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Abstract: This study evaluated the simple methods to improve the sorptional capability of polymeric sorbent to gold (III) ions. The polymeric sorbent is produced on the basis of phosphochlorinated industrial synthetic butadiene rubber SBR, as a result of the modification thereof by an oxidative chlorophosphorylation. In order to improve sorption efficiency of sorbents to gold ions, the non-covalent immobilization of organic sulfur containing compound to the polymer matrix was carried out. The gold embedded polymer matrix was characterized by SEM, EDX, XRD and visible absorption spectroscopy. The obtained results indicated that the polymeric sorbent contained metallic form of zero-valent gold. It was found that the average particle size of elemental gold crystals was 28, 31 nm. These results support that the sorption of gold ions by non-covalent organic sulfur compounds immobilizated polymer sorbent could be an alternative method for recovery of metallic zero-valent gold from gold solution.

Keywords: Sorption, Nanogold, phosphochlorinated polymer.

1. Introduction

Determination of gold ions in environmental and geological samples requires very often preconcentration and separation by polymer sorbents due to the high concentration of interfering matrix components and the low content of this metal. Solid phase extraction technique offers for this purpose high enrichment factor, rapid phase separation and the ability of combination with different detection techniques [1, 2]. The combined method of biosorption and incineration could be effective for concentration and recovery of gold from the solutions [3]. Nanogold composite materials have a large applications as nanoelectronic devices, catalyst,

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sensors, optical, anti corrosion materials and etc. [4, 5, 6, 7, 8, 9]. The biomaterials, carbon nanotubes, chitosans, activated carbons, resins, inorganic materials are used as a matrix in synthesis of gold nanoparticles by uptake of Au (III) ions from aqueous solutions. It is very clear that physicochemical properties of polymers (e.g. sensing and separation) are enhanced by anchored nanoparticles, while polymers provide excellent platforms for dispersing nanoparticles for e.g. high catalytic performances [10, 11, 12]. For nanogold produce is used chemical, physical, biochemical methods [13, 14, 15, 16, 17, 18]. Gold nanoparticles have been prepared chemically through the reduction of hydrogen tetrachloroaurate using suitable reducing agents in the presence of stabilizers. Several works have been published regarding the successful application of polymers with metal binding sites as stabilizers in the preparation of gold nanoparticles with a controlled size and narrow size distribution [19, 20]. The one of the main condition of the matrix selection is to be financally afordable. Here we report the uptake of gold ions from the aqua solutions using porous sorbent immobilized sulfur containing organic reagent which sensitive toward of gold. In the result of sorptional proses the gold nanoparticles formed in the sorbent phase. The synthesized polymer sorbent has a porous matrix which can used to form nanogold particles by sorption-reducing method.

2. Materials and Methods

The polymeric sorbent is produced on the basis of phosphochlorinated synthetic butadiene rubber SBR, as a result of the modification thereof by an oxidative chlorophosphorylation reaction method as reported [21]. Derived product has crosslinked structure, deep brown color, undissolved in acid, alkali and other organic solvents. The spherical of polymer granules have a porous structure with an average bead diameter 250-400 microns, which can sorb chemical compounds by physical adsorption such as Van der Waals forces, dipolar bonds.

The stock gold solution was prepared by dissolving 1g metallic gold in aqua regia ((HCl+HNO₃) (3:1)), following dilution with deonizined water to 1000 ml. Chloroauratic acid solutions in $[AuCl_4]^-$ ion form with an average concentration of 100-500 mq/l were prepared by diluting from the stock solution in 1 g/l concentration. Au(III) concentration after uptake of gold ions using polymer sorbent in the solution was determinated by Atomic absorption spectroscopy, at a wavelength of 242,4 nm.

Despite the fact that this sorbent is used for uptake of heavy metal ions due to having an phosphate acid groups, can make a property to sorb gold ions from aqous water. This concern has been achieved for non covalent immobilization as a result of treating of polymer sorbent with organic reagent which can interreact with gold ions. Thiourea is used as a immobilizated reagent. The immobilization was attempted in a batch process by mixing dried polymer cubes with thiourea solution heating up to 80° C. The thiourea immobilized

polymer cubes was then filtered, and dried at 50 ^oC under the vacuum to constant weight. The sorption of gold ions from an aqueous gold (III) solution was attempted in a batch process for 1 h at room temperature. After sorption process polymer is dried for 24 h at 50 °C. Residual amount of gold ions in solution were analyzed by Atomic absorption spectroscopy. SEM, EDX, XRD analyses are used for gold forming investigation in the polymer phase.

3. Results and Discussion

The adsorption efficiency was studied for varying the initial concentration, acidity, amount of sorbent, distribution coefficient at 22 ⁰C for an initial period of 60 min. The effective adsorption efficiency was obtained at pH 6,8. Increasing the concentration of the gold ions in the solution the adsorption efficiency increases which is describe as L type sorption isotherm (Fig. 1).



