboscis

Cyclopia is a rare facial abnormality characterized by the failure of the embryonic prosencephalon to properly divide the orbits of one eye into two cavities. Instead of the nose, a proboscis is usually present, i.e., a trunk-like appendage in the midline of the face originating above the eye level. Cyclopia and proboscis occur in association with holoprosencephaly.20,21

Ultrasound of the fetal face reveals a single orbit that is best demonstrated in the coronal or axial transparent view (Fig. 9A). The proboscis is best seen in a lateral surface view (Fig. 9B).

NOSE ABNORMALITIES

Flat Nose (lat Profile)

A flat profile is a known marker for a chromosomal abnormality (trisomy 21) in the second trimester.22-25

One of the main advantages of 3D ultrasound technology is the ability to provide the operator with a true median plane of the face, using the multiplanar mode.1 After identification of the precise profile in the multiplanar mode, several fetal measurements can be performed to confirm a flat profile.24,26

A frontal fetal facial angle of >145° identified in the second trimester should raise suspicion for trisomy 21.24

A flat face can also be demonstrated with the surface mode from a lateral view (Fig. 10).

Every flat profile requires a targeted ultrasound examination with a diligent search for additional anomalies, and fetal karyotyping should be performed.

Absent Nasal Bone

Missing ossification of the two bilateral nasal bones is a further marker for trisomy 21.27,28

3D ultrasound allows the identification of an absent nasal bone in both the multiplanar and the transparent mode (Fig. 11).
ABNORMALITIES OF THE MOUTH

Cleft lip/Cleft Palate

Orofacial clefts are one of the most common congenital anomalies. The group of orofacial anomalies is heterogeneous. It comprises “typical” orofacial clefts [cleft lip (CL), cleft lip and cleft palate (CLP) and cleft palate only (CP)], as well as “atypical” clefts (median, transversal, oblique and other Tessier facial clefts).

Both typical and atypical clefts can occur as an isolated anomaly, as part of a sequence of the primary defect, or as multiple congenital anomalies. More than 300 syndromes are associated with facial clefting. The most common chromosomal aberrations with orofacial clefts are trisomy 13, trisomy 18 and trisomy 21.

Ultrasound demonstration of different cleft lips is most successful in the surface mode (Fig 12). The detection of cleft palate is more difficult, particularly in the presence of an isolated cleft palate. Several 3D techniques have been described for the detection of cleft palate. The defect can be demonstrated in different modes: the multiplanar mode, tomographic mode, surface mode, and Omniview/VCR mode. The defect is most readily demonstrated when the fetus is yawning and the oral cavity is filled with amniotic fluid (Fig. 13).

Fetal karyotyping is recommended with a view to the risk for a chromosomal abnormality.

Epignathus

Epignathus is a rare type of teratoma arising in the oral cavity. In most cases, it is a benign tumor. However, it is associated with high mortality and morbidity rates because of severe airway obstruction and other malformations. Rare malignant cases have also been described in the literature.

The 3D surface mode shows a solid tumor in front of the fetal mouth (Fig. 14).

Macroglossia

Macroglossia is a disorder characterized by an abnormal enlargement of the tongue. It may occur as an isolated and sporadic trait, as a familial trait, or in association with Beckwith-Wiedemann syndrome or Down syndrome.

Ultrasound demonstration of macroglossia is best achieved in the surface mode and demonstrates an enlarged tongue in the open mouth (Fig. 15).

Chin Anomaly

Micrognathia/Retrognathia

Both micrognathia and retrognathia involve abnormal, arrested development of the mandible. Micrognathia...
refers to the size of the mandible whereas retrognathia refers to its position in relation to the maxilla (Paladini). Micrognathia is frequently associated with a number of different syndromes, skeletal and neuromuscular diseases, as well as with chromosome anomalies (primarily trisomy 18).

The sonographic diagnosis can be made on a subjective or objective basis. The subjective diagnosis is carried out by evaluating the midsagittal view of the facial profile and by assessing the geometric relationship between the mandible and the rest of the profile (Fig. 16). For an objective diagnosis, both the inferior facial angle (IFA) and the jaw index are used.

When micro-/retrognathia is detected, a detailed search for additional fetal abnormalities and fetal karyotyping is required.

CONCLUSION

3D ultrasonography allows a comprehensive evaluation of the fetal face with different display modes. In contrast to 2D ultrasound, 3D ultrasound enables a detailed demonstration of the soft tissue of the fetal face and thus contributes to a better understanding of the malformation by both the physician and the future parents.

REFERENCES

10. Francois J. Genetics of cataract. Ophthalmologica 1982;184;6171


