

Role of the nanocrystallinity of the chemical ordering of $\text{Co}_x\text{Pt}_{1-x}$ synthesized by wet chemistry

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Abstract

$\text{Co}_x\text{Pt}_{1-x}$ nanoalloys have been synthesized by two different chemical processes either at high or at low temperature. Their physical properties and the order/disorder phase transition induced by annealing have been investigated depending on the route of synthesis. It is demonstrated that the high temperature chemical synthesis allows stabilization the fcc structure of the native nanoalloys the soft chemical approach yield mainly to poly or non crystalline structure. As a result the approach of the order/disorder phase transition is strongly modified as observed by in situ annealing HR-TEM studies. The control of the nanocrystallinity allows to decrease significantly the chemical ordering temperature as the ordered structure is observed at 420 °C. This allows preserve the nanocrystal from the coalescence usually observed during the annealing.

A. Introduction

Metallic nanoalloys will initiate important development in nanotechnologies due to their specific chemical and physical properties (i.e. in catalysis, magnetism, optics, etc) [1,2]. It is now well known that these properties are mainly controlled by the fine tuning of structural parameters such as the size, the bimetallic composition and segregation processes [3]. Concerning their fabrication, the bottom up approach, either physical or chemical, is ideal to design this specific class of nanomaterials due to its versatility, facility and low cost. However, the realization of well-controlled bimetallic nanoparticles is not always straightforward from the know-how developed for monometallic nanoparticles.

In a sustainable approach, wet chemistry is well-adapted to produce such nanoalloys in large amount [4]. However at the nanometer scale, as the properties are strongly dependent on the size and the surface state (raw or passivated), it is crucial to develop method where the crystallinity, the polydispersity in size and composition are finely controlled. Nevertheless, depicted the large amount of work made on the synthesis of nanoalloys by the chemical way [1,2], there are still open questions considering the control of composition and especially the segregation process. For example, in the chemical approach, nanoparticles are always passivated by an organic molecule and mainly obtained in the disordered A_1 phase where both metals are randomly dispersed in the crystalline lattice (Figure 1A).

Among the nanoalloys, magnetic nanocrystals (NCs) are promising materials due to their strong potential in the development of applications especially for high-density data storage [5-8]. In this last domain, bimetallic alloys such as CoPt, FePt or CoRh represent a particularly interesting class of materials for the improvement of the recording on magnetic storage systems. Indeed, alloys such as CoPt (or FePt) have an ordered crystalline phase ($L1_0$) around the equi-atomic composition, which is intrinsic to the tetragonal symmetry (*fcc*) of the crystal structure (Figure 1C) and an ordered fcc structure ($L1_2$) form composition around Co_3Pt or $CoPt_3$ (figure 1D) [9,10]. The ordered $L1_0$ phase of the CoPt system is of particular interest thanks to the high coercivity (10 kOe) and the high magnetocrystalline anisotropy (4.9×10^6 J/m³). In fact, CoPt NCs in the $L1_0$ phase have large uniaxial magnetic anisotropy energy and so have the potential to exceed the superparamagnetic limit [11,12]. Moreover, these CoPt magnetic NCs are monodomain magnetic particles for a typical size below 10 nm [13]. Beyond this typical size the nanocrystals become polydomain magnetic and this leads to domain wall formation [14]. In the case of $CoPt_3$, the $L1_2$ phase present also a very magnetocrystalline anisotropy (2×10^6 J/m³). Therefore, high control of the chemical

composition and size of the alloy NCs is essential for optimizing the magnetic nanoscale behavior.

Wet chemistry is a very useful method to obtain large amount of NCs, however, CoPt NCs synthesized by colloidal route have an A_1 -disordered structure. A thermal annealing is required to ordered it toward $L1_0$ or $L1_2$ structure, depending on the composition [15]. For supported nanoparticles, the thermal assistances at high temperature ($> 300^\circ\text{C}$) can lead to coalescence and sintering effects, which increase the size, modify the shape and destroy their organization [15, 16]. Indeed, the coalescence driving force and the change of shape is the minimization of surface energy by elimination of interfaces and appearance of grain boundaries and defects [15]. Thus, the elaboration of well-defined nanoparticles in the $L1_0$ or $L1_2$, ordered phase remains very difficult. We report here a new approach for the synthesis of $\text{Co}_x\text{Pt}_{1-x}$ nanoalloys allowing control the nanocrystallinity and so the order/disorder phase transition of the bimetallic CoPt nanocrystals. The enhanced crystallinity allows decrease the transition temperature and so to minimize the coalescence effect usually reported during the annealing process. This is demonstrated comparing the evolution of same nanoalloys obtained by wet chemistry either at high or at low temperature.

B. Experimental Method

B-1 Chemicals. Cobalt (II) chloride hexahydrate ($\text{CoCl}_2 \cdot 6\text{H}_2\text{O}$) was from Prolabo. Platinum (II) acetylacetonate ($\text{Pt}(\text{acac})_2$, 98%), Oleylamine (80-90%) and sodium borohydride (NaBH_4 , 99%) were from Acros. Sodium oleate (97%) and 1,2 hexadecanediol (98%) were from TCI. Oleic acid (90%), 1-octadecene (90%) Platinum (IV) chloride (PtCl_4 , 99,9%), Octylamine (99%) and tetrakis(decyl)ammonium bromide (TDAB, 99%) were from Aldrich. Cobalt (II) chloride hexahydrate ($\text{CoCl}_2 \cdot 6\text{H}_2\text{O}$) was from VWR.

B-2 Apparatus. A JEOL (100kV) model JEM-1011 transmission electron microscope (TEM) was used to determine the size of CoPt nanocrystals. The average compositions of the nanocrystals are determined by energy dispersive spectrometry (EDS) analysis using a scanning electron microscope (SEM, JEOL 5510 LV) with IXRF Systems 500 digital processing. Magnetic measurements have been obtained by using a VSM (vibrating sample magnetometer) operating in a Quantum Design PPMS.

B-3 In situ transmission electron microscopy study

Variable temperature in situ experiments were performed in a JEOL 2100F field emission transmission electron microscope operated at 200 kV, equipped with a Gatan 652 high-temperature sample holder. The CoPt nanoparticles synthesized by two different methods were cast dropped on 3-5 layer thick graphene substrates supported on standard nickel grids. The so-prepared TEM samples were loaded in the holder and annealed in vacuum in the temperature range between room temperature and 700°C at pressure below 2×10^{-7} Torr. The annealing experiments were carried out step-wise: the temperature between steps was increased at the rate of $20^\circ\text{C}/\text{min}$. Once the temperature of interest was reached the nanoparticles were stabilized for ~ 30 minutes without exposure to the electron beam. Electron irradiation during imaging was kept intentionally low below $2 \text{ A}/\text{cm}^2$ to prevent any uncontrolled electron beam induced structural changes.

