

DECREASED EXPRESSION OF OESTROGEN RECEPTOR β IN THE REPRODUCTIVE TRACT OF PREGNANT RELAXIN-DEFICIENT (*Rlx*^{-/-}) MICE

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The peptide hormone relaxin (RLX) is reported to directly affect uterine oestrogen receptors (ERs) in the rat (1). Treatment of immature ovariectomised rats with porcine RLX causes a decrease in uterine ER β mRNA levels within 6 h. However, RLX has no effect on ER α expression. As both ER β 1 and ER β 2 inhibit ER α -mediated transcriptional activity, this RLX-induced downregulation in ER β could be a prerequisite for oestrogen to exert its effects on target tissues. The aim of the current study was to use relaxin-deficient (*Rlx*^{-/-}) pregnant mice to investigate if relaxin deficiency results in alterations in either ER β or ER α mRNA expression in reproductive tissues. Cervix and vagina tissues were obtained from adult C57/Blk6J wild-type mice at five stages of gestation (Days 7.5, 10.5, 14.5, 17.5, 18.5 pc) and *Rlx*^{-/-} littermates on Days 7.5, 14.5 and 18.5 pc. Q-PCR with TaqMan probes in the Opticon 2 thermal cycler (MJ Research, GeneWorks) was used to quantify ER α and ER β gene expression. ER α mRNA levels were significantly ($P < 0.05$; ANOVA) increased in the cervix/vagina on Days 17.5 and 18.5 pc in *Rlx*^{+/+} mice. The increase in ER α in *Rlx*^{+/+} mice was negatively correlated with a significant decrease in ER β expression from Day 14.5 pc. In contrast, there was no decrease in ER β gene expression in the cervix/vagina of *Rlx*^{-/-} mice; ER β mRNA levels were significantly ($P < 0.05$) higher compared to *Rlx*^{+/+} mice on Days 14.5 or 18.5 pc. However, there was no corresponding reduction in ER α expression in the cervix/vagina of the *Rlx*^{-/-} mice, so that ER α mRNA levels were still elevated at term despite the maintenance of high ER β expression. In summary, these data show changes in ER β expression in the cervix/vagina of relaxin-deficient mice, which may subsequently affect ER α -mediated transcriptional activity.

(1) Pillai *et al.* (2002) *Biol. Reprod.* **67**, 1919–1926.