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Abstract

The preparation, characterization and the magnetic properties of gold and gold–iron oxide glyconanoparticles (GNPs) are described. Glyconanoparticles were prepared in a single step procedure in the presence of aqueous solution of thiol functionalized neoglycoconjugates and either gold salts or both gold and iron salts. Neoglycoconjugates of lactose and maltose disaccharides with different linkers were used. Iron-free gold or gold–iron oxide GNPs with controlled gold–iron ratios were obtained. The average core-size diameters are in the range of 1.5–2.5 nm. The GNPs are fully characterized by 1H NMR spectrometry, transmission electron microscopy (TEM), and UV–vis and X-ray absorption (XAS) spectroscopies. Inductive plasma-atomic emission spectrometry (ICP) and elemental analysis gave the average number of neoglycoconjugates per cluster. The magnetic properties were measured in a SQUID magnetometer. The most remarkable results was the observation of a permanent magnetism up to room temperature in the iron-free gold GNPs, that was not present in the corresponding gold–iron oxide GNPs.

Introduction

The application of methodologies to produce nanoparticles with bioresponsive properties has opened the way for producing useful tools for molecular diagnostics, therapeutics, and biotechnology.1–6 Metal, semiconductor, and magnetic colloidal nanoparticles are presently under intensive study for potential applications. The development of functional nanobioconjugates is mainly based on proteins and nucleic acids. However, carbohydrates as biologically important molecules have not received as much attention as protein- and gene-oriented nanotechnology.

We recently prepared, in a single step, sugar-protected gold nanoclusters. These tools are gold glyconanoparticles (GNPs) incorporating carbohydrate antigens for studying carbohydrate-mediated interactions.7–9 The dispersion of these glyconanoparticles are stable for months in physiological solutions, and they have been shown to be excellent platforms for basic studies of carbohydrate interactions10,11 and potential tools for biotechnological and biomedical applications.12 In addition, they have an exceptionally small core size, which confers on them singular physical properties as materials (due to quantum confinement effects).

In an attempt to confer magnetism to these biofunctional gold nanoparticles and obtain new contrast agents based on stable nanoparticles protected with carbohydrates, we prepared gold–iron oxide glyconanoparticles by modification of the procedure used for preparation of the gold glyconanoparticles8 and studied the magnetic behavior of AuFe GNPs. For the sake of comparison, pure Au GNPs were also measured.

Synthesis of magnetic nanoparticles with size control and suitably coated with polymers or other coating material to be dispersed into water is of interest for cell labeling and separation and for magnetic resonance imaging.13–17 Commercially available iron oxide particles have been used in cell sorting and separation.18 Magnetic nanoparticles coated with cross-linked dextran and functionalized with peptides and oligonucleotides have been prepared for intracellular labeling19,20 and as nanosensors.21 Biofunctional small (SPIO) and ultra-small (USPIO) iron oxide nanoparticles have been also prepared and used as a contrast agent in cellular labeling.20 However, synthesis of magnetic particles with the desired size that are highly stable in aqueous solution without undergoing aggregation after some time and able to target a determined tissue or organ constitutes, even today, a problem to be solved. Surfactant-protected gold–iron core–shell nanoparticles have been recently synthesized by means of reverse micelles.22 Attachment of biologically active molecules to this type of nanoclusters has, however, not been described.

We now report on a straightforward synthesis in aqueous solution of exceptionally small gold–iron oxide magnetic glyconanoparticles (AuFe–GNPs), which form stable colloidal solutions in water (ferrofluids) and present the study of their properties in comparison to those of the pure gold glyconanoparticles (Au–GNPs) previously prepared in our laboratory.8 The singular (magnetic) properties found in both pure gold and gold–iron GNPs are also described.

