

Euro Pediatrics 2018: The genetic background of human neural tube development in the aspect of prenatal diagnosis - Krzysztof Piotrowski - Pomeranian Medical University

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Congenital neural tube defects (NTD) are communal malformations both as an isolated form and a part of genetic syndromes. Extremely fast development of molecular genetics confirms that almost all NTD are genetically dependent in terms of aberrations in different regions of a chromosome or single gene mutations. On the other hand, NTD are an important component of diverse genetic diseases, including monogenic and metabolic disorders with mutations (often called polymorphism) genes responsible for the condition of the MTHFR gene. The genes participating therein are located near on each chromosome, mainly on pathways, along with ligand genes and co-factors, transcription factors or individually. Several mechanisms on NT development are based on the balance between apoptosis, proliferation and migration. Crucial genes regulatory fetal development, including the creation of neural tube and the forming of vertebral continuity are primary home box genes grouped in 4 clusters HOX1-4. Other genes condition the forming of different structures. The most important pathways are Shh, Wnt, FGF, Notch, and BMP. Instead of recognition that periconceptional folic acid supplementation in reproductive age women decreases the risk of fetal neural tube defects (NTDs), these malformations remain as the second most collective serious fetal birth defect in the United States, are mostly surpassed only by congenital heart defects. Methods castoff to detect neural tube defects are now a component of routine obstetrical care, and include both second trimester maternal serum alpha-fetoprotein (MSAFP) levels and fetal ultrasonographic evaluations. Once an NTD is finds as a various management options are available for families, including consideration of pregnancy termination, in utero fetal surgery, as well as referral to a tertiary care center for management and delivery. Therefore, it is useful for the practicing obstetricians to be aware of these complex diagnostic and management options for optimal care of the obstetric patient and fetus.1 In this chapter, the etiologies, antenatal diagnosis, and management, as well as information regarding prevention are find out. All pregnant women have the possibility to undergo prenatal screening/diagnosis for genetic conditions and/or birth defects. Specific signs for genetic counselling and prenatal diagnosis options include a history of chromosome abnormality, Mendelian genetic disorder, or metabolic disorder; increased risk for neural tube defect; abnormal maternal serum screening test; or a fetal anomaly suspected/diagnosed on ultrasound. Fruitful prenatal diagnosis requires a known condition associated with a structural abnormality visible in the fetus, a biochemical abnormality in the

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