





Volume 282 Supplement 1 July 2015

Poster Sessions

Table of Contents

Poster Session 1

Sunday 5 July & Monday 6 July 08:30–19:30, Foyer Convention Center

- 56 Gen EX S1, Chromatin Structure and Epigenetic Modifications and Maintenance of the Genome
- 70 Gen Ex S2, Turning Signals into Messages the Complexity of Gene Regulation
- 89 Gen Ex S₃, Translational Control and Protein Turnover
- 98 Mem Biol S1, Organelle Dynamics and Communication
- 107 Mem Biol S2, Autophagy and Degradation
- 110 Mem Biol S₃, Redox-Regulation of Biological Activities
- 129 Chem Biol S1, Probing Cellular Function with Small Molecules
- 158 Chem Biol S2, Targeted Cancer Therapy
- 160 Chem Biol S4, RNA-Based Disease Mechanism and Therapy
- 166 Mol Neu S1, Neuronal Ion Channels and their Role in Disease
- 168 Mol Neu S2, Mechanisms of Nervous System Development and Regeneration
- 172 Mol Neu S₃, Degeneration and Ageing of the Nervous System
- 184 Sys Biol S2, Molecular Clocks
- 187 Sys Biol S₃, Comprehensive Models of Metabolism and Signaling
- 198 Struct Biol S1, Mechanisms of Membrane Transport
- 205 Struct Biol S2, Channels and Transporters
- 206 Struct Biol S3, Protein-Mediated Membrane Deformation and Penetration

Poster Session 2

Tuesday 7 July & Wednesday 8 July 08:30–19:30, Foyer Convention Center

- 209 Gen Ex S4, RNA Processing and Modifications
- 215 Gen Ex S5, Non-Coding RNAs in Gene Regulation
- 220 Mem Biol S4, Extrinsic and intrinsic regulation of cellular growth control
- 228 Mem Biol S5, Lipid Signaling & Dynamics
- 240 Chem Biol S2, Targeted Cancer Therapy
- 274 Chem Biol S₃, Functional Glycobiology from Mechanism to Disease
- 281 Chem Biol S5, Signal Transduction in Tumor Development,
 Differentiation and Immune Escape
- 293 Mol Neu S4, Molecular Architecture and Assembly of the Synapse
- 297 Mol Neu S5, Control of Neuronal Function by Regulating Protein Homeostasis
- 302 Sys Biol S1, Interspecies Communications
- 304 Sys Biol S4, Functional Networks Regulating Cellular Stress Response and Ageing
- 315 Sys Biol S5, Systems Biology in Stem Cells
- 316 Struct Biol S2, Channels and Transporters
- 324 Struct Biol S4, Monitoring Protein Conformational Dynamics and Movement
- 329 Struct Biol S5, Advances in Structural Biology from Subcellular to Molecular Resolution
- 353 FEBS Education Session
- 380 Late-breaking abstracts

P10-043

Modeling experimental atherosclerosis in rabbits for investigation of antioxidant proteins expression

L. Skvortsova¹, E. Khussainova¹, A. Ali¹, Z. Berkimbayeva¹, L. Djansugurova¹, A. Mansharipova², B. Bekmanov¹

¹Institute of General Genetics and Cytology, Almaty, Kazakhstan,

Ischemic heart disease (IHD) is the main cause of human mortality, lifetime and its quality worldwide. In most cases, anatomical "ground" of its development are fatty deposits in the coronary arteries walls (atherosclerosis), appearing during pathophysiological unbalance of lipid and carbohydrates metabolism. It is well established that oxidative stress and reactive oxygen species (ROS) play an important role in coronary atherosclerosis initiation and subsequent development with inflammation system involvement. Naturally occurring defense antioxidant system was noticed to protect against ROS aggressive and disruptive capabilities. However, involvement of many antioxidant proteins remains contradictory and uncompletedin coronary atherosclerotic processes.

In understanding of the basic processes of atherosclerotic pathogenesis, model objects as rabbits are the most suitable system, as biochemical parameters of lipid-carbohydrate metabolism

²Kazakh-Russian Medical University, Almaty, Kazakhstan