Experimental Section

General Methods. All commercially available chemicals were used without further purification. Hydrogen tetrachloro-
aurate tetrahydrate, sodium tetrahydroborate, iron trichloride, lactose, and maltose were obtained from Aldrich. UV spectra were obtained by a UV/vis Perkin-Elmer Lambda 12 spectrophotometer. 1H NMR spectra were acquired on a Bruker DRX-300 or DRX-300 spectrometer, and chemical shifts are given in ppm (δ) relative to D2O. TEM examination of the samples was carried out at 200 kV with a Philips CM200 microscope. For the TEM characterization, a 10 μL drop of the aqueous solutions (0.1 mg/mL) of the glyconanoparticles was placed onto a copper grid coated with a carbon film. The grid was left to dry in air for several hours at room temperature. Particle size distributions of the nanoclusters were evaluated from several micrographs using an automatic image analyzer. EDX analysis was performed with a Philips DX4 spectrometer attached to the microscope. Elemental analysis was carried out in a LECO CHNS-932 apparatus after drying analytical samples under vacuum for 24 h. ICP analysis was performed in an Iris Advantage apparatus (Termo Jarrell Ash, Franklin, MA).

AFM measurements were made with a Topometrix explorer AFM (Veeco Instruments, Sunnyvale) in noncontact mode in air. The noncontact mode was used to avoid any compression of the sample that may have occurred with contact mode AFM. Rectangular silicon cantilevers with integrated tips of spring constant 4.5 N/m (MikroMasch, Tallinn, Estonia) were used, oscillating at just below their natural resonant frequency (normally 150 kHz but measured for each cantilever) throughout the work. Untreated glyconanoparticles were prepared for imaging by dissolving in pure water (18 MΩ, Sigma-Aldrich, U.K.). Glyconanoparticles were deposited onto freshly cleaved mica from dilute (70 μL·mL−1) solution (5 μL per sample) and left to air-dry.

The magnetization measurements were done in a SQUID magnetometer (Quantum Design MPMS 5). The powder samples were fixed to the sample holder by means of a Kapton tape. Due to the low magnetization of the sample, an accurate determination of the magnetization was performed by measuring the sample holder and the Kapton tape used in each measurement. The magnetic signal (a diamagnetic one) arising from both, the sample holder and the Kapton tape used in each measurement, could no longer be detected by thin-layer chromatography (TLC). The residue was dispersed in water and centrifuged to eliminate insoluble materials, mainly gold and iron oxide precipitates. The brownish-clear solution was lyophilized, and the lacto(1-AuFe) GNPs obtained were free of salts and starting material (absence of signals from disulfide and Na+ ions in 1H and 23Na NMR spectroscopy). Average diameter and number of gold atoms: 2 nm, 314. 1H NMR (500 MHz, D2O): δ 4.49 (brd, 1H, H-1′), 4.40 (brs, 1H, H-1), 4.10−3.30 (m, 23H), 2.92 (m, 0.5H). UV (H2O, 0.3 mg·mL−1): λ 500 nm, broad band. ICP: 51% Au, 2.8% Fe. Average formula: (C20H37O14S)95Au266Fe48.

**Preparation of Maltose Gold—Iron Oxide Glyconanoparticles Malto(2-AuFe).** To a solution of the disulfide 2 (80 mg, 0.075 mmol, 3 equiv) in methanol (11.5 mL) a solution of FeCl3 (2 mg, 0.013 mmol, 0.25 equiv) in water (0.5 mL) was added followed by addition of a solution of HAuCl4 (17 mg, 0.05 mmol, 1 equiv) in water (2 mL). The mixture was left 5 min at room temperature, and then an aqueous solution of NaBH4 (1 M, 52 mg, 1.38 mmol, 27.5 equiv) was added in small portions (250 μL) with rapid stirring. The brown suspension formed was stirred for an additional 2 h, and the solvent was removed under vacuum. The brown residue was dispersed in NANOpure water (~15 mL), loaded into a centrifugal filter device (CENTRIPLEX YM30, MICROCON, MWCO = 30 000), and subjected to centrifugation (3000g, 40 min). The dark glyconanoparticle residue was first washed with MeOH (5 mL) and then with water several times (8 × 5 mL) until the excess amount of neoglycoconjugate could no longer be detected by thin-layer chromatography (TLC). The residue was dispersed in water and centrifuged to eliminate insoluble materials, mainly gold and iron oxide precipitates. The brownish-clear solution was lyophilized, and the malto(2-AuFe) GNPs obtained were free of salts and starting material (absence of signals from disulfide and Na+ ions in 1H and 23Na NMR spectroscopy). Average diameter and number of gold atoms: 1.6 and 2.5 nm, 140 and 586. 1H NMR (500 MHz, D2O): 0.532 (s, 1H, H-1′), 4.37 (s, 1H, H-1), 4.00−3.30 (m, 13H), 2.70 (s, 2H, CH2S), 1.85−1.20 (m, 17H). UV (H2O, 0.3 mg·mL−1): λ 500 nm, broad band. ICP: 28% Au, 0.27% Fe.