B-4 Synthesis of of CoPt nanoalloys by the polyol process.

There is numerous synthesis of CoPt by the polyol process using mainly acetylacetonate derivative for both cobalt and platinum [17]. However, they are quite complex, as the procedure needs different step and always injection of the precursor at high temperature. Here in an attempt to control the crystallinity, a novel easy synthesis of CoPt nanoalloys by the so-called polyol process is achieved by mixing platinum(II) acetylacetonate ($\text{Pt}(\text{acac})_2$) and homemade cobalt(II)oleate ($\text{Co}(\text{oleate})_2$). The choice of oleate derivate is driven by the fact that above 100°C , $\text{Pt}(\text{acac})_2$ reacts with oleic acid to form stable complexes such as $\text{Pt}(\text{oleate})_2$ [18]. It has been mentioned, in the chemical synthesis of nanoalloys, the importance to use precursor having similar chemical structure in order to control the composition [19]. This is the reason why we use $\text{Co}(\text{oleate})_2$ as precursor. It should be noticed that $\text{Pt}(\text{oleate})_2$ is not available commercially and very difficult to synthesize. Conversely $\text{Co}(\text{oleate})_2$ could be easily synthesized. The synthesis of the cobalt-oleate complex is adapted from the one made for $\text{Fe}(\text{oleate})_3$ [20]. It is prepared by mixing cobalt (II) chloride hexahydrate (30 mmol in 30 ml of ultrapure water) and sodium oleate (60 mmol in a mixture of 40 ml of ethanol and 70 ml of hexane). The mixture is refluxed at 66°C for 4h and the organic phase is washed three times with ultrapure water in a separatory funnel. The remaining hexane in the organic phase is evaporated under vacuum in a rotavapor at 32°C . The paste of cobalt-oleate complex is then dissolved in 130 ml of hexane and again evaporated at 32°C . The purity and the atomic composition of the compound have been checked by quantitative Energy dispersive Spectrometry, EDS.

In a typical synthesis procedure of CoPt nanoalloys, platinum acetylacetonate (1.25×10^{-1} mmol), cobalt oleate (2.5×10^{-1} mmol) and 1, 2 hexadecanediol (5.6×10^{-1} mmol) are dissolved in a mixture of 10 ml of 1-octadecene, oleic acid (1.88×10^{-1} mmol) and oleylamine (1.88×10^{-1} mmol) in a three-necked round bottom flask, with magnetic stirring under

nitrogen at room temperature for 1h. In the reported polyol synthesis of CoPt, dioctyl-ether is often used as solvent but the obtained nanoparticles present crystalline defects [21]. In order to control the nanocrystallinity, we use another solvent with a higher boiling point: 1-octadecene (b.p. 317 °C). Hence the mixture is heated to the boiling point of the solvent and is refluxed for 30 min and then cooled to room temperature, giving a black dispersion, indicating the formation of the nanoparticles. The NCs are isolated by centrifuging and washed with a large excess mixture containing hexane (4 %) ethanol (43 %) and acetone (43 %). The NCs, capped by oleic acid, can be dispersed in organic solvents such as chloroform, hexane or toluene.

B-5 Synthesis of Solutions of CoPt nanoalloys by liquid-liquid phase transfer method.

Cobalt-platinum nanocrystals are made by a slightly modified synthesis as reported by Demortière *et al.* [19]. A 168.45 mg sample of PtCl_4 is dissolved in 5 mL of water and 10 mL of HCl to obtain PtCl_6^{2-} . A 80 mL volume of toluene containing 2 g of TDAB was mixed with this platinum complex solution for 24h. After the phase transfer, the aqueous phase is separated and discarded and we obtain the $\text{PtCl}_6(\text{TDA})_2$ precursor. As for the platinum salt, 118.9 mg of CoCl_2 was used to produce the $\text{CoCl}_4(\text{TDA})_2$ precursor. The following step of synthesis is made under nitrogen in a glove box. 10 mL of platinum organic phase and 10 mL of cobalt organic phase were added to 20 mL of toluene. After addition of octylamine (1 mL), an aqueous solution of NaBH_4 (378 mg, 10 mL of H_2O) is subsequently introduced into the mixture with rapid stirring. The organic phase containing the CoPt nanocrystals was separated after 21 h of stirring. The remaining toluene in the organic phase is evaporated (until 1 mL) under vacuum in a rotavapor at 35°C. The NCs are isolated by centrifuging and washed with a large excess ethanol and redispersed in 4 mL of toluene with 20 μL of octylamine. After one night, a final centrifugation is performed to eliminate unstable nanocrystals or aggregates in the solution.

C- Results and Discussion

C-1 Characterization of Co_xPt_{1-x} Nanoalloys.

Single phase methods has been developed in the last decade to synthesize metallic and bimetallic nanocrystals (ref). The "polyol process" is one of these methods. In this process, the diol or polyalcohol (as ethylene glycol for example) is the solvent, where the metal salts is dissolved, but it acts also as a reducing agent. This reaction is performed at high temperature (typically 100-200 °C). FePt [22] or NiPd [24] nanoalloys have been synthesized by the polyol process. As example, the use of iron acetylacetonate ($Fe(acac)_3$) and platinum acetylacetonate ($Pt(acac)_2$) in ethylene glycol or tetraethylene glycol, generates FePt NCs that show partially ordered fct structures [24]. Oleic acid or oleic amine are often used as capping agent and added directly in the chemical bath to limit the growth process. Furthermore, this high temperature process often allows to reach a better crystal quality and to avoid boron contamination often observed in monometallic nano-crystals obtained by single phase borohydride reduction [25].

