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Spin transition nanoparticles made electrochemically†

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Materials displaying novel magnetic ground states signify the most exciting prospects for nanoscopic devices for nanoelectronics and spintronics. Spin transition materials, *e.g.*, spin liquids and spin glasses, are at the forefront of this pursuit; but the few synthesis routes available do not produce them at the nanoscale. Thus, it remains an open question if and how their spin transition nature persists at such small dimensions. Here we demonstrate a new route to synthesize nanoparticles of spin transition materials, gas-diffusion electrocrystallization (GDEx), wherein the reactive precipitation of soluble metal ions with the products of the oxygen reduction reaction (ORR), *i.e.*, *in situ* produced H_2O_2 , OH^- , drives their formation at the electrochemical interface. Using mixtures of Cu^{2+} and Zn^{2+} as the metal precursors, we form spin transition materials of the herbertsmithite family—heralded as the first experimental material known to exhibit the properties of a quantum spin liquid (QSL). Single-crystal nanoparticles of $\sim 10\text{--}16\text{ nm}$ were produced by GDEx, with variable Cu/Zn stoichiometry at the interlayer sites of $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$. For $x = 1$ (herbertsmithite) the GDEx nanoparticles demonstrated a quasi-QSL behavior, whereas for $x = 0.3$ ($0.3 < x < 1$ for paratacamite) and $x = 0$ (clinoatacamite) a spin-glass behavior was evidenced. Finally, our discovery not only confirms redox reactions as the driving force to produce spin transition nanoparticles, but also proves a simple way to switch between these magnetic ground states within an electrochemical system, paving the way to further explore its reversibility and overarching implications.

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Introduction

The synthesis of materials exhibiting spintronic properties is at the forefront of pursuit of novel magnetic ground states (*i.e.*, quantum magnets). Spin transition materials refer to frustrated antiferromagnets, in which their spins are not aligned in a regular pattern (geometrical spin frustration), including spin glasses, quantum spin liquids (QSL), and high-temperature superconductivity materials.^{1,2} For instance, quantum spin liquids (QSL) were first realized by herbertsmithite,³ with chemical formula $\text{ZnCu}_3(\text{OH})_6\text{Cl}_2$. Due to its characteristics as a spin-liquid, herbertsmithite may have applications in

quantum computing devices (*e.g.*, for storage and memory purposes), and in quantum communications.^{1,4}

The copper ions in herbertsmithite are arranged in triangular grids known as kagomé lattices, wherein the spins are uncoupled.⁵ Each spin tends to be anti-parallel to its neighbors, but the geometric arrangement of the sites where the spins sit makes an ordered alignment impossible. So, the spins collapse into a fluctuating state.^{3,5,6} This non-trivial and disordered arrangement, results in magnetically-frustrated materials due to the low spin of the copper ions ($S = 1/2$). The entangled interactions continue to fluctuate, even at the lowest temperature of absolute zero.^{7,8} In such condition, the spins in other magnetic states would be already frozen, but in the quantum state, spins are never locked into a particular configuration.⁶ Compounds featuring such lattices are predicted to show no long-range magnetic order due to competing for antiferromagnetic interactions between nearest-neighbor spin centers,^{2,4-7,9-11} thus being also candidates for quantum spin transitions.

Before these materials can be effectively integrated into spintronic technologies, the competitive synthesis of herbertsmithite remains a significant challenge. A major synthesis problem is the production of sizeable amounts at relevant rates, with controllable physicochemical features. Moreover,

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envisioning small-scale applications, its synthesis at the nano-scale has not been reported. Thus, it is uncertain if its spin-liquid properties are preserved at such scale. This is also the case for other spin-liquid materials, as well as other spin-transition materials, such as spin-glasses.

The first method reported for synthesizing crystalline $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$ (wherein $x = 1$ for herbertsmithite, and $x = 0$ and $0.3 < x < 1$ for its polymorphs, clinoatacamite and paratacamite, respectively) was issued in 2012 (herbertsmithite was originally found in nature only in 2004).⁶ Herbertsmithite and its polymorphs are typically synthesized by hydrothermal, or solvothermal methods (e.g., 458 K–473 K). Related cuprate compounds of the form $\text{MCu}_3(\text{OH})_6\text{Cl}_2$ with divalent cations (i.e., $\text{M} = \text{Mg}^{2+}$, Co^{2+} , Fe^{2+} , Mn^{2+} , and Ni^{2+})² and trivalent cations, $\text{MCu}_3(\text{OH})_6\text{Cl}_3$ ($\text{M} = \text{Y}^{3+}$ or the lanthanides Nd^{3+} and Sm^{3+}),^{12,13} have been achieved by unconventional solid-state reactions at higher temperature (e.g., 463 K). An ionothermal method, initially designed for the fabrication of new zeolitic solids, has also been described for the preparation of materials with kagomé lattices.¹⁴ However, as in the case of the methods previously described, the synthesis rates are rather slow (weeks- to months-scale). As an example, it took ten months to grow a 1 mm monocrystal of herbertsmithite, in a very complicated hydrothermal reactor.^{9,15}

Although there is one work presenting the formation of nanoscale clinoatacamite by hydrothermal synthesis at 363 K–368 K, there are no equivalent methods available for producing nanoscale herbertsmithite. Furthermore, the study where nanoscale-clinoatacamite is reported does not evidence its magnetic (spin transition) properties, nor the control of other material features, such as crystallite size and stoichiometry, besides requiring the addition of a toxic organic buffer i.e. 2-(*N*-morpholino) ethanesulfonic acid.¹⁶

In general, all these methods are limited to meager production rates and have restrained the production of herbertsmithite (and its polymorphs) to limited samples of macroscale materials (i.e., mm-scale). Green synthesis routes are also preferred, but they are also not available thus far. The development of such routes will be a key step forward for innovations, growth, and competitiveness in the digital miniaturization arena—e.g., using quantum technologies in computing and electronics.

It has been experimentally proven for many materials that micro- and nano-scale dimensioned particles can differ in properties vs. their macro-scale counterparts. Especially, spin transition materials may display changes in their magnetic properties with size, and the effects at the nanoscale are mainly unknown.

In this work, we present the synthesis of spin transition nanoparticles. Specifically, using copper and zinc chlorides as precursors, our first goal was to achieve the synthesis of $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$ compounds.

By using different concentrations and ratios of Cu and Zn, different values of x are targeted. The electrochemical method developed, gas-diffusion electrocrystallization (GDEx), results in these materials by the reactive precipitation of intermedi-

aries issued from the oxygen reduction reaction (ORR) at a gas-diffusion cathode, with the metal precursors provided. GDEx has been formerly described for the synthesis of iron oxide nanoparticles with a controlled saturation magnetization.¹⁷ The materials resulting from the present research were investigated by X-ray powder diffraction (XRD), together with quantitative phase analysis (QPA, Rietveld method), infrared spectroscopy (IR) and inductively coupled plasma optical emission spectrometry (ICP-OES). The morphology and size of the nanoparticles were measured using transmission electron microscopy (TEM).

