



Evaluation of Anti Mullerian Hormone (AMH) assay Roche® on umbilical cord blood: Determination of reference values in newborn girls and boys.

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Background

AMH concentration is now well studied for prepubertal boys (Plotton 2009, 2012) or women in reproductive medicine (Anderson, 2015). AMH reference values have been determined with the Roche® automated technique, but we dispose very few data about AMH reference value for the newborns. These value could be helpful for the management in neonatal period of neonate cases with Disorders of Sex Development (DSD).

Objective

Determine AMH reference values on umbilical cord blood for girls and boys neonates.

Methods

This study of AMH values on umbilical cord blood was realized after approval of the ethics committee between May 13th, 2015 and February 4th, 2016.

AMH assay was realized by using an automated Electrochemiluminescence Immunoassay (Cobas Roche®).

We analyzes AMH on Heparin syringes for blood gas systematically collected after births.

Personal history and clinical data for all the sample were collected simultaneously at the systematic examination (3d day or before the exit): gestational age and history of syndromic malformations or urogenital anomalies like cryptorchidism, hypospadias or micropenis diagnosed on the clinical exam.

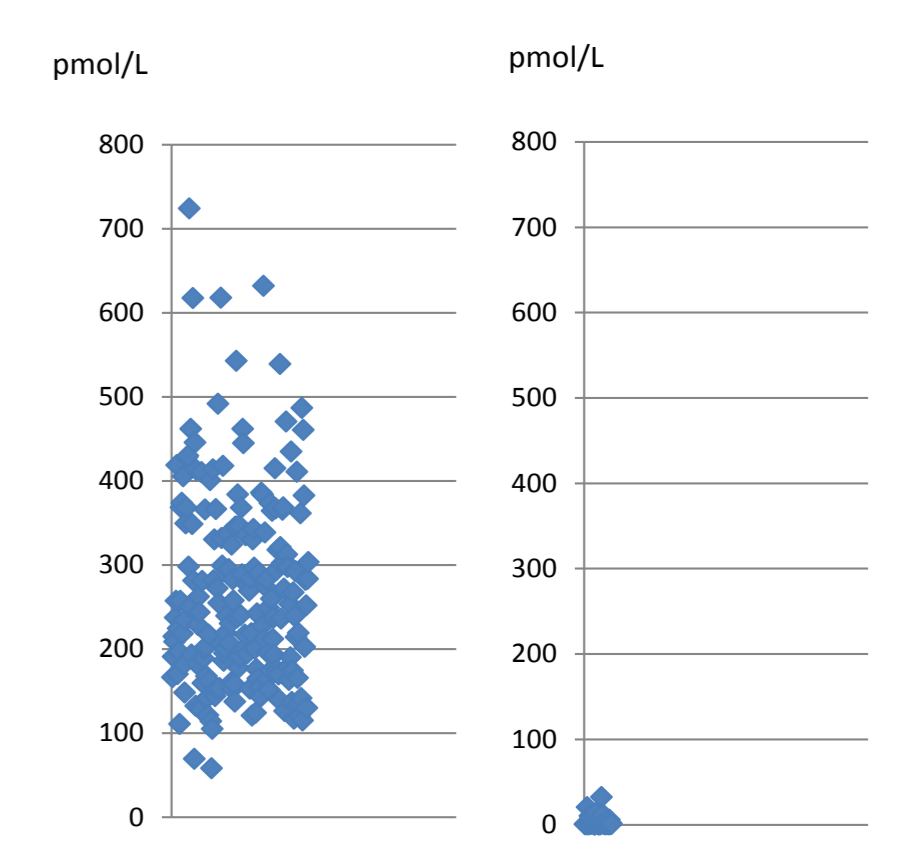
We excluded samples of neonates presenting genitourinary abnormalities.

During this period, we also evaluates AMH in 7 cases of DSD neonatal (hypospadias with intrascrotal or inguinal gonad) for which we identify an molecular explanation.

Results

□ 230 blood samples of veinous umbilical from neonates for which no clinical abnormalities were include. This study - The samples of 192 boys and 38 girls allowed of reference interval for AMH 20 the first hours of life (See Table and graph)

Graph repartition of AMH values in boys and girls neonate



pmol/L	n	Median	1st Quartil	3d Quartil	Girls min-max
Boys	192	249.95	178.7	338	58.5-724.3
Girls	38	1.03	0.56	2.59	0.28-32.5

AMH reference values between boys and girls significantly different ($p < 0,0001$)

□ Among the 7 neonates, we identified :

- 3 cases of DSD with defective androgen (DA) action (mutation Androgen Receptor mutation (1), of 5 alpha reductase (2))
- 4 cases with testicular dysgenesis (TD): 1 SF1 mutation case, 3 mosaicism (45X, 46XY)

pmol/L	n	Mean	min-max
DA	3	544	292-777
TD	4	98.8	72.4-140

Conclusions

These preliminary data of normal values on venous umbilical cord blood appear to be useful for etiological issue and the management of DSD at birth.

References

Anderson et al. Fertil steril, 2015
Plotton et al. Horm Research, 2009
Plotton et al. Horm Research, 2012