However, only few report on the synthesis of CoPt nanoalloys by the polyol route, due to the difficulty to synthesize stable cobalt precursor derivate from acetylacetonate. Thus, the synthesis described here presents an original way using cobalt oleate and platinum acetylacetonate as precursor. Figure 2 shows Co_xPt_{1-x} nanoalloys obtained by the polyol process with two different compositions as revealed by the EDS analysis (figure 2G and 2H). They are easily obtained by tuning the initial salt composition: $Co_{32}Pt_{68}$ nanoalloys, in average composition are obtained by mixing 1.04×10^{-1} mmol cobalt oleate to 1.25×10^{-1} mmol platinum acetylacetonate (i.e., molar ratio 1.04-1.25). (figure 2 A, C, E, G), while $Co_{52}Pt_{48}$ is obtained when the molar ratio is 2.5-1.25 (Figure 2 B, D, F, H). In both case, TEM images show well-dispersed spherical NCs (figure 2A and 2B). It can be noticed that the NCs are homogeneous in size, shape and electronic contrast indicating the absence of segregation or

core-shell structure [16]. A slight decrease in the average size keeping the same polydispersity is observed with the composition of the CoPt (2.9 nm with a polydispersity of 13 % for $\text{Co}_{32}\text{Pt}_{68}$ and 2.5 nm with a polydispersity of 13 % for $\text{Co}_{52}\text{Pt}_{48}$). Furthermore, HRTEM observations reveal well faceted and crystallized nanoparticles as evidenced by clearly resolved lattice fringes (Figure 2C and 2D). The selected area electron diffraction patterns (Figure 2E and 2F) are similar to platinum one [26], which indicate the formation of the chemically disordered (A_1 , disordered fcc structure (Figure 1A)) face-centered-cubic (*fcc*) CoPt structure, whatever the composition is. Diffraction patterns appear more diffuse as the cobalt composition increases, which is consistent with an increase of the chemical disorder. This is characteristic of an alloy by substitution where platinum and cobalt atoms are randomly distributed on the platinum lattice: the more cobalt content, the less structural order [19].

The second method used to form nanoalloys involves transfer of the metal ion from a polar phase to a non-polar phase using a transferring agent. It has been developed for synthesis of metallic nanoparticles by Brust et al. in the 90's [27]. This Liquid-Liquid Phase Transfer (LLPT) method, also called two-phase synthesis, has been largely used to synthesize metallic nanoparticles as silver, gold, platinum or palladium. It typically involves the transfer of the metal precursor (metallic ions) from an aqueous solution to an organic solution containing a capping molecule as alkanethiol or amine. The transfer is assisted by a phase transfer agent such as tetrakis(decyl)ammonium bromide (TDAB). Reduction of metallic precursor is then carried by adding an aqueous solution of reducing agent (mainly NaBH_4) under vigorous stirring. We have reported previously the synthesis of $\text{Co}_x\text{Pt}_{1-x}$ nanoalloys by using the LLPT method [19]. In order to compare with the previous one, nanoalloys with an average composition of the $\text{Co}_{34}\text{Pt}_{66}$ and an average size of 2.2 nm have been synthesized (figure 2). TEM and HRTEM images confirm the formation of nanoalloys and the electron diffraction

patterns are indexed to the face-centred cubic (fcc) phase (A_1 phase, Figure 1A).

C-2 Magnetic properties of the native $Co_{30}Pt_{70}$ nanoalloys

In the following, we chose to focus our study on the particles with the composition around $Co_{30}Pt_{70}$ obtained by the two process of synthesis. For this composition a crystalline phase $L1_2$ corresponding to the fcc-ordered $CoPt_3$ is expected (see Figure 1B) [28]. The ordered $L1_2$ phase of the Co_xPt_{1-x} system present also a high coercivity and magnetocrystalline anisotropy. As for the $L1_0$ characteristic of $CoPt$, it can be obtained by annealing process on the native nanocrystals in the disordered phase. Magnetic measurements have been performed by using a VSM apparatus on the two previous samples of $CoPt$ NCs in the A_1 crystalline phase obtained either by polyol process ($CoPt_{polyol}$) or by the liquid-liquid phase transfer ($CoPt_{LLPT}$) with a composition close to $Co_{30}Pt_{70}$. The susceptibility versus temperature behavior is measured by a Zero Field Cooled (ZFC)/ Field Cooled (FC) experiment. In the ZFC measurement, the sample is cooled down to 3 K without an applied field starting from a high temperature where all the particles are in the superparamagnetic state. Afterwards, the magnetization is measured as function of the increasing temperature in an applied field of 100 Oe. In the case of an ideal system of perfectly monodisperse particles, the magnetization measured in the ZFC curve drops upon cooling from a maximum, usually defined as being the blocking temperature, T_B , to zero in a few degrees. As shown in Figure 4A, the two ZFC curves exhibit a maximum at 15 K (Table 1). This confirms the superparamagnetic behaviour at room temperature of both samples as expected for 2-3 nm size $CoPt$ NCs in the A_1 phase and the ferromagnetic behavior at 3K as seen on the hysteresis loop (Figure 4B). Indeed, from the T_B values and the average size, the particle magnetic anisotropy energy KV can be estimated by the relation $K_{eff}V = k_b T_B \ln(\tau/\tau_0) \approx 28 k_b T_B$, where V is the particle volume, K_{eff} is the effective magnetic anisotropy energy per volume unit and $\tau_0 \approx 10^{-9}$ - 10^{-11} s [29].

Table 1: Average diameter, polydispersity, blocking temperature (T_B), coercive field (H_C) and magnetic remanence normalized ($M_{R-Normalized}$) of $(Co_{30}Pt_{70})_{Polyol}$ and $(Co_{30}Pt_{70})_{LLPT}$.

	Average diameter	Polydispersity	T_B	H_C	$M_{R-Normalized}$
$(Co_{30}Pt_{70})_{Polyol}$	2.9 nm	11 %	15 K	4000 Oe	0.45
$(Co_{30}Pt_{70})_{LLPT}$	2.2 nm	13 %	15 K	5000 Oe	0.30

These values take into account the volume anisotropy but also the shape and surface anisotropies. Thus the magnetic anisotropy is estimated to $K_{eff} = 5.7 \cdot 10^4 \text{ J.m}^{-3}$ for 2.9 nm CoPt obtained by the polyol route and $13 \cdot 10^4 \text{ J. m}^{-3}$ for the 2.2 nm CoPt obtained by the LLPT method. These values are very low compare to that of bulk $CoPt_3$ in the $L1_2$ phase ($K_a = 2 \times 10^6 \text{ J.m}^{-3}$) however, there is a significant difference between the two values obtained for the two sample both in the A_1 phase. It is mainly due to the surface effect contribution. This behavior is extremely important around 2 nm, where 70% of atoms are located at the surface. In fact the magnetic energy is due both to K_s , surface anisotropy and to K_v the volumic anisotropy. As the size decrease the surface contribution increase. The high influence of surface atoms, which have coordination weaker than the atoms of the core, induces an important modification of the magnetic anisotropy energy [30]. It should be noticed that the shape of the ZFC curve depends on the nature of the CoPt nanoalloys. For those synthesized by the LLPT method, $CoPt_{LLPT}$, the width is significantly larger compare to those obtained by the polyol process, $CoPt_{Polyol}$. This reflects a larger distribution of the magnetic anisotropy energy (MAE). As the size distributions are similar (table 1), this effect arises from another source. Let us consider the crystalline structure of the native $Co_{30}Pt_{70}$ nanoalloys depending on the synthetic route. Table 2 summarizes the distribution of the crystalline structure for both samples as deduced from HRTEM studies (see figure 2 and 3). It is clear that the crystalline distribution is larger in the case of $CoPt_{LLPT}$ than for $CoPt_{Polyol}$.