A second goal was to achieve well-controlled magnetic properties linked to the nano-sized $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$ particles. Zero-field-cooled (ZFC) and field-cooled (FC) thermal dependence of magnetization (M) were used to distinguish magnetic phase transitions in these materials down to 2 K. Also, hysteresis loops at 2 K and 8 kA m^{−1} (100 Oe) were investigated for the different compositions.

We demonstrate that it is possible to use our novel electrochemical method for the production of spin transition nanoparticles (i.e., herbertsmithite, clinoatacamite, and paratacamite), in a fast, upscalable, atom- and electron-efficient, energy efficient, and environmentally-friendly way, setting a new precedent towards the industrial era of these materials.

Experimental

Gas-diffusion electrocrystallization reactor

The electrochemical reactor design described in Prato *et al.*,¹⁷ was used in this study. As a working electrode, 10 cm² of VITO CoRE® multilayered carbon-based gas-diffusion cathodes were used to enable the GDEx process. The VITO CoRE® electrodes consisted of a current collector (stainless steel gauze), an active layer (i.e., porous electrically-conducting matrix), and a hydrophobic gas diffusion outer layer (porous PTFE). The composition of the active layer was 20% PTFE and 80% active carbon. Norit®SX1G (878 m² g^{−1}, Norit Americas Inc., USA) was employed as the active carbon source. The counter electrode (anode) consisted of a 10 cm² of platinum sheet laser-welded to titanium (Ti) plate current collector. The anode and cathode compartments were separated by an ion-permeable separator (Zirfon® Perl UTP 500, Agfa, Belgium).¹⁸ A 3 M KCl saturated Ag/AgCl reference electrode (+212 mV vs. SHE) (REF 321, Radiometer Analytical, Hach, USA) was inserted in the proximity of the working electrode, *via* an external connector chamber, filled with 3 M KCl. A long cotton thread (e.g., 30 cm, packed within the external connector chamber) verged the reference and the working electrodes, through a small channel, with the purpose of establishing a microchannel enabling a continuous capillary-suction of the electrolyte. The thread was not treated, since due to its manufacturing process the wax layer that naturally covers cotton fibres is removed, making it more polar and hydrophilic due to exposure of the fibrous structure of cellulose, rich in hydroxyl (OH[−]) groups.¹⁹



All potentials here reported are referred *versus* the standard hydrogen electrode (SHE).

Operation of the gas-diffusion electrocrystallization reactor

Experiments were conducted on a multichannel potentiostat (VMP-3, Bio-Logic SAS, France). The GDEx process was controlled chronoamperometrically at $-0.15\text{ V}_{\text{SHE}}$ and room temperature ($18\text{ }^{\circ}\text{C}$) of synthesis. The pH in the catholyte was monitored using a sensor for continuous measurement of pH in liquid media (SE555X/1-NMSN pH sensor, Knick, Germany). A pH data logger was connected to the Bio-Logic potentiostat to simultaneously follow up the pH *vs.* charge (*Q*) evolution.

The data logger consisted of a pH transmitter (A1491N-P1-10-0000, Knick, Germany) with a converter (P15000 H1, Knick, Germany) which transforms the 4–20 mA signal delivered by the transmitter into a 0–10 V signal.

All experiments were performed in batch mode with recirculation. The catholyte was recirculated at 15 L h^{-1} throughout the cathode compartment, where gas-diffusion electrocrystallization of Cu^{2+} and Zn^{2+} occurred. The total liquid volume of catholyte and anolyte was 500 mL, which was continuously stirred in the recirculation reservoir (*i.e.*, a borosilicate glass bottle) using a polygonal, PTFE-coated rotating magnetic stirring bar (*i.e.*, 200 rpm). The air was fed through the cathodic gas compartment at a flow rate of 200 ml min^{-1} . A mass gas flow meter and controller (GF40 Bronkhorst hi-tech B.V, Netherlands) was set in place. An overpressure of 26 mbar over a water column (see Fig. 1 for a schematic layout) was applied at the gas exhaust.

The GDEx-driven synthesis was carried out using a mixture of Cu^{2+} and Zn^{2+} ions as the metal precursors, and O_2 (in air) as the oxidant gas through the gas-diffusion cathode. An acidified supporting electrolyte was employed as an anolyte and catholyte, composed by a NaCl aqueous solution (0.5 M), adjusted at pH 3 by adding HCl (1 M). In order to tune the formation of stable $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$ products with the desired value for *x*, different $[\text{Cu}^{2+}]_0/[\text{Zn}^{2+}]_0$ ratios were evaluated. $\text{CuCl}_2 \cdot 2\text{H}_2\text{O}$ 99.999% (Sigma-Aldrich, Germany) and ZnCl_2 99.999% (Sigma-Aldrich, Germany) were supplemented, respectively.

Oxygen (in air), provided through the gas chamber of the electrochemical cell (right side chamber, Fig. 1A), was electrochemically reduced at the active porous carbon layer of the gas-diffusion cathode (Fig. 1B). The products of the electrochemical ORR, profusely available at the electrochemical interface, reacted with the metals ions in solution (*e.g.*, Cu^{2+} , Zn^{2+}), which are transported to the hydrophilic porous carbon on the cathode, *via* the aqueous electrolyte. When these metal ions met the ORR products (or highly reactive intermediaries) supersaturation was reached, resulting in the nucleation of hydroxides or oxides. Additive OH^- concentration and supersaturation kept ongoing; thus secondary nucleation and crystal growth proceeded during the transient period of residence of the primary nuclei formed within the cathodic interface. The resulting precipitate progressively striped out, as a result of the flow and polarization conditions.²⁰ The solid precipitate was collected in the electrolyte solution and left to sediment under stagnant conditions. Subsequently, the precipitate was washed with deionized water with a pH set at that of the final synthesis

