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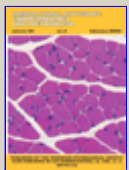
JOURNAL ARTICLE

Effects of finasteride on the rat ventral prostate

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The objectives of this study were to compare changes in the ventral prostate (VP) of young adult Sprague Dawley rats after 28 days of treatment with finasteride (F), a potent 5 alpha-reductase inhibitor, with those caused by castration (Cx). VP concentrations of DHT were reduced to 20.2% and 6.6% of controls (1,947 +/- 207 pg/VP, mean +/- SE) by F (5 or 20 mg/kg/day) and to 2.6% of controls by Cx. VP weights were reduced 49% and 54% by F and 88% by Cx. DNA/VP fell 25% and 15% with F treatment and 72% after Cx, whereas RNA and protein/VP were reduced 37-51% by F and 91-93% by Cx. The RNA/DNA and the protein/DNA ratios fell to 30-36% of controls after F treatment and to 70% of controls after Cx. The mRNA concentrations of the C3 subunit of prostatein 28S ribosomal RNA fell after treatment with F (5 mg/kg/day) and after Cx, whereas the mRNA for TRPM-2, an androgen-suppressed protein associated with apoptosis, was increased only after castration. To examine further the effects of F on the rate of DNA synthesis, 7-day regressed adult rats were treated for 3 days with testosterone propionate +/- F, and incorporation of 3H-thymidine in minced ventral prostates was determined. F inhibited 3H-thymidine incorporation. We conclude that Cx causes a greater reduction in cell number/VP and a greater reduction in RNA and protein cell than F and that the differences between F treatment and castration probably result from differences in prostatic concentrations of T. (ABSTRACT TRUNCATED AT 250 WORDS)

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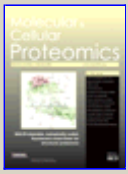
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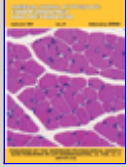
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