

(E-mail: sattarahmady@yahoo.com)

Proteinopathies or protein conformational diseases such as Alzheimer's, Parkinson's diseases and type II diabetes, are conditions that arise from the misfolding and aggregation of proteins in non-native conformations. Non-enzymatic mechanisms such as glycation or oxidation are a modifying factor that leads to proteinopathy, by affecting the structure and function of proteins which have essential role in diabetes. Identification of anti-glycation compounds is attracting considerable interest. In this study, deferiprone, deferasirox and desferal, three iron chelators used in the treatment of β -thalassemic patients, were chosen to explore their effects on the fructation of hemoglobin. Our results indicated that deferasirox cannot prevent AGE and carbonyl formation but it reduces the functional changes of hemoglobin, heme losses and helix depletion due to fructation. Deferiprone and desferal, on the other hand, prevent AGE formation and inhibit changes in the structure and function of hemoglobin during the fructation process.

Keywords: Glycation, Hemoglobin, Iron chelators, Proteinopathies.

Abstract No.215

Nanoflowers of Cobalt: Synthesis, Characterization and Application for the Electrochemical Oxidation and Determination of Sulfite and Nitrite

*Hossein Heli*¹, Iman Eskandari², Naghmeh Sattarahmady³,
Ali Akbar Moosavi-Movahedi⁴*

1. Laboratory of Analytical and Physical Electrochemistry, Department of Chemistry, Science and Research Branch, Islamic Azad University, Fars, Shiraz, IR
2. Department of Chemistry, Islamic Azad University, Firouzabad Branch, Firouzabad, IR
3. Department of Medical Physics, Shiraz University of Medical Sciences and Pharmaceutical Science Research Center, Shiraz University of Medical Sciences, Shiraz., IR
4. Institute of Biochemistry and Biophysics, University of Tehran, Tehran, IR
(E-mail: heli@ibb.ut.ac.ir)

Cobalt hexacyanoferrate (CoHCF) nanostructure was synthesized by anodic oxidation of metallic cobalt nanoflowers in a solution of $K_3Fe(CN)_6$. The synthesized CoHCF sample was then employed to prepare a modified carbon paste electrode. The modified electrode was characterized electrochemically in a phosphate buffer solution at

physiological pH. Two redox transitions were appeared in the voltammograms which were related to the redox processes of Co^{II}/Co^{III} and Fe^{II}/Fe^{III} in the solid state of CoHCF. The modified electrode was successfully applied to the electrooxidation of nitrite and sulfite and these substrates were oxidized electrocatalytically on the modified electrode surface via the active Fe^{III} species. The catalytic rate constants, the electron transfer coefficients and diffusion coefficients involved in the electrocatalytic oxidation of the compounds were reported. The modified electrode was applied to the amperometric determination of nitrite and sulfite.

Keywords: Cobalt hexacyanoferrate, Cobalt nanoflowers, Electrocatalysis, Modified electrode, Nanoflowers, Nitrite, Sulfite.

Abstract No.216

Measuring the Activity of Cytomegalovirus Promoter Using an in Vivo

*Fateme Mortazavi*¹, Masood Torkzadeh², Saman Hosseinkhani²,
Mokhtar Jalali Javaran²*

1. Biotechnology Department, Kerman Graduate University of Technology and International Center for Science High Technology and Environmental Sciences Kerman, Kerman, IR
2. Biochemistry Department, Tarbiat Modares University, Tehran, IR
(E-mail: fatimamortazavi@gmail.com)

Transgenic technologies are dependent upon genetic tools among which sufficiently strong promoters for the construction of expression genes are very important. Reporter genes are essential for the quantitative analysis of gene elements that potentially regulate gene expression. Several kinds of reporter genes have been developed and luciferase reporter gene is the most favored for functional analysis of promoters and enhancers, due to rapid, sensitive and reproducible assay system. In this study we investigated the activity rate of cytomegalovirus promoter of mammalian virus in model plant cell suspension of *Nicotiana tabacum* via firefly luciferase. The sequence of firefly luciferase (codon optimized luciferase gene LUC+) from pgl3 control vector was introduced in to plox vector via appropriate primers and two restriction enzymes ($BamH_1$ and Xho_1). The luciferase gene was cloned in Plox vector so cytomegalovirus promoter used to drive the expression of firefly luciferase in suspension cells of *Nicotiana tabacum*. The plox vector which contains luciferase reporter was transformed to cells. Measurement of luciferase activity was done in intact cells. Our result show that the cytomegalovirus promoter can be