# **ABSTRACTS**

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# Stable gold nanoparticles – synthesis, bioconjugation and application

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#### Introduction

Gold nanoparticles (GNPs) are good biocompatible materials due to their special physical and chemical properties<sup>1)</sup>. There is more and more research where GNPs are used in the medical imaging and in the cancer therapy. Nanotechniques, the usage and production of nanomaterials, have been developed in our team for many years<sup>2-4)</sup>. Furthermore, we synthesise various compounds which have anticancer activity and can be developed into novel anticancer treatment drugs<sup>5, 6)</sup>. Therefore, it is not surprising that we decided to combine these two paths of our studies.

Indeed, there are many examples described in the literature of chemically<sup>7, 9)</sup> and biologically<sup>8, 10)</sup> functionalized GNPs; however, these conjugates are very often instable. Although GNPs are rather stable within the period of many months, modified nanoparticles – conjugates GNPs to drug, dye or antibody are often instable within a long period, particularly in the presence of high salts and proteins, which is the essence of the human body<sup>8, 11)</sup>.

There exist some reports where glycerol is used as an efficient reducing agent<sup>12–14)</sup>. Glycerol is an environmentally friendly and completely biocompatible reagent. Thus we decided to use glycerol and to synthesise GNPs with various concentration of cross-linked polyvinylpyrrolidone (PVP) as the stabilizing agent. We also wanted to determine the size, shape, dispersity and

stable GNPs with diameters smaller than 60 nm because these GNPs have the most appropriate physical and biological properties<sup>1)</sup>. The next step will be coupling nanoparticles with our potential drugs. These bioconjugates will be examined in the *in vivo* biological studies.

stability of the obtained GNPs. Our aim was to obtain

# **Experimental methods**

# Preparation of gold nanoparticles

Gold nanoparticles were obtained using wet bottom-up approach. All glassware was cleaned in aqua regia, rinsed with water and dried prior to the use. All solution was prepared with ultrapure water. A mixture (10 ml) containing glycerol: H<sub>2</sub>O solution (1:4 v/v) and various amounts of cross-linked (i.e. *crospovidine*) PVP (0%, 0.015%, 0.03%, 0.05% and 0.1% w/v) were prepared. Then, 20 µl of tetrachloroauric acid (HAuCl<sub>4</sub>) was added (final concentration: 0.5 mM). After 2-hour preincubation, NaOH was added (1 mM) and the reaction was left to react overnight (16 hours). The mixture was kept in the room temperature and mixed during the whole process of preparation.

# Characterization of gold nanoparticles

The transmission electron microscopy (TEM) images of the resultant GNPs were obtained on a JEOL 2000 FX microscope and for GNP-conjugates they will be also obtained on a high resolution (HRTEM) JEM3010 microscope (both equipped with a EDS detector). The UV-Vis absorption spectra of the GNPs were obtained by using a UV-Vis V-530 spectrophotometer (Jasco). The size distribution profiles (by DLS) of GNPs were received by a Zetasizer Nano ZS (Malvern) (light source – 633 nm).

# Preparation of nanoparticles-drug conjugates

Gold nanoparticles-drug conjugates will be obtained by mixing GNPs (dissolved in a water:glycerol solution) and the drug solution in a proper ratio at room temperature.

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Conjugation time was previously fixed as 1 hour. Thiosemicarbazones will be the most common group amount of used drugs. They will be both obtained in our group of novel compounds and the others, investigated in clinical or preclinical trials such as Dp44mT or triapine. Because of their lack of solubility in water, thiosemicarbazones will be dissolved in DMSO.

# Biological tests in vivo

For GNP-drug conjugates the biological tests will be conducted. In this study the bare GNP and the drug alone will be used as the reference. The conducting of GNP-drug combination studies will provide a possibility to observe a potential synergistic effect. Biological studies of obtained nanoparticles and theirs conjugates will be carried out by using cell cultures and the MTT assay.

Two groups of cell line culture will be used. Firstly, the group of cancer cells consisting of HCT116 (human colon adenocarcinoma cells), CT26-CEA (murine colon cancer cell line with stable expression of carcinoembryonic antigen) and T41 (murine breast cancer cell line); secondly, the group of normal cells such as human fibroblasts (NHDF), hCMEC/D3 (human model of the blood brain barier) and RAW 264.7 (murine macrophages from blood). The comparison between these two groups of cells will make it possible to determine not only cytotoxicity but also therapeutic index of the examined conjugates.

All cell lines will be cultured in a proper medium (supplemented according to the requirements of each cell line) under standard conditions at 37 °C in a humidified atmosphere at 5% CO<sub>2</sub> and will be subcultured every 3–4 days as required. After a 24- or 48-h incubation with the drug, GNPs or their conjugate (in different concentrations), the MTT assay will be conducted.