Fig. 1. The gold (III) adsorption isotherm

This behavior is most likely attributed to an instability of the produced compound species formed between gold (III) ion and the reagent immobilized polymer of the produced complexion assoside of $[AuCl_4]^-$ with the sorbent site of the thiourea noncovalent immobilized polymer. No significant increase in the efficiency was obtained with use of metal solution concentration beyond 0,8 mmol/l. The sorption efficiency stabilized after determined concentration. It can be attributed to the saturation of active sites on the polymer matrix. The value of maximal sorption capacity is calculated as a 0,647 mmol/g⁻¹.

Adsorption efficiency increases with increase in the sorbent amount from 0,01g to 0,350g. It can be attributed to an increase amount of immobilized reagent which can react with gold ions. No significant increase in the efficiency was obtained with use of sorbent beyond 0,350g. It can be attributed to molar matching between gold ion and the active species in polymer sorbent (Fig. 2).



Fig. 2. The effect of adsorbent dose

The distribution coefficient decreases with the increase concentration of gold ions from aqueous solution which can be attributed to the free movement limitation due to the metal ions overage in aqueous media (Table. 1).

Initial conc. of Au (III) ion	25	45	55	75	100	150
Eq.conc. of Au(III) ion	3,11	6,76	9,1	17,05	22,5	72,5
Distribution coefficient of Au(III) ion	14076	11313	10087	6797	6890	2137

Table 1. Calculated results of Au (III) distribution coefficient in solution.

The XRD pattern for the polymer sorbent after uptake of gold (III) ions from aqueous media is shown in Fig.3.



Fig.3 . X-ray diffraction pattern of nanogold embedded polymer matrix.

From the XRD pattern we get the d spacing values which are calculated from the observed reflections respectively. These peaks match well with the d spacing of pure metallic gold in cubic Fm3m space group. From the XRD pattern we indicate the characteristic diffraction peaks for nanogold particles which consist as

a single phase in the polymer matrix. The XRD pattern reveals the peaks with characteristic d spacing for pure metallic gold as obtained from JCPDS –International Center for Diffraction Data. The results are in good agreement with the literature report [22, 23, 24]. Elemental crystals size of the average price consisting gold particles were calculated by special program (Evaluation) on the basis of the X-ray analysis results. It was found that the average particle size of elemental gold crystals is 28,31 nm. The obtained results indicate that nanoparticles agglomerate and form to the gold particles in polymer matrix due to reduction of Au (III) ions by thiourea reagent.

Based on the obtained results and the date reported earlier, using thiourea non covalent immobilizated polymer sorbent for the uptake of Au (III) ions from aqua media, the pure gold particles embedded in nano form in the polymer matrix. Thus, the presence of pure metal-nanogold in the polymer matrix could be confirmed.

This result was also confirmed from the Scanning electron microscope (SEM), Energy Dispersive Spectroscopy (EDS) investigation(Fig. 4).



Fig. 4. SEM image and EDS spectra of gold particles embedded polymer matrix

Fig.4 shows the SEM image of the gold nanoparticles in polymer matrix. The SEM micrograph with of $3000 \times$ shows the presence of gold particles in the surface of the polymer matrix. A similar result has been reported in [25, 26]. In addition, energy dispersive X-ray (EDS) measurements and quantitative elemental analyze made on the sample shows the expected primary metal signals along with weak S signals, which suggests the presence of sulfur-containing reagent in reduction of gold. Gold nanoparticles were immobilized into the pores of polymer by shaking it with aqueous solution containing H[AuCl₄]⁻ complex.

These results support that the sequential process comprising of the loading of polymer sorbent followed by sorption of gold ions, could be an alternative method for recovery of metallic zero-valent gold from the gold solution.

4. Conclusion

The use of polymeric sorbent is able to sorbs not only heavy metals, and gold (III) ions as a result of the treating of thiourea solutions with the same sorbent. The polymeric sorbent is produced on the basis of chemical modification of industrial synthetic butadiene rubber SBR, as a result of the oxidative chlorophosphorylation reaction. The thiourea loaded polymeric sorbent permits uptake of gold (III) ions from the aquous solution effectively. The SEM, XRD, EDS investigations show that after sorptional process Au (III) ions converted to a gold nano particles in the pores of polymer sorbent. The developed method could be extended for further applications on the low-cost procedures for the quantitative uptake of gold ions and formation of gold nanoparticles in the polymer matrix. We contemplate that our produced nanocomposite material has a large application as a catalyst in chemical reactions, sensor, acidproof polymer materials. Work is continuing to investigate the influence of size of gold particles to the physic and chemical properties of nanocomposities.

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