1. ABSTRACT

Charles University in Prague, Faculty of Pharmacy in Hradec Králové

**Department of Biological and Medical Sciences** 

Oxidative stress and endothelial condition in myocardium of mice expressing high levels of soluble

endoglin in plasma

Author of diploma thesis: Iveta Dusílková

Supervisor: PharmDr. Jana Rathouská, Ph.D.

Background: In recent years, soluble endoglin is among others considered as a marker of endothelial

dysfunction and plays a crucial role in many cardiovascular diseases. This study aimed to evaluate the

expression of specific markers of endothelium protection, inflammation and oxidative stress in cardiac

wall of transgenic model of mice with high plasma levels of human soluble endoglin, which was fed a

high fat diet.

Methods: For this work, we used a female transgenic mouse model on a CBAxC57BL/6J background

with high plasma levels of human soluble endoglin, fed a high fat (40%) and high cholesterol (1.25%)

diet. As a control group, we used their female littermates showing undetectable levels of human

soluble endoglin. The expression of the selected molecules eNOS, peNOS, VCAM-1, HO-1, SOD-3 and

catalase was evaluated by Western blot analysis. Total cholesterol levels were detected by biochemical

analysis. The levels of human soluble endoglin were detected by ELISA analysis.

Results: Biochemical analysis didn't show any significant difference in total cholesterol levels between

both groups of mice. Western blot analysis also failed to demonstrate significant difference in the

expression of selected proteins in the cardiac wall between the group of mice with high soluble

endoglin levels and the control group.

**Conclusion:** From these results, there is possible to suggest that high levels of soluble endoglin in

combination with high fat diet do not affect the expression of markers of endothelium protection,

inflammation and oxidative stress in cardiac wall of transgenic mouse strain. However, final

conclusions are the subject of ongoing studies.

Keywords: soluble endoglin, endothelial NO synthase, phosphorylated endothelial NO synthase,

vascular cell adhesion molecule 1, heme oxygenase 1, superoxide dismutase 3, catalase, high fat diet,

Western blot, oxidative stress, heart