

# Progesterone synthesis by human luteal cells: Modulation by estradiol

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To assess the role of estradiol (E2) upon progesterone (P4) synthesis, a well defined human midluteal cell system was used. A dose-dependent inhibition of P4 synthesis with and without hCG was induced by E2. In addition, E2 had a dose related cumulative effect on pregnenolone as compared with control experiments (2-fold,  $P < 0.05$ ) as well as in hCG- stimulated conditions (3-fold,  $P < 0.005$ ). On the other hand, the concentrations of 20 $\alpha$ -hydroxyprogesterone obtained in all experimental conditions were similar to control values, indicating that the catabolism of P4 was not modified. 3 $\beta$ -Hydroxysteroid dehydrogenase activity was significantly diminished ( $P < 0.05$ ) in the presence of E2. Finally, the kinetic studies on P4 synthesis from pregnenolone showed a competitive type of inhibition with a  $K_1$  of  $2.22 \times 10^{-6}$  mol/L. These data indicate an inhibition of 3 $\beta$ -hydroxysteroid dehydrogenase on human corpus luteum by E2.