**Preparation of Maltose Gold—Iron Oxide Glyconanoparticles Malto(2a-AuFe and 2b-AuFe).** To a solution of the disulfide 2 (100 mg, 0.094 mmol, 3 equiv) in methanol (14 mL) a solution of FeCl3 (2.5 mg, 0.01 mmol, 0.25 equiv) in water (0.63 mL) was added followed by addition of a solution of HAuCl4 (21 mg, 0.04 mmol, 1 equiv) in water (2.5 mL). The mixture was left 5 min at 60 °C, and then an aqueous solution of NaBH4 (1 M, 65 mg, 1.7 mmol, 27.5 equiv) was added in small portions with rapid stirring. The brown suspension formed was stirred for an additional 2 h, and the solvent was removed under vacuum. The brown residue was dispersed in NANOpure water (~15 mL), loaded into a centrifugal filter device (CENTRIPLEX YM30, MICROCON, MWCO = 30 000), and subjected to centrifugation (3000g, 40 min). The dark glyconanoparticle residue was first washed with MeOH (5 mL) and then with water several times (8 × 5 mL) until the excess amount of neoglycoconjugate could no longer be detected by thin-layer chromatography (TLC). The residue was dispersed in water and centrifuged to eliminate insoluble materials, mainly gold and iron oxide precipitates. The brownish-clear solution was lyophilized, and the malto(2a-AuFe) and Malto(2b-AuFe) GNPs obtained were free of salts and starting material (absence of signals from disulfide and Na+ ions in 1H and 23Na NMR spectroscopy). Average diameter and number of gold atoms: 1.6 and 2.5 nm, 140 and 586. 1H NMR (500 MHz, D2O): 0.532 (s, 1H, H-1′), 4.37 (s, 1H, H-1), 4.00−3.30 (m, 13H), 2.70 (s, 2H, CH2S), 1.85−1.20 (m, 17H). UV (H2O, 0.3 mg·mL−1): λ 500 nm, broad band. ICP: 28% Au, 0.27% Fe.
Fe. Anal. Calcd for (C_{23}H_{44}O_{11}S)_{191}Au_{191}Fe_{10}: C, 37.9; H, 5.9; ppm.

but avoiding the use of Fe-contaminating material. ICP analysis starting material (absence of signals from disulfide and Na+ the precipitates. The brownish-clear solution was lyophilized, and eliminate insoluble materials, mainly gold and iron oxide (TLC). The residue was dispersed in water and centrifuged to tailor glyconanoparticles with a variety of carbohydrate ligands and very useful magnetic properties for future applications as ferrofluids in molecular imaging and tumor targeting. The so-prepared glyconanoparticles have less than a 2 nm diameter core. They formed stable dispersions, and even after several months, flocculation or aggregation was not observed.

Thiol-functionalized neoglycoconjugates of the disaccharides lactose (lacto, Galβ1–4Glcβ1-OR) and maltose (malto, Glcα1–4Glcβ1-OR) have been synthesized to prepare the magnetic nanoparticles in situ (Scheme 1). The disaccharides lactose and maltose were chosen to compare the gold–iron oxide with the pure gold lacto(1-Au) and malto(2-Au) GNPs previously prepared.8 The synthesis of lactose and maltose neoglycoconjugates 1 and 2 was carried out as previously described by glycosylation of the conveniently protected sugar derivatives with 11-acylthio-3,6,9-trioxa-undecanol and 11-acylthio-undecanol, respectively.8 Different linkers have been used to test the influence of their hydrophilic or hydrophobic nature in the properties of the whole material. Neoglycoconjugates 1 and 2 were isolated as disulfide forms and used in this form for the preparation of AuFe-protected glyconanoparticles. A water/methanol solution of FeCl3 and HAuCl4 in a ratio of 1:4 was reduced with NaBH4 in the presence of disulfides 1 and 2 to give stable dispersions of lacto(1-AuFe), and malto(2-AuFe), and malto(2b-AuFe) GNPs in a single step. Pure gold lacto(1-Au) and malto(2-Au) GNPs (Fe content below 0.01 ppm) were also prepared, and special care was taken to avoid contamination by iron-containing materials.

