INFLUENCE OF AGE ON PEAK FREQUENCY OF LH AND TESTOSTERONE IN A HEALTHY POPULATION J.P.Deslypere, J.L.Vandalem°, A.Vermeulen^{*}. Dept. of Endocrinology State University of Ghent^{*} and Liège[°], Belgium.

Aging influences the hypothalamic-pituitary-testicular axis in a complex fashion. Although the primary lesion is essentially at the testicular level, the delayed pituitary response to GnRH suggests the existence of discrete change in hypothalamopituitary function. We studied peak frequency of LH and T secretion in young and elderly males. Blood samples were taken every 20 minutes for 24 hrs in 20 young (<60 yrs) and 13 elderly (>70 yrs) monks living together in a monastery. Mean T was significantly lower in the elderly (402±181 ng/d1) than in the young monks (535±134), while the opposite was true for LH (2.8±2.1 ng/m1 and 1.7±0.3 respectively) (P<0.05). Peak frequency, analysed by pulsar, of T and LH was significantly higher in 20-40 yrs old monks (n=10) than in elderly monks (LH :3.9±1.0 and 3.0±0.7; T :3.7±0.8 and 1.7±1.0 pulses/12 hrs) (P<0.01). No differences could be found in peak amplitude for both hormones between young and old monks. Percutaneous 5α-androstanolone (DHT) administration resulted in a more important decrease of LH and T levels in elderly than in young males (T : Δ post-pre young 123±123 ng/d1 - Δ old 204±78; LH : Δ young 1.1±1.5 ng/m1 - Δ old 2.0±1.6) (P < 0.01) indicating a greater sensitivity to feedback in elderly males in comparison to young males.

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ADRENAL STEROIDS STIMULATE THE GROWTH OF DIMETHYLBENZANTHRACENE (DMBA)- INDUCED RAT MAMMARY TUMORS. Spinola, P.G., Marchetti, B. and Labrie, F., MRC Group in Molecular Endocrinology, Laval University Medical Center, Quebec, GIV 4G2 - Canada.

There is convincing evidence that adrenal steroids, especially androst-5-ene-38,178diol (Δ^5 -diol), play a role in the etiology of breast cancer. Dehydroepiandrosterone (DHEA) and DHEA-sulfate (DHEA-S) have been postulated as being involved in the development of breast cancer through the formation of their metabolite Δ^5 -diol. In order to further investigate the role of adrenal steroids in hormone-sensitive cancer growth, we have studied the effect of treatment with Δ^5 -diol (2 mg, twice daily) or DHEA (2 mg, twice daily) on the growth of DMEA-induced mammary tumors in the rat. From 3 to 4 months after DMBA administration, at a time when tumors had an average area of 3.76 ± .87 to 4.51 ± 0.89 cm², groups of animals were kept as intact controls or were ovariectomized (OVX) and received either Δ^5 -diol or DHEA for 24 days. While very few new tumors developed in OVX animals (average = 0.07 ± 0.07 per animal), an average of 0.40 ± 0.24 new tumor appeared in intact rats. In OVX animals treated with Δ^5 -diol or DHEA, the numbers were 1.85 ± 0.82 and 0.39 ± 0.17, respectively. An even more striking effect was observed on average tumor area that decreased from 4.70 ± 0.95 cm² in intact animals to 0.75 ± 0.27 cm² in OVX rats. Values of 9.79 ± 2.25 and 3.93 ± 0.86 cm² were found in Δ^5 -diol- and DHEA-treated animals, respectively. Δ^5 -diol and DHEA led to stimulation of the level of progesterone receptors in both the uteri and DMEA-induced tumors. The present data show that the two C₁₉ steroids Δ^5 -diol and DHEA can exert stimulatory effects analogous to those of estrogens on DMEAinduced tumor growth in the rat, thus suggesting a role of these adrenal steroids in breast cancer and other estrogen-sensitive diseases in the human.