Moving forward: early diagnosis of fetal spinal defects

Moving prenatal diagnosis and screening to earlier stages of pregnancy has probably been the most important progress in the field of Fetal Medicine in the recent years. Screening for chromosomal abnormalities is the most obvious example; the sensitivity of screening for Down syndrome by maternal age only (1980s – early 1990s) was only 30%, which rose to 60% by second-trimester biochemical tests (late 1980s – early 1990s) and is nowadays as high as 95% with the implementation of first-trimester screening with ultrasound and biochemical markers [1].

A by-product of the advent of first-trimester screening for chromosomal abnormalities is that examiners became increasingly familiar with early fetal anatomy. As a result, nearly half of all major fetal structural defects can currently be detected at the 11–14 weeks’ scan if a structured examination of the fetal anatomy is performed, and this includes virtually all cases of acrania, alobar holoprosencephaly, exomphalos, gastroschisis, megacystis and body stalk anomaly [2]. However, the detection rate of spinal defects in early pregnancy has been notoriously lower than that, as only 14% of the cases with open spina bifida were picked up at the first-trimester scan in the Syngelaki et al. series [2].

These rates are very likely to improve through the examination of a novel first-trimester ultrasound marker, the so-called intracranial translucency (IT). Intracranial translucency, which essentially represents the early fourth cerebral ventricle, was first described by Chaoui et al. in 2007 as an anechoic space between the brain stem and choroid plexus [3] (Fig. 1A).

In cases with spina bifida the fourth ventricle is compressed through caudal displacement of the brain stem, and IT is usually obliterated. Additional measurements and ratios of the posterior fossa have also been proposed in order to improve sensitivity [4,5], and abnormal measurements (obliteration or dilation) have been reported in various CNS defects, such as diastematomyelia [6], Dandy-Walker complex [7] or Blake's pouch cyst [8]. Similar to nuchal translucency, IT can be measured using a semi-automated algorithm, with high reproducibility [9]. A large prospective multicentre study in Germany very recently reported 100% sensitivity (11/11) for open spina bifida by structured examination of the first-trimester posterior fossa [10].

In the case of Fig. 1B, obliteration of the fetal IT at 11+2 weeks (CRL 48 mm) led to the identification of a lumbar defect measuring 5.2 \times 2.4 \text{mm} (Fig. 2A,B) before the disruption in the architecture of the spine becoming visible (Fig. 2C), in a couple with no known risk factors for CNS defect. The parents were counselled about the diagnosis and opted for termination of pregnancy at 12 weeks.

The rationale for early detection is that it offers the parents time to digest the news and realize their feelings, facilitates them in their decision-making process and lessens the degree of psychological and physical strain should they opt for a termination. One might argue that the earlier the diagnosis, the easier the parents might decide to terminate pregnancy, which for some might pose an ethical question. Although there is not nearly enough data for the case of structural defects, the paradigm of chromosomal abnormalities shows that gestational age at diagnosis mostly influences parents’ decision for mild and moderate rather than severe ab-
normalities [11]. Anyhow, moving earlier in pregnancy appears to be the way forward, and, as many experts believe, a combination of maternal characteristics and history together with specific biophysical and biochemical tests in the first trimester will enable providers to predict many pregnancy complications [12] and produce a triage model which will result in tailored subsequent obstetric care.

References
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Fig. 2 Disruption of the continuity of the skin in the lumbar area in the fetus of Figure 1B, in a coronal plane (A) and axial plane (B), with apparently normal architecture of the lumbar spine on 3D reconstruction (C).