The 1H NMR in deuterated water, TEM micrographs, and the corresponding particle size distribution histograms for the AuFe GNPs are shown in Figure 1. The corresponding data for the Au GNPs were already described.8 For the lactose glyconanoparticles, lacto(1-AuFe), the corresponding histogram gives a mean diameter for the gold–iron oxide cluster of 2 nm, which corresponds to an average number of 314 atoms (Figure 1a). In the case of malto(2-AuFe) two different fractions could be obtained depending on the isolation procedure (see Experimental Section). In the first procedure a mixture of two different nanostructures could be observed on the micrographs: dispersed nanoclusters and areas with linearly ordered nanoparticles (Figure 1c). The corresponding histogram gave a bimodal particle size distribution centered at 1.6 and 2.5 nm. The two types of nanostructures could be isolated separately following the second procedure. A methanol-insoluble brown residue was isolated which corresponds to malto(2a-AuFe) GNPs. The micrographs have a particle size distribution centered at 1.8 nm,
as indicated by the corresponding histogram (Figure 1e). From
the methanol solution a white residue was obtained that was
not detected in the methanol solution of the lacto(1-AuFe). The
isolated white solid, malto(2b-AuFe), also formed water-stable
dispersions and contains very small AuFe nanoparticles perfectly
aligned and encapsulated in what seems to be organic material
as indicated by the TEM micrographs. The corresponding
histogram gave a particle size distribution of less that 1.5 nm
diameter, and the distance between the ordered lines is around
3.4 nm (Figure 1g). Elemental analysis of malto(2b-AuFe)
indicates that more than 2000 maltose molecules surrounded
the AuFe cluster. The amphiphilic character of the maltose
neoglycoconjugate 2 (with a lipophilic spacer C11) may be the
origin of these supramolecular aggregates in water solution.
Neoglycolipids such as n-dodecyl β-maltoside, n-octyl β-glucoside, and phenyl-alkyl β-glucosides are known to form micelles and other types of aggregates. Encapsulation of Au nanocrystals in a nanotube formed by a β-glucolipid has been recently described. Capsules were not observed for lacto(1-AuFe). Neoglycoconjugate 1 has the much less lipophilic linker tetra(ethylene glycol) and therefore is not expected to behave in an amphiphilic manner.

Elemental and inductive coupled plasma-atomic emission spectrometry (ICP) gave an average composition of 95 lactose molecules, 266 gold atoms, and 48 iron atoms for lacto(1-AuFe), 190 maltose molecules, 190 gold atoms, and 10 iron atoms for malto(2a-AuFe), and 2050 maltose molecules, 67 gold atoms, and 10 iron atoms for malto(2b-AuFe), which corresponds to an average Au:Fe ratio of 5:1, 20:1, and 6:1, respectively. In the case of malto(2b-AuFe) the 2000 maltose molecules in excess accounts for formation of the organic nanocapsules.

Atomic force microscopy (AFM) images confirm this encapsulation (Figure 2). These images strongly differ from those usually obtained for isolated lacto(1-Au) and malto(2-Au) glyconanoparticles. In addition to small particles of mean height 1.3 nm, broad flat aggregates of around 6, 12, or 18 nm height were observed which correspond to the distance of two, four, or six consecutive maltose neoglycoconjugates, indicating that growth of these aggregates is controlled by the maltose neoglycoconjugate. These results, along with the elemental analysis, suggest the presence of two components in the sample: the organic neoglycoconjugate aggregates and the nanoparticles with a metal core.