Table 2: Distribution of crystallinity of $\text{Co}_{30}\text{Pt}_{70}$ nanoalloys depending on the synthesis route

	Monocrystals (%)	Polycrystals (%)	Undetermined (%)
($\text{Co}_{30}\text{Pt}_{70}$)_{Polyol} (236 particles)	33	31	36
($\text{Co}_{30}\text{Pt}_{70}$)_{LLPT} (525 particles)	21	23	56

This cannot be explained by the size difference as the shape equilibrium calculation does not predict a drastic difference of stability between crystalline fcc structure and the non crystalline one as decahedron in this range of size. [31] However it should be remained that the polyol process occurs a higher temperature than the liquid-liquid phase transfer method, which is known to favor a better crystallinity. As a matter of fact, it should be noticed that the percentage of undetermined structure and polycrystals is very high for $\text{CoPt}_{\text{LLPT}}$, compare to that observed obtained by the polyol method (see table 2). These amorphous or quasi-amorphous materials have a different magnetic anisotropy compare to the crystalline one, which could explain the larger distribution of MAE observed figure 4A. This is confirmed by the hysteresis loop measured in the ferromagnetic regime at 3K (figure 4B and Table 1). The coercivity at 3K is higher for $\text{CoPt}_{\text{LLPT}}$ than for $\text{CoPt}_{\text{Polyol}}$, which is consistent with a higher magnetic anisotropy. Indeed, the coercivity increases with K_{eff} [32]. However, the reduced remanence, M_r/M_s , is lower for $\text{CoPt}_{\text{LLPT}}$. This is surprising if we consider only the anisotropy value, but could be expected if we take into account the high level of amorphous or quasi-amorphous materials, which are softer magnet compare to the crystalline structure. It should also be considered that for $\text{CoPt}_{\text{LLPT}}$, some NCs are always in the superparamagnetic state even at 3K due to the larger anisotropy distribution. This explains also the fact that magnetization at saturation is not reached even at 5T compare to the $\text{CoPt}_{\text{Polyol}}$.

C-3 Ordering of the $\text{Co}_{30}\text{Pt}_{70}$ nanocrystals.

Thus the fcc- CoPt NPs in the disordered A_1 phase exhibit a low magnetocrystalline anisotropy and soft magnetic properties as well. Therefore, high temperature annealing is needed (around 685 °C in the bulk phase) for the transformation from the fcc disordered A_1 to

the fcc ordered $L1_2$ phase. We have seen previously the influence of the nanocrystallinity on the magnetic properties. Here we will consider the effect of the nanocrystallinity on the ordering by comparing the order/disorder phase transition induced by annealing of CoPt nanoalloys. This has been investigated by an in situ annealing, in real time and on sub-nanometer scale, using HR-TEM studies on both samples $(Co_{30}Pt_{70})_{llpt}$ et $(Co_{30}Pt_{70})_{polyols}$ deposited on a graphene coated TEM grid. (see experimental section).

Table 3: Mean diameter and polydispersity of $(Co_{30}Pt_{70})_{polyol}$ nanoalloys at different temperature of the annealing process.

	25°C	300°C	420°C	550°C	680°C
Mean diameter	2.9 nm	2.7 nm	3.0 nm	3.3 nm	3.2 nm
Polydispersity	11 %	12 %	17 %	15 %	12 %

It is shown on figure 5 a typical structural evolution of the $(Co_{30}Pt_{70})_{polyols}$ NCs during the annealing process. It should be noticed that compar to the native NCs there is a slight and continuous increase of the average size with the annealing temperature (Table 3). This has been already observed during annealing process of CoPt and attributed to an Ostwald ripening to the benefit of larger nanoparticles [33]. HR-TEM patterns at 300°C show lattices fringes of the NCs, characteristics of the disordered A_1 phase with a lattice spacing equal to 2.14 Å (Figure 5A). This is confirmed by the power spectrum, where only one pair of reflection corresponding to the (111) plane is observed. The deduced lattice parameter is equal to 3.70 Å, which is consistent with a A_1 fcc structure. In fact no variation of the lattice parameters is observed compare to the value a obtained at room temperature (figure 2C). drastic structural evolution is observed begins as the temperature rises 420°C, (Figure 5B): some nanocrystals exhibit different fringe patterns characteristics from the ordered fcc $L1_2$ phase with a lattice parameters of 3.85 Å as deduced from power spectrum (figure 5-B3 and Table 4). Indeed the corresponding power spectrum clearly shows characteristics reflections corresponding to the (001) plane, which confirm the formation *at low temperature* of ordered $CoPt_3$ nanoalloys. It

should be noticed that no sintering effect are observed as the NCs still remain isolated on the graphene coated TEM grid. Further increase of the temperature induces the ordering of more and more nanocrystals (Figure 6 and Table 3) as well as a continuous increase of the lattice parameters. This isotropic expansion of the lattice parameter is characteristic of the ordering of CoPt_3 indicating a homogeneous order-disorder transformation [34]. The bulk value is equal to 3.9 Å [34], in our case it is reach at 680 °C, which indicates that the ordering is total.

Table 4: Structural evolution of ($\text{Co}_{30}\text{Pt}_{70}$) obtained by polyol process during the annealing. Only nanoparticles with visible atomic planes were taken in count on the percentage of disordered nanoparticles (A_1) and ordered nanoparticles ($L1_2$). The ratio ($L1_2/A1$) is given for each temperature. The lattice parameters are deduced from the power spectra shown on figure 5.

Temperature	300°C	420°C	550°C	680°C
Number of nanoparticles studied with observable atomic planes	161	161	191	77
Lattice parameters (Å)	3.70	3.85	3.86	3.90
% A_1	100	83	59	43
% $L1_2$	0	17	41	57
$L1_2/A_1$	0	0.20	0.69	1.33

Table 4 shows also the statistical evolution of the ordering of the NC. At 680°C, 57 % of the observed NCs present the $L1_2$ structure. In fact there is a linear evolution of this percentage with the annealing temperature as shown of figure 6 in relation with the evolution of the lattice parameters. This indicates a continuous and monotonic transition from the ordered to the disordered fcc structure of CoPt_3 . Indeed, this low temperature transition is surprising. Even if it has been reported, and calculated, a decrease of the transition temperature in the $\text{Co}_x\text{Pt}_{1-x}$ phase diagram in case of nanocrystals compare to bulk materials [31, 35], this value close to 180°C below the bulk value (around 760 °C [28]), for 3 nm in size nanocrystals. Here the huge decrease of 350°C of the order/disorder transition is very significant.