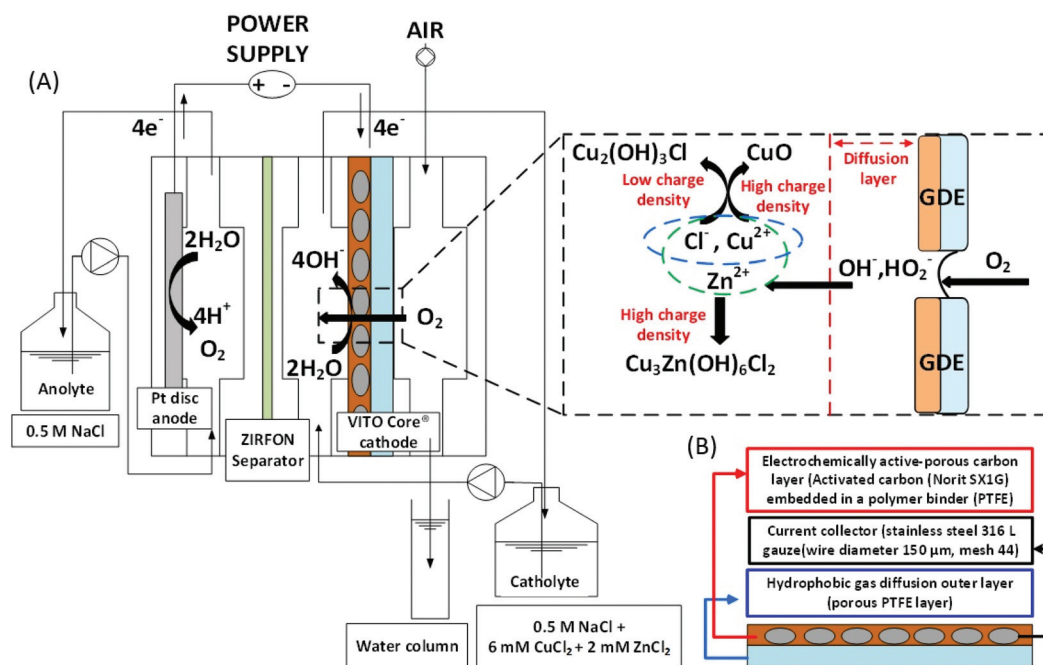


Fig. 1 (A) Gas-diffusion electrocrystallization (GDEx), represented for a batch setup. The potentiostat employed is indicated as a power supply, due to its function. (B) Schematic representation of the parts of the multilayered gas-diffusion electrode employed as cathode.



solution, re-centrifuged (Jouan CR422, France) at 3000 rpm, washed three times, and dried afterwards under a nitrogen atmosphere, resulting in a particulate powder.

Particle size, morphology and structure characterization

Electronic microscopy. Particle size and morphology were measured by transmission electron microscopy (TEM) using a JEOL JEM 2100FII operated at 200 keV. For the observation of the sample in the microscope, the particles were dispersed in butanol, and a drop of the suspension was placed onto a copper grid covered by a carbon film. The mean particle size and distribution were evaluated by counting more than 100 particles using Digital Micrograph™. After that, data were fitted to a normal distribution to obtain the mean particle size (d_{NP}) and standard deviation (σ).

X-ray powder diffraction (XRD). The XRD patterns were obtained with a diffractometer (Empyrean, Malvern Panalytical, United Kingdom) using CoK α radiation ($\lambda = 1.7928 \text{ \AA}$) with 40 mA–45 kV and a finer step size of 0.013° in the same scan range. Quantitative phase analysis (QPA) by the Rietveld refinement method with the HighScore Plus software (Empyrean, Malvern Panalytical, United Kingdom) was carried out for the quantitative analysis of the phase distribution (%), using the measured diffraction profile and a calculated profile crystal from the inorganic crystal structure database (ICSD).^{21,22}

Stoichiometric analysis. An amount of 50 mg of the precipitate powder sample was transferred into a digestion vessel. Then, 6 mL of HCl and 2 mL of HNO₃ were added. The screw cap of the digestion tube was gently turned and placed on a heating block. The temperature was slowly increased to $(105 \pm 5)^\circ\text{C}$ and kept for 120 min. The tube was cooled down to room temperature, and water was added to the volume mark. The copper and zinc content in the digested solution was determined by inductively coupled plasma optical emission spectrometry (ICP-OES, Agilent, 5100, USA) in an axial view using ICP-OES equipped with a baffled cyclonic spray chamber and a conical nebulizer. The atomic emission lines of 324.754 nm and 213.857 nm were used for copper and zinc, respectively. The metal content in the precipitate sample and the phase percentage calculated by QPA were used to determine the stoichiometric coefficient x on the interlayer site of $\text{Cu}_{4-x}\text{Zn}_x(\text{OH})_6\text{Cl}_2$.

Infrared spectroscopy (IR). IR measurements were performed on a Thermo Scientific, Nicolet iS5 (Waltham, MA, USA), with a diamond plate, and used to distinguish between clinoatacamite and related Zn-polymorphs. The spectra were recorded in the range of $4000\text{--}400 \text{ cm}^{-1}$ with a resolution of 2 cm^{-1} .

Magnetic characterization and analysis

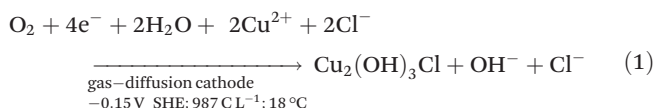
The magnetic characterization was carried out on powder samples in a glycerin capsule and performed in a SQUID magnetometer Quantum Design MPMS-5S, with 5 T maximum applied field and temperature range from 2 K to 400 K. The ZFC–FC curves were measured at 8 kA m^{-1} (100 Oe) from 2 K to 300 K, with a step of 0.25 K from 2 K to 6 K, 0.5 K from 6 K

to 12 K and more than 1 above 12 K. Before each ZFC measurement, a 3 T demagnetization field was applied in order to avoid any remanent magnetization in the superconductor coils. The hysteresis loops up to 5 T were measured at 2 K, 3 K, 4 K, 6 K, and 15 K.

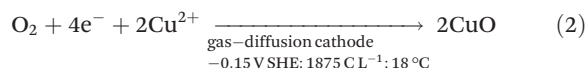
Results and discussion

Charge-driven stoichiometric and structural control

GDEx allows the selective preparation of $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$ materials. By varying the $[\text{Cu}^{2+}]_0/[\text{Zn}^{2+}]_0$ ratio in the GDEx catholyte (wherein $[\text{Cu}^{2+}]_0$ and $[\text{Zn}^{2+}]_0$ correspond to the initial concentration of copper and zinc ionic precursors, respectively), together with a systematic control of the electric charge, the stoichiometric composition of the $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$ products and their structural control, between $\text{Cu}_2(\text{OH})_3\text{Cl}$ and CuO , can be customized (Fig. 2). For instance, the formation of $\text{Cu}_2(\text{OH})_3\text{Cl}$ and CuO , for $x = 0$, and $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$ within the range of $0.3 < x < 1$ can be individually targeted, by controlling the operational conditions of GDEx. At $x = 0$, the initially-colorless solution of copper chloride (*i.e.*, at pH 3) turns into an opalescent and greenish-colored dispersion upon reaching pH ~ 5 . The color change (see Fig. ESI 1†) can be attributed to the precipitation of $\text{Cu}_2(\text{OH})_3\text{Cl}$ (see the green line (a) in Fig. 2), by following reaction (1), in analogy to the mechanism proposed by Engelbrekt *et al.*¹⁶



A plateau of precipitation can be distinguished at pH 4.7–5.5, where 1630 C L^{-1} were consumed from a starting pH of 3. Upon reaching a pH of 5.5, the Cu^{2+} ions were fully removed from the solution (see Fig. ESI 1†), followed by a sharp increase of pH, up to 11—which was reached at a charge consumption of 1875 C L^{-1} . Tenorite (CuO) formed rapidly under alkaline conditions (black line), according to reaction (2).



