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FETO-PLACENTAL STEROID METABOLISM IN GROWTH RETARDED HUMAN FETUSES

Women bearing fetuses with intrauterine growth retardation (IUGR) frequently have low serum estriol (E₃) levels in the gestation. One origin of the low E₃ production in these pregnancies may be reduced secretion of fetal adrenal cortex-derived neutral steroid precursors of estriol synthesis in the placenta. However, there is evidence that placental conversion of the neutral precursors to estrogen may be low in association with a fetus with IUGR. In order to investigate the relative importance of these two possible causes of low E₃ production, umbilical venous, and maternal venous neutral steroid and estrogen levels were assessed in a series of pregnancies in which the fetus had IUGR.

The significantly low E₃ values in both umbilical and maternal samples are postulated to result not only from the reduced fetal adrenal dehydroepiandrosterone sulfate (DHAS) secretion, but also underactive 16 α -hydroxylase activity in fetal liver or low efficiency of 16 α -OH-DHAS relative to DHAS, as a substrate for placental conversion to an estrogen.

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