Figure 7 and Table 5 resume the same experiment but with $\text{Co}_{30}\text{Pt}_{70}$ made by the liquid-liquid

phase transfer method. behavior of the nanocrystals strongly differs from both structural and ordering point of view.

Table 5: Structural evolution of $(Co_{30}Pt_{70})$ obtained by LLPT process during the annealing. Only nanoparticles with visible atomic planes were taken in count on the percentage of disordered nanoparticles (A_1) and ordered nanoparticles ($L1_2$). The ratio ($L1_2/A1$) is given for each temperature. The lattice parameters are deduced from the power spectra shown on figure 6.

Temperature	300°C	400°C	550°C	680°C
Number of nanoparticles studied with observable atomic planes	51	101	99	30
Lattices parameters (Å)	3.49	3.60	3.84	3.63
% A_1	100	97	79	87
% $L1_2$	0	3	21	13
$L1_2/A1$	0	0.03	0.27	0.15

Similarly to the previous case, no ordering is observed 300 °C however it is clear that significant coalescence occurs, even if isolated nanocrystals are still present on the TEM grid. This effect probably arises from the nature of the passivating agent on the CoPt surfaces, depending on the chemical route. Indeed, alkyls amines are used in this soft synthetic method when CoPt polyols are coated by carboxylate derivatives (see experimental method). In case of $CoPt_{llpt}$, this is due to the growth process at room temperature, which is totally inhibited if chemically bond agents are used as a capping agent [36]. As a result, there protection is not strong enough to avoid a coalescence process. However, operating in situ on deposited NCs on the TEM grid limit the progression of this process during the annealing and allows us to study the ordering of $(Co_{30}Pt_{70})_{llpt}$. From a crystalline point of view, no transition is observed at 300 °C whatever are the size of the NCs. The lattice parameter is still corresponding to the (111) plane of the A_1 disordered fcc structure (Table 5). At 400 °C only a few part of NCs present a structural evolution towards the $L1_2$ fcc ordered phase: 3% of the observed NCs when it was 17% for the $(Co_{30}Pt_{70})_{polyols}$. It is not only a size effect as the corresponding

transition is not observed for the smallest NCs. Further increase of the annealing temperature yields to an increase of the number of NCs in the $L1_2$ phase, but their proportion is always very low compare to the $(Co_{30}Pt_{70})_{polyols}$. (Figure 6 and Table 5). At 680 °C only 15% of the observed NCs present a $L1_2$ structure.

This strong difference of chemical ordering depending on the chemical route can be understood if we take into account the difference of crystallinity reported above. It has been reported in similar annealing process of CoPt nanoalloys that the A_1 to $L1_0$ (or $L1_2$) is a multi-step process [33]. There is first a transition from polycrystalline, or non-crystalline phase as decahedron, to a fcc truncated octaedron structure. Simultaneously, the NCs size increases and their shape becomes more and more isotropic. Further increase of the temperature induces the chemical ordering of the NCs keeping a fcc octaedron structure. Indeed it has been reported that the noncrystallinity of the nanocrystals is a limiting factor to achieve the chemical order. Furthermore agglomeration of the NCs during the annealing process often yield to high angle grain boundaries, which also prevent or slow down the chemical ordering and the homogeneous crystalline transition as numerous energetic barrier should be overcome to be the correct crystalline structure [15]. Thus chemical ordering can only be reached at higher temperature. In our case, on the one hand $(CoPt)_{llpt}$ present an important part of non crystalline or amorphous structure (Table 2), on the other hand the passivating agent is not strong enough to prevent the coalescence conversely to the case of $(CoPt)_{polyol}$. As a result the chemical ordering appears at lower temperature and on larger scale for CoPt nanoalloys made by the polyol process compare to similar NCs synthesized by the LLPT process. This is probably reinforced by the fact that the polyol process is a high temperature synthetic method, which favors a better crystallinity of the NCs. Thus the first step of the order transition considered above is limited or inexistent. As a matter of fact, this assumption is confirmed if we consider the chemical ordering of $Co_{50}Pt_{50}$ synthesized by the polyol process (Figure 8).

Again $\text{Co}_{50}\text{Pt}_{50}$ NCs present mainly a disordered fcc A_1 crystalline structure, with a truncated octahedron shape, and only few non-crystalline structure are observed. As a consequence, similar behavior as reported above for $\text{Co}_{30}\text{Pt}_{70}$, is observed during the in situ annealing process. For a 50-50 composition the ordered structure, as deduced from the phase diagram, is the tetragonal fct $L1_0$ phase (see Figure 1). It can be obtained by annealing as the chemical synthesis only yield to the disordered A_1 structure. Similar experiments have been undertaken on these NCs. There is a slight coarsening due to the Ostwald ripening of the NCs as their size increase from 2.5 to 3.0 nm but without coalescence. Furthermore, ordering occurs also at low temperature (420 °C) and the nanoparticles keep their ordered $L1_0$ structure at low temperature as observed on figure 8E and 8F. Our previous study on NCs synthesized by the LLPT method with same size but passivated also by alkyl amin chains shown a huge coalescence and a chemical ordering only above 500 °C [16]. Again, we observe the beginning of chemical ordering at lower temperature for $(\text{CoPt})_{\text{Polyol}}$ than for $(\text{CoPt})_{\text{LLPT}}$ whatever the composition.

D- Conclusion

We have reported here a new synthesis of bi-metallic $\text{Co}_x\text{Pt}_{1-x}$ nanocrystals by the polyol process, allowing a large control of the composition of these nanoalloys. Structural analysis performed by TEM and HR-TEM show that, compare to conventional soft chemical approach, the nanocrystallinity of these nanoalloys strongly differs as fcc structure is mainly observed even for very low size. The decreases of the crystalline polydispersity induce a decrease of the distribution of magnetic anisotropy energy as observed by magnetic measurement. Furthermore, the order/disorder transition induced by annealing and observed in situ by HR-TEM studies, clearly demonstrate a decrease of the chemical ordering temperature compare to the bulk value but also to nanocrystals obtained by soft chemistry.

The absence of coalescence during the annealing allows us to develop a new approach to obtain periodic 2D self-assembly of chemically ordered nanoalloys. Hence it is clearly demonstrated the importance of the chemical route on the crystallinity of the nanocrystals. This strongly influences the physical and chemical properties of the nanocrystals.