The formation of $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$ with $x = 0.3$ can be individually targeted with a combination of low charge consumption (778 C L^{-1}), and a relatively low concentration of Cu^{2+} (7.3 mM) and Zn^{2+} (2.3 mM) as the metal precursors as shown in the red line (c) of Fig. 2.

The pH vs. charge evolution of sample $x = 1$, represented by the blue line (d) in Fig. 2, is distinct from that with $x = 0$, represented by the black line (b) in Fig. 2.

As zinc ions are substituted onto the interlayer site, a shorter plateau of precipitation is distinguished at pH 4.7–5.5, wherein 1000 C L^{-1} had been consumed from a starting pH of 3. This is attributed to the insertion of chloride ions as part of the chemical structure in the product form, which—concern-



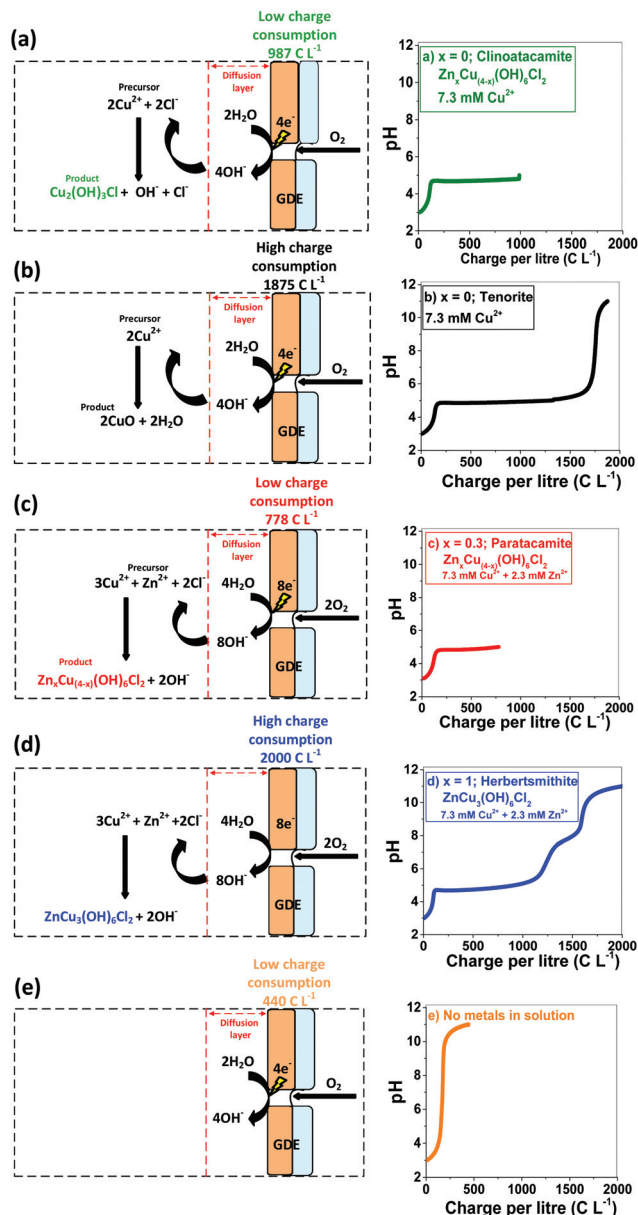
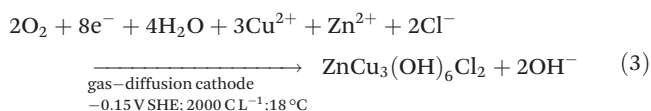


Fig. 2 A suggested mechanism for the synthesis of spin transition nanoparticles. Effect of charge consumption on structural control of (a) $\text{Cu}_2(\text{OH})_3\text{Cl}$ vs. (b) CuO and stoichiometric composition control of spin transition compounds ($\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$) for (c) $x = 0.3$ (paratacamite) and (d) $x = 1$ (herbertsmithite). (e) The control experiment without metals in solution.

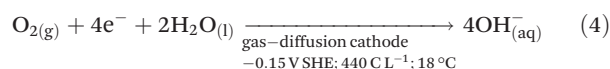
ing CuO implies a reduced consumption of hydroxide ions, as apparent from comparing reaction (2) vs. reaction (3).



The extent of copper and zinc removal from the liquid phase is shown in Fig. ESI 2.† At a higher Cu^{2+} concentration of 574 mg L^{-1} and a Zn^{2+} concentration of 91 mg L^{-1} , 100% of

Cu^{2+} and 53% of Zn ions are removed from the liquid phase at a charge consumption of 2460 C L^{-1} , while 300 mg L^{-1} of Cu^{2+} and 106 mg L^{-1} achieved 100% of Cu^{2+} and 44% removal of Zn ions at a charge consumption of 1810 C L^{-1} .

The orange line in Fig. 2 was obtained, as a control experiment without metals in solution. The charge needed to raise the pH from 3 to 11 was 440 C L^{-1} . The generation of peroxides and free radicals are transient intermediaries, formed together with OH^- ions—as per the established mechanism of O_2 reduction at non-catalyzed carbon electrodes.^{23,24} After oxygen diffuses to the electrocatalytic surface (*i.e.*, activated carbon) of the gas diffusion electrode (GDE), O_2 is reduced (see Fig. 2e). The imposed cathodic polarization conditions (*e.g.*, $-0.15 \text{ V}_{\text{SHE}}$) drive this electrolysis mainly to OH^- ions *via* a 4 electron (4e^-) transfer (eqn (4)), as described in reactions proposed by Šljukić *et al.*²³



From the charge consumed, it is estimated that a profuse amount of OH^- was produced (*e.g.*, $>1300 \text{ mol m}^{-3}$) resulting from the oxygen reduction reaction, from early stages of the GDEx process.²⁰ This would immediately result in a pH of 14 within the porosity of the gas-diffusion cathode,²⁴ facilitating the onset for hydroxide supersaturation and hence for reactive precipitation (*i.e.*, crystallization) at the electrochemical interface.