The UV/vis spectra of these GNPs show weak surface plasmon absorption around 500 nm. The absorption bands of lacto(1-AuFe) and malto(2a-AuFe) (Figure 3a and 3b) strongly decrease in comparison with the absorption spectrum of 10 nm gold nanoclusters. This is probably due to the very small dimension of the nanoclusters (below 2 nm). However, it is worth highlighting that the surface plasmon band of malto(2b-AuFe) increases (Figure 3c), probably due to the short distance between the metallic glyconanoparticles along the lines (Figure 1g).

Figure 4 shows the XANES spectra of the Au L3-edge and the Fe K-edge for lacto(1-AuFe) and malto(2-AuFe). The XANES region of the XAS spectra contains information about oxidation states and local structure around the absorbing atom, and it can also be used like a fingerprint technique to obtain results only by comparison with references samples. A threshold resonance at the Au L3 absorption that is just below the one for the bulk gold was observed (Figure 4a). In the case with particles 2 nm in diameter, the lattice contraction effect was almost totally compensated by the charge transfer to the capping molecules. This effect was also observed in the case of malto(2-Au) samples. Figure 4b shows the spectra at the Fe K edge for the lacto(1-AuFe) and malto(2-AuFe) samples together with the spectra measured for a metallic Fe foil, γ-Fe2O3, and FeO4 references. It is clear that the spectra of the samples are similar to the oxide ones, and furthermore, the intensity of the features at 7113 eV and of the white line at 7132 eV seems to indicate that the Fe atoms contained in the sample are more similar to the Fe2O3 structure.

High-resolution TEM was performed to better characterize the structure of the inorganic core. However, after several trials we were not able to obtain high-resolution micrographs of these samples, probably due to the small size of the nanoparticles and the large amount of organic material covering and protecting the inorganic cores, making these measurements extremely difficult. Nevertheless, due to the low amount of iron oxide included in the inorganic cores (2.8–0.04% Fe) it is improbable to have a core–shell structure. Iron must be dissolved into the gold phase.

Magnetic characterization of the gold and gold–iron GNPs reveals singular properties in these nanoclusters. Magnetization curves as a function of the applied field at different temperatures as well as the dependence of magnetization with the temperature have been measured for the gold–iron oxide and iron-free gold GNPs. It was very remarkable to observe a ferromagnetic-like behavior not only for the AuFe GNPs but also for the pure gold GNPs (Figure 5). In the AuFe samples a ferromagnetic behavior...
at 5 K with small hysteresis cycle was observed (Figure 5a, 5c, and 5e). The coercive field values at 5 K were around 15, 45, and 80 Oe for lacto(1-AuFe), malto(2a-AuFe), and malto(2b-AuFe), respectively. The samples were far from being saturated, with magnetization values at 1 T of around 0.125 (0.24 emu/g AuFe), 0.044 (0.16 emu/g AuFe), and 0.008 emu/g (0.64 emu/g AuFe) for lacto(1-AuFe), malto(2a-AuFe), and malto(2b-AuFe), respectively. These values were obtained after subtracting the diamagnetic contribution of the sample holder (Kapton). As shown in Figure 5f, both the measurements of the Kapton tape and the Kapton plus the sample are completely different. In the case of the malto(2b-AuFe) sample, the diamagnetic contribution associated with the organic component has been subtracted for determining the magnetization of the nanoparticles. Although the Fe content determined in the samples (3.0%, 0.4%, and 0.04% for lacto(1-AuFe), malto(2a-AuFe), and malto(2b-AuFe), respectively) would account for the magnetization values, it is difficult to attribute the magnetic hysteresis observed to the presence of iron, in particular to Fe$_3$O$_4$. For instance, the blocking temperature of 5 nm Fe$_3$O$_4$ particles is around 45 K. From this value, and assuming spherical particles, it can be estimated that 2 nm pure Fe$_3$O$_4$ nanoparticles would exhibit a blocking temperature of around 3 K. Taking into account that in our case Fe$_3$O$_4$ is the minor component of the nanoparticles, a blocking temperature lower than 3 K should be expected. As can be seen in Figure 5, all AuFe GNPs exhibit hysteresis at 5 K and, in particular, the one with the lower Fe content (malto(2b-AuFe)) exhibits hysteresis at 300 K.