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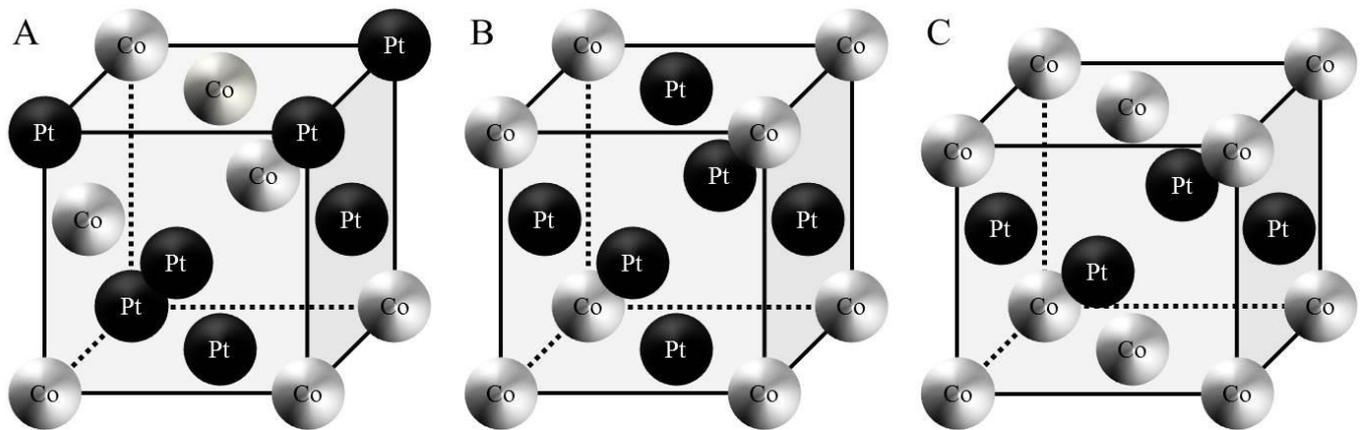


Figure 1 : Representation of $\text{Co}_x\text{Pt}_{100-x}$ structure on three different phases: (A) disordered phase A1 with cubic symmetry (B) ordered phase $L1_2$ with cubic symmetry ($X=25$) and (C) ordered phase $L1_0$ with face centered tetragonal symmetry ($X=50$).

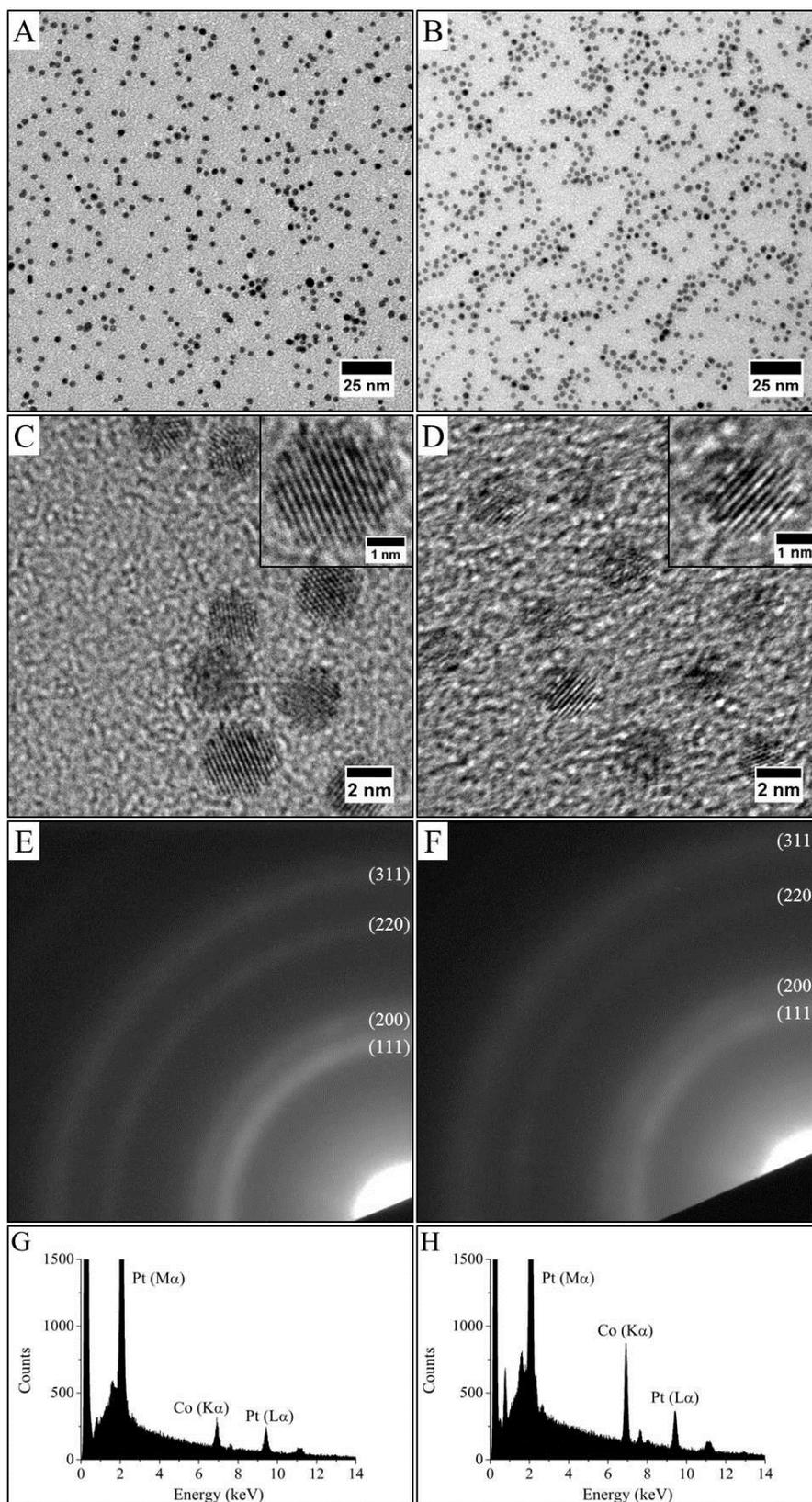


Figure 2: TEM images (A-B), HRTEM images (C-D), electronic diffraction (E-F) and EDX analysis (G-H) of $(Co_{30}Pt_{70})_{Polyol}$ (left images) and $(Co_{50}Pt_{50})_{Polyol}$ (right images).