Spin transition nanoparticles characterization

The morphologies of the $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$ particles made by GDEx, at different charge consumption extents, in which $x = 0$ for clinoatacamite and $0.3 < x < 1$ for paratacamite and $x = 1$ for herbertsmithite, are shown in Fig. 3(a.2, c2, d2). The experimental evidence for $x = 0$, rendered (a2, a3) clinoatacamite ($\text{Cu}_2(\text{OH})_3\text{Cl}$), which was produced as a mixture of faceted nanocrystals and large spiky nanowhiskers ($16 \pm 0.4 \text{ nm}$), when using a charge consumption of 987 C L^{-1} , while only tenorite (CuO) microwhiskers of $1.5 \mu\text{m}$ (b2, b3), with needle-structures, were found at a high charge consumption of 1876 C L^{-1} . This finding is consistent with that of Engelbrekt *et al.*,¹⁶ who demonstrated differences in the morphology of CuO microparticles concerning $\text{Cu}_2(\text{OH})_3\text{Cl}$ nanoparticles, which depended strongly on the reaction conditions by pH-controlled precipitation. Conversely, by adding the copper and zinc precursors in an adequate ratio, the formation of highly crystalline nanoparticles with sizes that seemingly correspond to single crystals of $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$, is achieved. For samples with a stoichiometric coefficient of $x = 0.3$ (c2, c3), the nanocrystals size was $(14 \pm 0.2 \text{ nm})$, obtained with a charge consumption of 778 C L^{-1} . When the stoichiometric coefficient was $x = 1$ (d2, d3), the nanocrystals size was $10 \pm 0.1 \text{ nm}$, with a charge consumption of 2000 C L^{-1} .

The frequency of the spin transition nanoparticles sizes and the corresponding normal distribution fits are shown in Fig. ESI 3.† The addition of Zn^{2+} had a significant effect on the



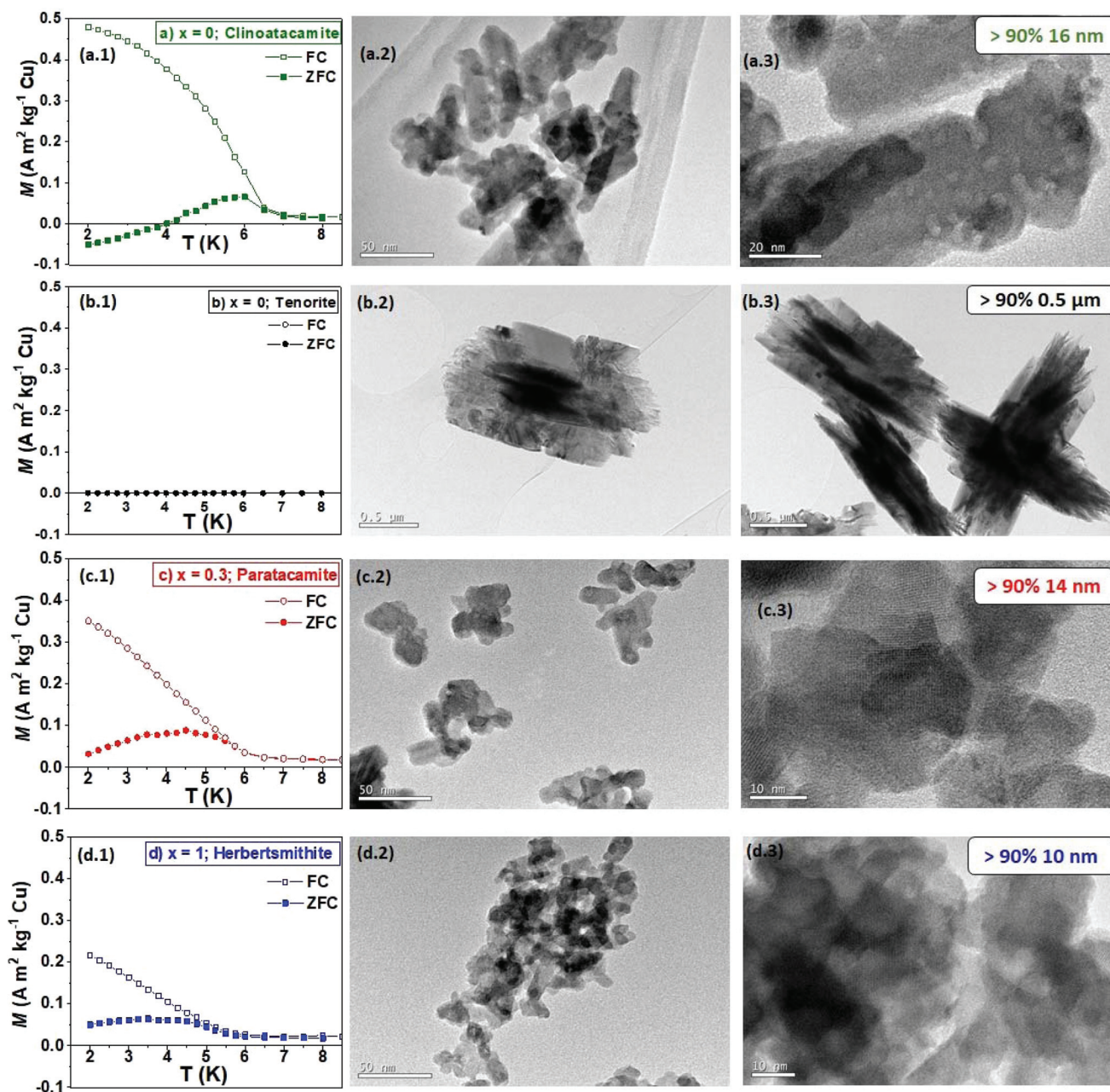


Fig. 3 Temperature dependence of mass magnetization (M) at 8 kA m^{-1} (100 Oe) for $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$ and CuO as measured under ZFC–FC conditions; (a1–a2) for clinoatacamite; (b1–b2) for tenorite; (c1–c2) for paratacamite and (d1–d2) for herbertsmithite. TEM imaging for $\text{Cu}_{4-x}\text{Zn}_x(\text{OH})_6\text{Cl}_2$ products made by GDEx using 6 mM Cu^{2+} and 2 mM Zn^{2+} as precursors.

particle size and polydispersity degree. When the content of zinc was increased from $x = 0.3$ to $x = 1$, particle sizes reduced from 14 to 10 nm, even at higher alkaline pH (*i.e.*, 11). The absence of Zn in the structure, under alkaline conditions, allows the rapid decomposition of $\text{Cu}_2(\text{OH})_3\text{Cl}$ to CuO micro-whiskers of $1.5 \mu\text{m}$ with needle-structures.

