

## CHAPTER 34

### MEDICAL SCIENCES PHARMACOLOGY

#### Doctoral Theses

309. MOHD. SHAHID  
**Effect of Remote Preconditioning on Myocardial Reperfusion Injury.**  
Supervisors : Dr. K. K. Sharma and Dr. M. Fahim  
Th 15375

#### *Abstract*

Demonstrates the protective effect of hind limb ischemia-induced preconditioning and to explore the possible mediators of remote preconditioning against myocardial ischemia/reperfusion injury. A new model of remote preconditioning by brief occlusion and reperfusion of femoral arteries was developed. To investigate the possible factors, which mediate preconditioning of the heart. A powerful protection against myocardial ischemia/reperfusion injury by RPC was obtained which is comparable to the previous classical IPC and other models of RPC. Protection was observed not only in terms of infarct size but also in myocardial mechanical and serum biochemical markers. Thus, brief occlusion and reperfusion of femoral arteries induces a strong preconditioning effect against myocardial ischemia/reperfusion injury in rats. This cardioprotective effect is mediated by the opening of mitoK<sub>ATP</sub><sup>+</sup> channels and increased NO synthesis. NO appears to be working upstream and is acting via activation of mito K<sub>ATP</sub><sup>+</sup> channels in this form of RPC. Indeed, exogenous NO donor, L-arginine could also trigger preconditioning of the heart which is as potent as established classical IPC and other forms of RPC. However, more systematic studies are required for information regarding the detailed molecular signaling cascade and temporal profile of RPC and minimum stimulus of ischemia of remote organ to induce potent cardioprotection.

1. Introduction. 2. Review of literature. 3. Materials and methods. 4. Results. 5. Discussion. 6. Summary and Conclusion. Bibliography
310. MOHAMMAD TAUSEEF  
**Evaluation of the Mechanism of Action of Aspirin as a Cardioprotective Agent in Experimentally Induced Hypercholesterolemic Rats**  
 Supervisors : Dr. K. K. Sharma and Dr. M. Fahim  
 Th 15374

*Abstract*

Testes the hypothesis that aspirin by its antioxidant effect, improves haemodynamic profile, baroreflex sensitivity and endothelial function in rat model of hypercholesterolemia. Hypercholesterolemia was induced in Wistar rats by feeding 1% cholesterol rich diet for 10 weeks. Demonstrates the vasoendothelial dysfunction during experimetnally induced hypercholesterolemia in rats. Hypercholesterolemic rats showed significant increase in total cholesterol, low-density lipoprotein-cholesterol, very low-density lipoprotein-cholesterol and atherogenic index, and significant decrease in high-density lipoprotein cholesterol. Significant rise in blood pressure, heart rate and attenuation of baroreflex sensitivity were found in hypercholesterolemic rat. Furthermore, hypercholesterolemic rats has also hown endothelial dysfunction and decreased sensitivity of  $\beta_2$  adrenoceptors in isolated aortic tissues, and free radical generation, evident by a significant increase in serum lipid peroxidation and significant reduction in serum reduced glutathione content. The study demonstrated that aspirin improves baroreflex response, prevents the rise blood pressure and heart rate, restore endothelial function and  $\beta_2$  adrenoceptors sensitivity possibly by reducing sympathetic activity due to its antioxidant effect in experimentally induced hypercholesterolemic rats.

*Contents*

1. Introduction. 2. Aims and objectives. 3. Review of literature. 4. Materials and methods. 5. Results. 6. Discussion. 7. Summary and Conclusion. Bibliography.

311. RISHI PAL

**Experimental Studies on the Role of Free Radicals in Emotional and Environmental Stress.**

Supervisors : Dr. Prof. A. Ray and Prof. B.D. Banerjee  
Th 15437

*Abstract*

Studies Pharmacological, Biochemical and Immunological techniques to evaluate the possible role of free radicals and antioxidant defense mechanisms in emotional and environmental (xenobiotic) stress in experimental animals.

*Contents*

1. Introduction. 2. Review of Literature. 3. Aims and Objectives. 4. Materials and Methods. 5. Results. 6. Discussion. 7. Summary and Conclusions. Bibliography and Appendix.