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## PHARMACOKINETICS OF LITHIUM DURING DELIVERY AND IN THE NEONATAL PERIOD. A PRELIMINARY DATA

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**Introduction:** Lithium has been used in the treatment of bipolar disorder in pregnant women. However, information on the pharmacokinetics of lithium during perinatal period is scarce.

Objectives: To study pharmacokinetics of lithium during delivery and in the neonatal period.

**Methods:** A prospective, observational and naturalistic study was conducted at the PERINATAL PSYCHIATRY PROGRAM CLÍNIC-BARCELONA, from 2005 to 2012. We included all consecutive cases of pregnant women with bipolar disorder I or II (n=22), and on maintenance treatment with lithium monotherapy (n=13) or polytherapy (n=9) during pregnancy who elected artificial feeding. Lithium plasma concentrations in maternal blood and umbilical cord were detected. Lithium plasma concentrations in infants (n=16) at delivery and in the neonatal period were obtained to calculate elimination half-life, which was estimated by lineal regression. Technique: AVL 9180 electrolyte analyser using a lithium-selective electrode (detection limit =0.10 mEq/L).

**Results:** Women did not fulfil diabetes criteria pre-pregnancy and during pregnancy. Attending to neonatal outcomes, infants exposed to polytherapy had a higher weight at birth (percentils) than those exposed to lithium alone [53.38 (33.40) vs. 70.22 (26.25)]. No statistically significant differences were found in umbilical cord:maternal plasma concentration ratio between those treated with lithium monotherapy and women treated with polytherapy (1.05 vs. 1.08). The lithium mean elimination half-life (SD) in infants was 6.73 (9.12) days.

**Conclusions:** Lithium crosses placental barrier almost completely. Elimination half-life in neonates exposed to lithium in utero was 6.73 days. Moreover, lithium treatment during pregnancy requires therapeutics monitoring in exposed dyads.