### Results and discussion

The gold nanoparticles were obtained as described above. During GNPs preparation changes in the colour of the mixtures from light yellow to light red or even pinkish violet were observed. Metal nanoparticles (especially Ag, Au and Cu) exhibit a phenomena known as surface plasmon resonance. The volume (size) and shape of a nanoparticle determines, *via* interactions with light, the observed colour of the GNPs. A change of size induces a shift of the peak maximum location and a change of shape (e.g. spherical nanocrystal into nanorode) causes the formation of new band at a longer wavelength. Figure 1 shows an exemplary series of changes in the UV-Vis spectrum during the reaction for the sample with 0.03% of PVP concentration.

There were five versions of the experiment, where different amounts of the stabilizing agent (PVP) were added. For each version of the experiment, size distribution profiles by dynamic light scattering (DLS) phenomena and TEM images were obtained. The data from these two techniques are shown in Table 1 and in Figure 2.

As it can be observed, PVP was found to

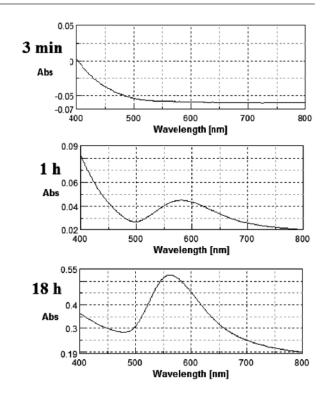


Fig. 1. Change in UV-Vis spectrum for 0.03% PVP concentration

be a very sufficient stabilizer of GNPs. Even with a small concentration of PVP (0.015%), gold nanoparticles were created. However, PVP was proved to be indispensable for obtaining nanoparticles in a similar size (i.e. monodisperse) (Fig. 2). Moreover, without the stabilizing agent, the reduction of gold proceeded with very low efficiency, which could be seen in the UV-Vis spectrum (data not shown). Otherwise, PVP with the highest concentration (0.1%) seems to be inappropriate for the preparation of monodisperse nanoparticles (Fig. 2). It is probably because of the incorrect ratio between the polymer and gold ions - the arising nanoparticle is randomly included into the PVP structure, which stops further growth of GNP. This hypothesis seems to be confirmed by TEM images (not shown). Therefore, for the next GNPs preparation experiments three middle concentrations of PVP (0.015–0.05%) will be chosen.

Although we managed to obtain monodisperse gold nanoparticles, the size of all obtained nanoparticles occurred to be too big for further medical application and investigation, where GNPs smaller than 60 nm are required. Thus, changes in gold nanoparticles preparation are necessary. The first option to be tested is to use the straight PVP polymer (povidone) instead of a cross-linked

Table 1. Summary of GNPs size at different PVP (cross-linked) concentrations

No.	PVP	Average particle size	Dispersity
	(% w/v)	by DLS (nm)	
1	0	none	polydispersity
2	0.015	230	nearly monodispersity
3	0.03	340	monodispersity
4	0.05	320	monodispersity
5	0.1	420/ 369/308/137	polydispersity

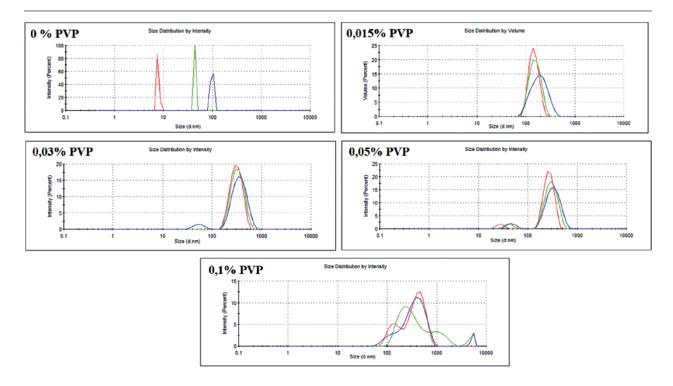


Fig. 2. Dispersity of the obtained GNPs with different PVP concentration (obtained with DLS technique)

one (*crospovidone*). Other biocompatible polymers such as polyethylenimine (PEI) or poly-L-lysine (PLL) should also be investigated.

#### **Conclusions**

In this research, stable gold nanoparticles were synthesised using polyvinylpyrrolidone (PVP) as the stabilizing agent. Then, using many different techniques, size, shape and dispersity of nanoparticles were investigated. The presented results suggest that the most appropriate concentration of cross-linked PVP for GNPs preparation lies within 0.01–0.05% range. The obtained data showed in which way these investigated physical properties depend on the concentration of polymers. However, further examination is necessary because of the fact that we did not manage to obtain gold nanoparticles in the required size.

In further studies, stable gold nanoparticles of an appropriate size will be conjugated with several approved or potential anticancer drugs. The chosen promising conjugates will be examined in a basic anticancer *in vivo* study with different cell lines.

These researches seem to be interesting and helpful in the design and improvement of medical application for nanoparticles.

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# Conflicts of interest: none.

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