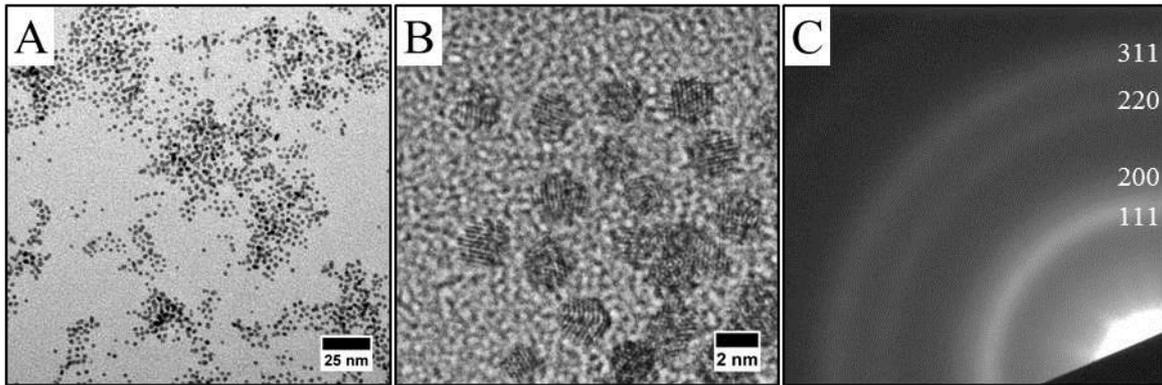


Figure 3: TEM images (A), HRTEM images (B) and electronic diffraction (C) of $(\text{Co}_{30}\text{Pt}_{70})_{\text{LLPT}}$

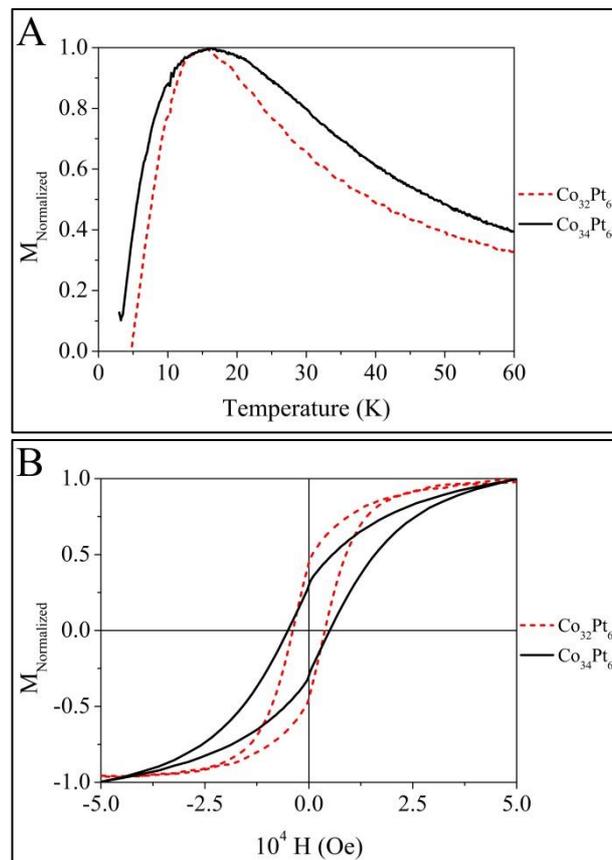


Figure 4: (A) Temperature dependence of the magnetization in the Zero-Field Cooled curves with applied field $H=100$ Oe and (B) hysteresis loop of $\text{Co}_{32}\text{Pt}_{68}$ (dash line) obtained with polyol process and $\text{Co}_{34}\text{Pt}_{66}$ obtained with LLPT synthesis (solid line).

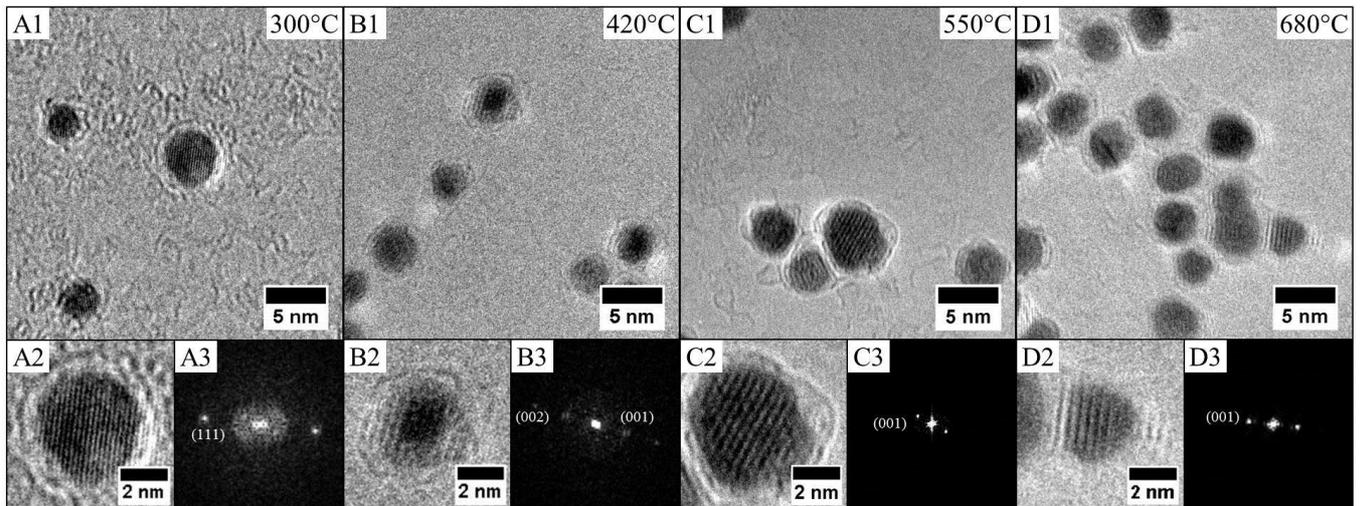


Figure 5: HRTEM images of $(\text{Co}_{30}\text{Pt}_{70})_{\text{Polyol}}$ at various annealing temperatures: (A) 300°C; (B) 420°C; (C): 550°C and (D) 680°C. For each temperature, HRTEM images of one nanoparticle (A2, B2, C2, D2) and the associated FFT (A3, B3, C3, D3).