Even more unexpected was the ferromagnetic behavior observed for the iron-free lacto(1-Au) and malto(2a-Au) GNPs with coercive field values of 100 and 125 Oe higher than those observed for the AuFe GNPs (Figure 5b and 5d). By increasing the temperature the magnetic hysteresis in lacto(1-AuFe) and malto(2a-AuFe) GNPs disappears and a paramagnetic behavior is observed, whereas a ferromagnetic component remains even at room temperature for malto(2b-AuFe) (Figure 5e). This result is quite unexpected since, as mentioned previously, the Fe content in lacto(1-AuFe) GNPs is much higher than that in malto(2a-AuFe), suggesting that the origin of the magnetic properties of the systems cannot be merely attributed to the presence of iron oxide. Moreover, the average size of malto(2a-AuFe) nanoparticles is smaller than that of the lacto(1-AuFe) nanoparticles; therefore a “weak” ferromagnetic behavior could be expected. In addition, if the magnetization is referred to the weight fraction of gold atoms which are bonded to the thiol group, instead of to the total amount of sample, the magnetization values obtained become similar and around 0.1 emu/g Au bonded for both set of samples, suggesting that this ferromagnetic behavior can be attributed to the Au content instead of to the Fe content. This is confirmed by the magnetic hysteresis observed even at room temperature in the iron-free lacto(1-Au) and malto(2-Au) GNPs (Figure 5b and 5d). Observation of a permanent magnetism in the pure gold GNPs was so unexpected (mannose and other sugar thiol-capped GNPs present also this magnetism) that the magnetic properties of thiol-functionalized gold nanoparticles were compared with detergent-protected gold nanoparticles. Gold nanoparticles with commercially available 1-dodecanethiol and an average size of 1.5 nm were prepared. The results confirmed that a ferromagnetic behavior was also presented even up to room temperature only for the thiol-protected Au nanoparticles. We proposed two factors to be in the origin of the ferromagnetic-like behavior: the small particle size (below 2 nm) and the change on the
electronic structure of the Au nanoparticle due to the covalent Au–thiol bond. From XANES data it was inferred that this behavior is related with a strong charge transfer between the Au atom and the S atom, localized 5d holes generated through the Au–S bond. The d-charge loss has been estimated to be a maximum as 0.05 e/atom, a low value that is in agreement with the low values of the measured magnetization. This effect seems to also be in the origin of the magnetic behavior observed in AuFe nanoparticles reported here. The fast decrease of the magnetization at low temperatures could be attributed to the small amount of Fe present in the nanoparticles in the lacto(AuFe) and malto(2a-AuFe) (Figure 5a and 5c). The hysteresis observed at low temperatures in the AuFe GNPs is more likely attributable to the surface Au atoms which are bonded to the thiol group.

The origin of permanent magnetism in thiol-protected Au GNPs is still unclear, and many questions still remain. A more detailed study to understand the differences in the magnetic behavior of AuFe and Au–GNPs is presently underway.

Conclusions

In conclusion, we developed a simple methodology to prepare exceptionally small, water-dispersible, and stable magnetic nanoparticles covalently linked to oligosaccharides. The methodology can be extended to the preparation of hybrid nanoparticles incorporating both carbohydrates and other molecules (fluorescence probes, biotin, etc.). Carbohydrate–receptor interactions can direct the magnetic glyconanoparticles to target cells and tissues, allowing their selective labeling. This kind of polyvalent magnetic glyconanoparticles complements the already available bioactive iron oxide magnetic nanoparticles. However, the most striking fact of these results is the singular observation of a permanent magnetism in the pure gold GNPs with coercive fields higher than those of the gold–iron oxide GNPs. The easy preparation and purification, their small core size and stability, and their dispersability in physiologically relevant conditions together with their exceptional magnetic properties make these particles potential candidates for diagnostic, tumor targeting, or magnetic resonance imaging applications. The response of different cellular lines against these magnetic glyconanoparticles and their longitudinal and transversal relaxation times, $T_1$ and $T_2$, in buffer and plasma are presently being determined.

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References and Notes

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