The synthesis of particles at the nanoscale requires very specific supersaturation conditions, such as (1) short nucleation and growth periods, (2) at high rates, and (3) avoiding their

immediate agglomeration.²⁵ Correspondingly, the synthesis of spin transition nanoparticles by GDEx, relies on these features, provided by the nature of the process and reagents employed, combined with the following aspects. The flowing conditions impose only a transient contact of metal ion precursors with the reactive species at the saturated electrochemical interface—rendering transient nucleation conditions.

The possibilities of growth are feeble, as encountering other metal ions is restricted by their high dilution. The rate of

interfacial processes typically scales with available surface area (*i.e.*, surface active sites) which is provided by the highly-porous GDE (*i.e.*, $>800 \text{ m}^2 \text{ g}^{-1}$), facilitating a high rate of production of HO_2^- , OH^- and their radicals, which quickly react with the metal ion precursors available at the interface.

Notably, the synthesis of $\text{ZnCu}_3(\text{OH})_6\text{Cl}_2$ is fast under the operational conditions here employed, with rates approaching 40 mg min^{-1} . The process is highly reproducible, and it involves mild synthetic conditions (*e.g.*, 291 K and atmospheric pressure), in contrast to all previous options reported thus far, such as hydrothermal or solvothermal methods (*e.g.*, $\sim 458\text{--}473 \text{ K}$).^{6,15}

The XRD patterns and IR spectra of 4 representative samples are shown in Fig. 4. The diffraction patterns of Fig. 5A are indexed with the following compounds; $\text{Zn}_{0.25}\text{Cu}_{3.75}(\text{OH})_6\text{Cl}_2$ (ICSD-no. 192076), $\text{Zn}_{0.85}\text{Cu}_{3.15}(\text{OH})_6\text{Cl}_2$ (ICSD-no. 424325), ZnO (ICSD-no. 26170), $\text{Cu}_2(\text{OH})_3\text{Cl}$ (ICSD-no. 64956) and CuO (ICSD-no. 67850).

Table S1† compares the results obtained from quantitative phase analysis (QPA) by Rietveld refinement method and ICP-OES of powder samples. The synthesis of $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$ compounds using less than 1000 C L^{-1} , resulted in a product with stoichiometric coefficient $x = 0.3$ and the lowest amount of impurities, such as ZnO and $\text{Cu}(\text{OH})_2$ (Table S1†). At higher charge consumption (2000 C L^{-1}), the synthesis of compounds with higher Zn content ($x = 1$) is differentiated, but a lower purity is obtained, due to the final pH of synthesis of 11 (see Fig. 2 and duplicate experiment data (d1, d2) in Table S1†).

The XRD traces of $\text{Cu}_2(\text{OH})_3\text{Cl}$ and $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$ are indistinguishable by X-ray analysis (see Fig. 4A(a) vs. A (c)) due to the difficulty to differentiate Cu and Zn. The crystal structure of $\text{Cu}_2(\text{OH})_3\text{Cl}$ has been determined by several authors,^{10,16,26} which reported that pure $\text{Cu}_2(\text{OH})_3\text{Cl}$ occurs in the form of three different polymorphs: atacamite, clinoatacamite, and paratacamite. While the first two polymorphs constitute the thermodynamically-stable phase of $\text{Cu}_2(\text{OH})_3\text{Cl}$ at ambient temperatures (orthorhombic and monoclinic structure type), paratacamite (rhombohedral structure) has to be stabilized by partial substitution of Zn or Ni for Cu.^{26,27} According to the experimental work of Malcherek and Schlüter,²⁶ the XRD diffraction pattern of $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$ is weak and therefore easily mistaken by that of $\text{Cu}_2(\text{OH})_3\text{Cl}$. Differences in the hydroxyl groups of clinoatacamite and herbertsmithite can be used for appropriate identification.

Thus, infrared spectroscopic (IR) studies of $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$ and CuO were undertaken to overcome the barrier of crystal structure identification encountered with XRD (Fig. 4B). Our data indicate that $\text{Cu}_2(\text{OH})_3\text{Cl}$ and $\text{ZnCu}_3(\text{OH})_6\text{Cl}_2$ have close lying bands in the hydroxyl stretching region. However, a buried peak at 3400 cm^{-1} is unique for the $\text{Cu}_2(\text{OH})_3\text{Cl}$ structure and can be used as a fingerprint for its detection. Also, the slight shifts in the three major bands of the hydroxyl stretching region, allow us to confidently identify our synthesized $\text{Cu}_2(\text{OH})_3\text{Cl}$ as clinoatacamite. The IR spectrum of the synthesized CuO compounds supports the phase purity observed by XRD, with no indication of

