SERUM LIPOPROTEIN CHANGES IN WOMEN OVER THE AGE OF 35 YEARS USING COMBINED ORAL CONTRACEPTIVES

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The reduction of steroid dose has led to the increase in safety of combined oral contraceptives (COCs) use. Since then modern "low-dose" formulations of COCs have been widely recommended not only for young women but also for women of late reproductive age.

Objective: To evaluate the effect of two COC formulations (30 g ethinylestradiol with 300 g norgestrel and 20 g ethinylestradiol with 150 g desogestrel) on serum levels of lipoproteins and apoproteins in women over the age of 35 years.

Methods: 45 women within the age interval of 35-49 years were enrolled in the study. Measurement of lipoproteins and apolipoproteins-A1 and apolipoproteins-B was performed at the commencement of the study and after 6 cycles of COCs use.

Results: There was a significant reduction in serum cholesterol and LDL-cholesterol levels in both groups. The coefficient of atherogenesis was decreased in both groups as well. There was a significant increase in HDL and apoprotein-A1 serum levels in women who used COC with 20 g ethinylestradiol and 150 g desogestrel $(1,77\pm0,15$ and $1,40\pm0,11$ mmol/l; p<0,05 and $2,37\pm0,14$ and $1,87\pm0,15g/l$; p<0,05 respectively). No increase in triglycerides was detected in this group $(1,46\pm0,19$ and $1,45\pm0,15$ mmol/l). **Conclusions:** COC formulation with 20 g ethinylestradiol and 150 g desogestrel has more beneficial effect on antiatherogenic lipoprotein profile as compared to COC formulation with 30 g ethinylestradiol and norgestrel as a progestagen component.

EFFECTS OF COMBINED ORAL CONTRACEPTIVES ON HEMOSTASIS IN WOMEN OF LATE REPRODUCTIVE AGE

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Objective: to investigate the effects of combined oral contraceptives (COCs) on hemostasis in women of late reproductive and perimenopausal age.

Methods: 64 women within the age interval of 35-49 years were accepted to the study. Three different formulations of COCs were used: 20 g ethinylestradiol with 150 g desogestrel, 30 g ethinylestrad with 150 g desogestrel and 30 g ethinylestradiol with 300 g norgestrel. 13 different parameters of hemostasis including resistance to activated protein C (APC resistance) and presence of Lupus anticoagulant were evaluated at the commencement of the study and after 2 and 6 cycles of COCs intake. The presence of Lupus anticoagulant was detected using tissue thromboplastin inhibition test. Test for identification of the factor V Leiden mutation based on polymerize chain reaction (PCR) was performed as well.

Results: There was a significant reduction of APC-ratio in heterozygous carriers of factor V Leiden as compared to women without this mutation $(2,1\pm0,1)$ and $(2,7\pm0,1)$; (2,0). The presence of lupus anticoagulant was identified more frequently in women with the factor (2,0). There was a significant reduction of APC-ratio in these women as well $(1,7\pm0,1)$ and $(2,1\pm0,1)$; (2,0). APC resistance was diagnosed in (2,0) of the participants without the factor (2,0) Leiden mutation before the treatment and in (2,0) after COCs use (2,0). The prevalence of APC-resistance was higher in COC users with Lupus anticoagulant as compared to those without this anticoagulant (2,0) - (2,0). Conclusions: Apparently APS-resistance and Lupus anticoagulant may be involved in the development of thrombophilia associated with COC use.