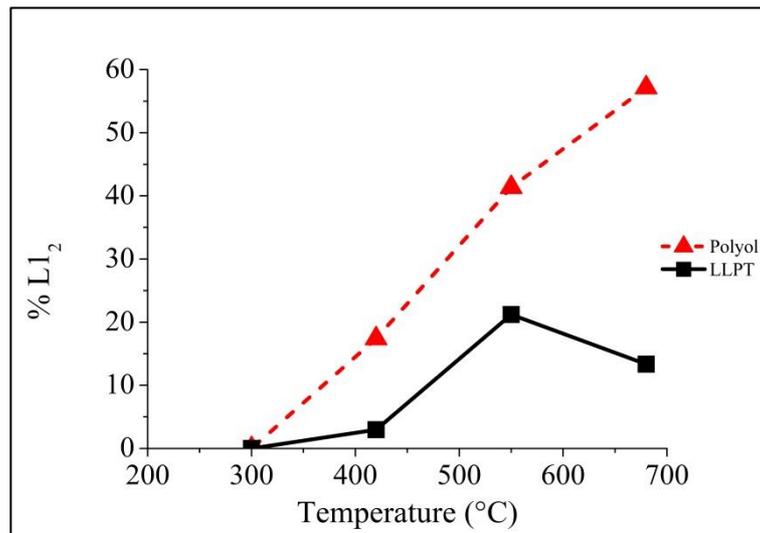


Figure 6: Evolution of ordering with temperature during the annealing process.

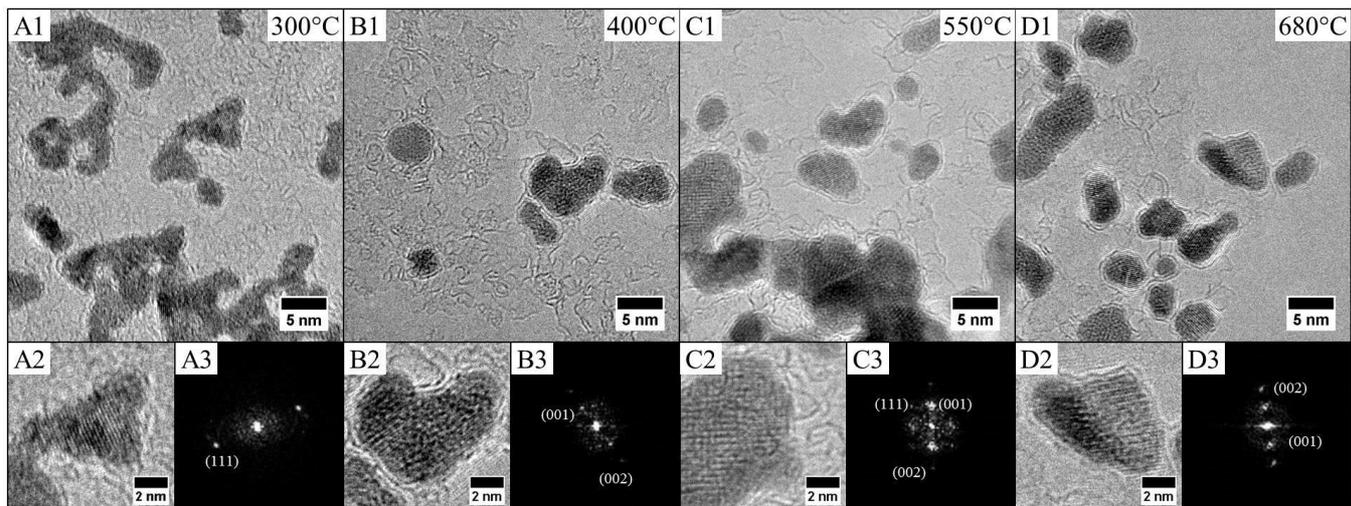


Figure 7: HRTEM images of $(\text{Co}_{30}\text{Pt}_{70})_{\text{LLPT}}$ at various annealing temperatures: (A) 300°C; (B) 400°C; (C): 550°C and (D) 680°C. For each temperature, HRTEM images of one nanoparticle (A2, B2, C2, D2) and the associated FFT (A3, B3, C3, D3).

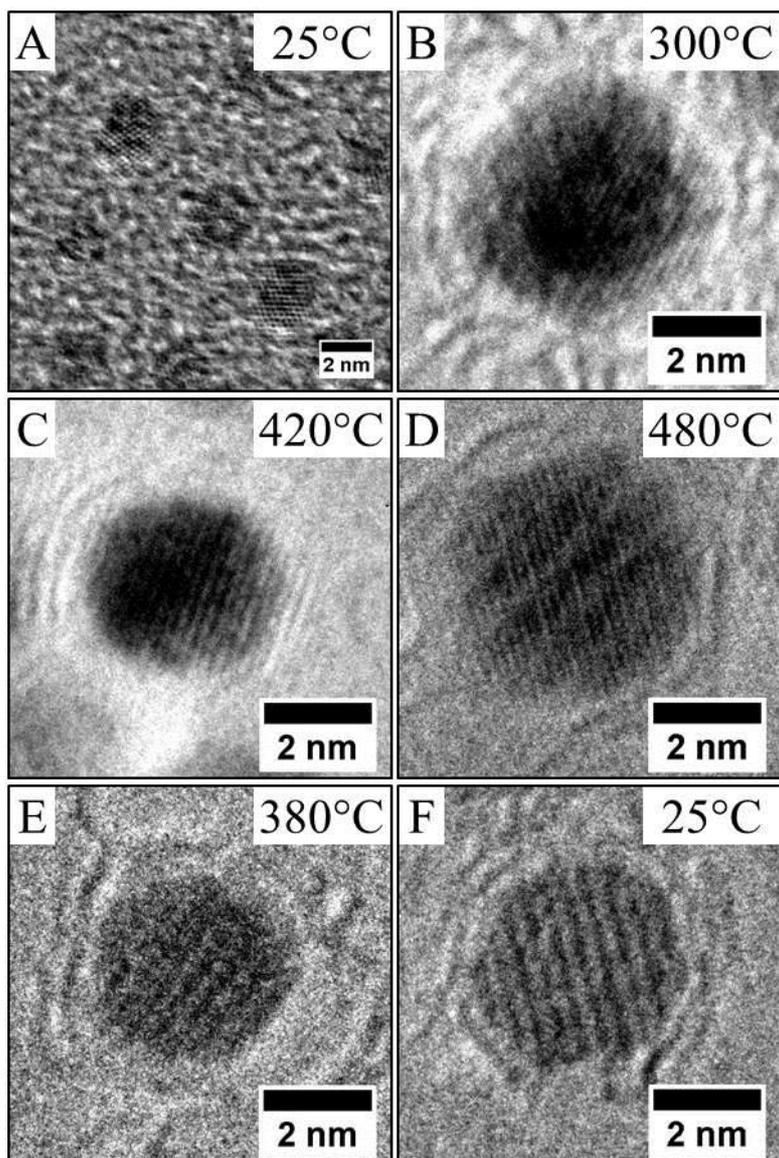


Figure 8: HRTEM images of $(Co_{50}Pt_{50})_{Polyol}$ during annealing treatment: (A) room temperature (RT), (B) 300°C, (C) 420°C on the increasing part of the annealing and (D) 480°C, (E) 380°C and (F) RT on the decreasing part of the annealing.