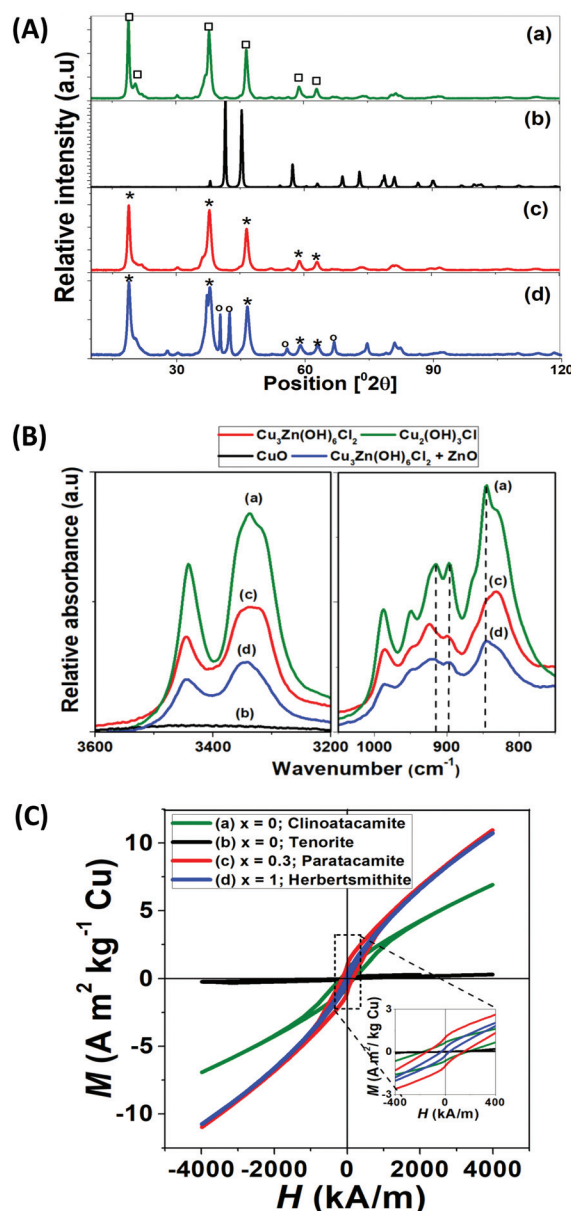


Fig. 4 (A) XRD patterns Co-K α ($\lambda = 1.7928 \text{ \AA}$) of solid products obtained using 7.3 mM Cu^{2+} and 2 mM Zn^{2+} as metal precursor. From top to bottom: Spin transition compounds with formula $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$ at $x = 0$, XRD patterns of (a) clinoatacamite ($\text{Cu}_2(\text{OH})_3\text{Cl}$) $x = 0$ was produced using a charge consumption of 987 C L^{-1} (green line), while only (b) copper oxide (CuO) $x = 0$, at a charge consumption of 1876 C L^{-1} (black line). (c) Paratacamite $x = 0.3$ (red line); (d) herbertsmithite $x = 1$ + impurity of zincite (OZnO) (blue line). The asterisk and circular markers show the most prominent peak positions of $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$ and ZnO phases respectively. The square marker represents the polymorph $\text{Cu}_2(\text{OH})_3\text{Cl}$. (B) FTIR absorption spectra of the synthesized $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$ with $x = 0$ (a) and $\text{Cu}_2(\text{OH})_3\text{Cl}$ and (b) CuO ; (c) $x = 0.3$ and (d) $x = 1$. The spectra graph is shown in two regions of the hydroxyl stretching ($3500\text{--}3200 \text{ cm}^{-1}$) and deformation ($1000\text{--}800 \text{ cm}^{-1}$). Characteristic frequencies are indicated with vertical dashed lines. (C) M against field over temperature sweep at 2.0 K . Stoichiometric coefficient on the interlayer site ranges of x between 0 to 1.



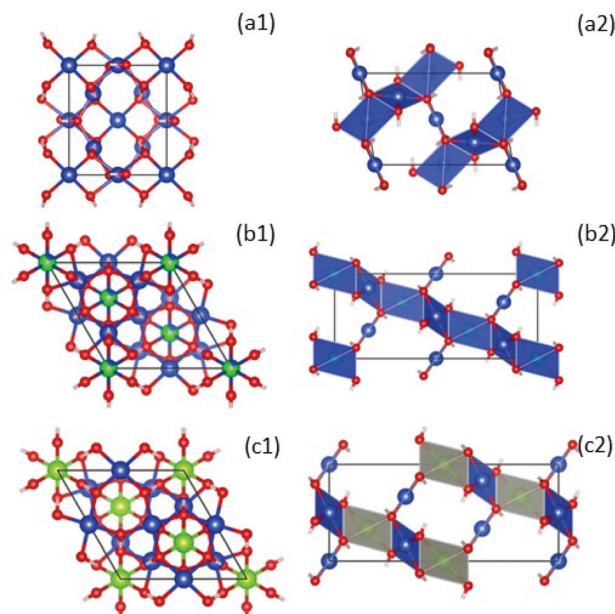


Fig. 5 Left: Crystal structures of compounds with formula $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$ at $x = 0$, clinoatacamite (a1),³¹ Zn-paratacamite (b1)¹⁰ and herbertsmithite (c1)⁹ looking down along the hexagonal c axis. Copper atoms are blue, zinc atoms are green, and oxygen (red). Chlorine and hydrogen atoms have been suppressed for clarity to emphasize better that the CuO_4 units. Right: A side view of the crystal structure using a polyhedral representation, emphasizing the stacking of the kagome layers due to the coupling of CuO_4 planar units by ZnO_6 octahedra in herbertsmithite (c2). This figure was made using the Vesta – JP Minerals software, <https://jp-minerals.org/vesta/en/>, retrieving the XRD data indexed with the following compounds: $\text{Zn}_{0.25}\text{Cu}_{3.75}(\text{OH})_6\text{Cl}_2$ (ICSD-no. 192076), $\text{Zn}_{0.85}\text{Cu}_{3.15}(\text{OH})_6\text{Cl}_2$ (ICSD-no. 424325), $\text{Cu}_2(\text{OH})_3\text{Cl}$ (ICSD-no. 64956), which correspond to the theoretical charts closest to the stoichiometries we achieved experimentally (i.e., see Rietveld analysis in Table S1†).

$\text{Cu}_2(\text{OH})_3\text{Cl}$, and the stability of $\text{ZnCu}_3(\text{OH})_6\text{Cl}_2$. Similar observations are reported using Raman spectroscopy and IR.^{16,28}

Magnetic properties of spin transition nanoparticles

The thermal and field dependence of magnetization (M) over a temperature sweep down to 2.0 K are also reported in Fig. 3 (a1, b1, c1 and d1) and 4C. The ZFC and FC thermal dependence of magnetization provides information about the magnetic properties of the spin transition nanoparticles at 8 kA m^{-1} (100 Oe) (see Fig. 3(a1, c1 and d1)). Tenorite (b1) is an antiferromagnetic semiconductor with Néel temperature (T_N) of about 230 K (Fig. ESI 4†). The ZFC–FC curves and hysteresis loops (Fig. 4C) are typical for CuO samples.²⁹

On the contrary, on cooling and below $T \sim 8 \text{ K}$, the spin transition nanoparticles with formula $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$ showed a transition, which also involves a strong antiferromagnetic (AFM) coupling down to 2 K, especially for clinoatacamite with $x = 0$. The strong AFM coupling of Cu atoms in the triangular lattice makes that the magnetic moments align against the applied field, giving as a result the negative magnetization reversal at low temperatures, especially in clinoata-

camite (a.1). As the temperature increases, the small ferromagnetic (FM) contribution coming from the frustrated geometrical surface becomes to dominate. This effect has been observed in different systems (see “the phenomenon of negative magnetization and its implications”)³⁰ and a similar behavior has been reported in $\text{Cu}_2(\text{OH})_3\text{Cl}$.^{31,32}

The negative magnetization shown for clinoatacamite allows us to differentiate between Zn-paratacamite compounds. The high magnetization value at FC in the clinoatacamite also confirms the coupling between a weak FM and a strong AFM interaction. These results are consistent with those of Colman *et al.*,² who indicated the presence of a small amount of ferromagnetic impurities.

The introduction of non-magnetic Zn in the pyrochlores lattice produces changes in the crystalline structure (from monoclinic to rhombohedral) and the replacement of Cu^{2+} by non-magnetic Zn^{2+} weakens the AFM coupling and finally leads to a system of uncoupled spin, as can be seen from ZFC–FC curves for $x = 0, 0.3$ and 1.

Fig. 5 shows the crystal structure of $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$ compounds, where clinoatacamite ($x = 0$) is a 3D distorted pyrochlore lattice antiferromagnet (a1; a2). The Zn^{2+} ions (green atoms) can be incorporated into the monoclinic clinoatacamite and converted to a rhombohedral structure with undistorted kagome layers forming a 3D framework structure.^{5,11}

The presence of Zn^{2+} ions in paratacamite (b1; b2) and herbertsmithite (c1; c2) decrease the ordering temperature from 6.5 K to 4.5 K, which is an indication of spin frustration.

Herbertsmithite ($x = 1$) is proposed as an ideal 2D kagome with $S = 1/2$, and no long-range magnetic ordering is observed down to 50 mK.

Nanoparticles with a stoichiometric coefficient of $x = 1$ showed less long-range antiferromagnetic (AFM) ordering or spin freezing than paratacamite nanoparticles ($x = 0.3$). On the contrary, the higher $T_N = 6.5 \text{ K}$ of $\text{Cu}_2(\text{OH})_3\text{Cl}$ with $x = 0$ compared to the $T_N = 4.5 \text{ K}$ of compounds with $x = 1$ supports the stronger AFM coupling of the former; besides, the huge magnetization under FC of samples with $x = 0$ confirms the presence of spin glass due to spin frustration of the magnetic moments at the surface.

Magnetization against field over temperature sweep at 2.0 K was recorded to determine the nature of the magnetic component (Fig. 4C). The hysteresis loops at 2.0 K of $\text{Cu}_2(\text{OH})_3\text{Cl}$ shows a kick at low fields confirming the presence of two magnetic phases, one FM and the other AFM (the majority one). The high coercive field ($H_c = 2500 \text{ Oe} = 200 \text{ kA m}^{-1}$) indicates also that the $\text{Cu}_2(\text{OH})_3\text{Cl}$ is a strong AFM.

Compounds with stoichiometry of $\text{Zn}_x\text{Cu}_{4-x}(\text{OH})_6\text{Cl}_2$ showed a paramagnetic-like behavior above the transition temperature and different magnetic couplings at low temperatures (see duplicate experiments in ESI 5†), where $\text{Cu}_2(\text{OH})_3\text{Cl}$ developed a stronger magnetic coupling than compounds with $x = 1$. The AFM behavior is weakened by the introduction of non-magnetic Zn^{2+} , indicating the formation of spin–spin correlations; which is a common feature of quantum kagome systems.^{2–4,6,13,33}



The less ordering temperature in herbertsmithite is a clear indication of spin frustration, which inhibits the tendency of a spin to order under cooling conditions. However, the Néel temperature has not been suppressed as described in the synthesis of a single crystal of herbertsmithite.³

This result may be explained by the fact that below a critical size, magnetic particles become single domain in contrast with the usual multidomain structure of microcrystalline samples, which can exhibit unique phenomena such as unusual coercivities,³⁴ quantum tunneling of the magnetization,^{35,36} and superparamagnetism.^{37,38} Another possible explanation for this is that the size reduction of herbertsmithite could affect the novel magnetic properties, due to the small volume or the high surface/volume ratio (spin canting effect),^{39,40} a phenomenon through which spins show a lack of full alignment at the surface. For example, Martínez *et al.*⁴¹ found a low-temperature surface spin-glass layer in Fe₂O₃ magnetic nanoparticles with a ferromagnetic core, indicating a magnetic disorder in a well-delimited surface layer of the nanoparticle. Aliyu *et al.*⁴² also found a low-temperature surface spin-glass layer in Pr_{0.5}Ca_{0.5}MnO₃ very crystalline nanoparticles with an antiferromagnetic core and a ferromagnetic shell of a single unit cell thickness; the origin of the glassy state is the magnetic frustration of the ferromagnetic order of the shell.

The ferromagnetic component has been recently explained by muon spin rotation (μ SR) studies, where a weak ferromagnetic moment appears in the paratacamite family, one around 18 K and the other one around 6 K, and where dynamics persist down to $T = 0$.^{8,31}

However more in-depth investigation is needed to shed light on a spin canting effect in our spin transition nanoparticles. Although Zn_xCu_{4-x}(OH)₆Cl₂ compounds with $x = 1$ showed less long-range antiferromagnetic (AFM) ordering than nanocrystals with a stoichiometric coefficient of $x = 0.3$ down to 2 K, further research is required to elucidate if spin canting is responsible for the ferromagnetic component observed, which in turn could increase the long-range antiferromagnetic (AFM) ordering of spin-transition nanoparticles.

Conclusions

The possibility of synthesizing spin transition nanoparticles using an electrochemical process with high and fast production rates is desirable for many reasons, notably the convenience of significantly reducing the temperature of synthesis and chemicals, providing a green route of synthesis.

The magnetic properties, particle size and rate of the reaction can be precisely tuned (directly or indirectly) by manipulating the charge consumption, as well as the composition of the electrolyte. This process allows the careful preparation of Cu_{4-x}Zn_x(OH)₆Cl₂ products. The synthesis is fast, reproducible and it involves mild synthetic conditions, running at 291 K and atmospheric pressure, in contrast previously reported methods.

We found faceted nanocrystals of around 10.4 ± 0.1 nm for Zn_xCu_{4-x}(OH)₆Cl₂ products with stoichiometric coefficient between $x = 0.3$ to 1. At a charge consumption of 987 C L^{-1} , we produced a mixture of faceted nanocrystals and large spiky nanowhiskers of 16.1 ± 0.4 nm with a stoichiometric coefficient of $x = 0$. Only microwhiskers of CuO (1.5 μm) with needle-structures were found at higher charge consumption of 1876 C L^{-1} .

Magnetic analyses demonstrate that nanoparticles with $x = 1$ in the formula Zn_xCu_{4-x}(OH)₆Cl₂ showed a small magnetic transition at $T_N = \sim 4.5$ K, with less long-range antiferromagnetic (AFM) ordering or spin freezing than nanoparticles with $x = 0.3$ down to 2 K.

ZFC-FC thermal dependence of M shows bifurcation during the transition at $T_N = \sim 6$ K in samples of stoichiometric coefficient between 0 and 0.3, which supports a long-range AFM ordering, spin freezing or spin-glass-behavior.

These results are in accord with recent studies of synthesis of spin transition compounds indicating that the presence of Zn allows the perfect kagomé distribution of the Cu²⁺ ions and consequently lead the absence of long-range ordering to the lowest measured temperature.

We finally suggest that the spin liquid behavior is sustained at the nanoscale in compounds of ZnCu₃(OH)₆Cl₂. Our discovery not only confirms redox reactions as the driving force to produce spin transition nanoparticles, but also proves a simple way to switch between these magnetic ground states within an electrochemical system, paving the way to further explore its reversibility and overarching implications.

Conflicts of interest

There are no conflicts to declare.

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