Upconversion luminescence in sub-10 nm β-NaGdF₄:Yb³⁺,Er³⁺ nanoparticles: an improved synthesis in anhydrous ionic liquids†

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Sub-10 nm β-NaGdF₄:18% Yb³⁺,2% Er³⁺ nanoparticles were synthesized in ethylene glycol and various ionic liquids under microwave heating. The products were characterized by powder X-ray diffraction, electron microscopy, and upconversion (UC) luminescence spectroscopy. After Yb³⁺ excitation at 970 nm, Er³⁺ ions are excited by energy transfer upconversion and show the typical green and red emission bands. The UC luminescence intensity was optimized with respect to reactant concentrations, solvents, and reaction temperature and time. The strongest UC emission was achieved for sub-20 nm core–shell nanoparticles which were obtained in the ionic liquid diallyldimethylammonium bistrifuoromethanesulfonilamide from a two-step synthesis without intermediate separation. Strictly anhydrous reaction conditions, a high fluoride/rare earth ion ratio, and a core–shell structure are important parameters to obtain highly luminescent nanoparticles. These conditions reduce non-radiative losses due to defects and high energy acceptor modes of surface ligands. A low power excitation of the core–shell particles by 70 mW at 970 nm results in an impressive UC emission intensity of 0.12% compared to the bulk sample.

1. Introduction

About a decade ago, the work of Haase and Güdel et al. initiated the investigation of upconversion (UC) luminescence from transparent solutions of nanoparticles.¹ ² They reported a synthetic route to cubic α-NaYF₄:Yb³⁺,Er³⁺ and hexagonal β-NaGdF₄:Yb³⁺,Er³⁺ nanoparticles dispersed in dimethyl sulfoxide. After 980 nm laser excitation of Yb³⁺, blue, green, and red UC emission from Er³⁺ was observed from clear, colorless solutions. Since then the number of publications on upconversion grew exponentially, about half of them investigating nanoparticles, most prominent with Yb³⁺ and Er³⁺ cooping.³ Auzel gave a comprehensive review on UC ions and processes.⁴ Hexagonal sodium rare earth fluorides, β-NaREF₄ (RE = rare earth), are among the most efficient UC host lattices.⁵–⁷ Two crystallographically distinct RE sites were structurally and spectroscopically characterized in β-NaGdF₄:Er³⁺.⁸ Both have nine-fold coordination, which implies non-centrosymmetric RE sites. Microscopic disorder gives rise to line broadening and a higher probability for spectroscopic overlap of the UC steps. Due to longer RE−F distances and a higher coordination number, hexagonal β-NaREF₄ has lower effective phonon energy than the cubic α-phase. Accordingly, the UC emission intensity of β-NaREF₄ is about an order of magnitude stronger compared to the respective α-NaREF₄ host lattice.⁹ UC mechanisms were investigated in detail,⁴ and recently it was shown that the Er³⁺ ⁴F₉/₂ → ⁴I₁₅/₂ emission in β-NaYF₄ originates from cross-relaxation but not from multi-phonon relaxation.¹⁰ Syntheses for sodium rare earth fluoride nanoparticles were reported by many groups.¹¹–¹⁴

UC nanoparticles were used in various applications, e.g. biological imaging, photodynamic therapy, and nano-thermometry.¹⁵–²⁰ Continuous improvements of the syntheses yielded nanoparticles of smaller size and stronger luminescence;²¹–²⁶ water-dispersible particles are more prone for biological imaging applications.²⁷–³⁰ UC luminescence nanoparticles provide several advantages as biological fluorophores. They show low light scattering, no auto bleaching, and near infrared excitation light has a deep tissue penetration depth. Both in vitro and in vivo experiments led to encouraging results without significant hints for any cytotoxicity of sodium rare earth fluoride nanoparticles.¹⁵

Syntheses of UC nanoparticles in high boiling solvents or under hydrothermal conditions are meanwhile well established.¹,³¹–³⁵ Often, oleic acid is used as coordinating ligand. Oxide ions from its carboxylic groups can substitute for fluoride
in the rare earth coordination sphere on the particle surface due to the similar size of O and F. High energy vibration modes of the ligands increase losses by multi-phonon relaxation.\textsuperscript{31} The high surface-to-volume ratio results in a significant number of (surface) defects in nanoparticles which decrease the luminescence compared to the bulk material. Accordingly, the UC emission from nanoparticles is still about three orders of magnitude lower than for the respective \( \mu \text{m}-\)size phosphor powder.\textsuperscript{31,33} A shell of undoped host material efficiently protects the active core and reduces quenching from surface defects and ligands.\textsuperscript{31,33} Finally, surface functionalization is required to achieve water-dispersible nanoparticles.\textsuperscript{31}

Microwave heating was successfully applied for syntheses in polar or ionic solvents.\textsuperscript{34} The homogeneous heating results in lower temperature and shorter time for the reaction.\textsuperscript{35,36} Ethylene glycol (EG) and ionic liquids (ILs) were proficiently used in microwave syntheses of luminescent materials.\textsuperscript{37-43} Uniform nucleation of particles was achieved which results in a narrow particle size distribution. ILs with complex fluoride anions such as BF\textsubscript{4}\textsuperscript{−} have a dual use as solvent and fluoride source. Cubic \( \alpha \)-NaYF\textsubscript{4}:Yb\textsuperscript{3+},Er\textsuperscript{3+} nanoparticles were obtained from a microwave synthesis in BF\textsubscript{4} ILs.\textsuperscript{39,44} A synthesis in the imidazolium IL 1-butyl-3-methylimidazolium tetrafluoroborate, [C\textsubscript{6}mim] [BF\textsubscript{4}], yields phase pure hexagonal nanoparticles.\textsuperscript{44} The ions determine the heating efficiency\textsuperscript{45} and the properties of the IL, e.g. thermal stability, dielectric constant, ionic conductivity, and water content.\textsuperscript{45-47} The dielectric heating depends on the dipole relaxation and ionic motion of the IL and increases with the microwave absorption. A comparison between tetrafluoroborate (BF\textsubscript{4}) and bis(trifluoromethanesulfonyl)amidine (NTf\textsubscript{2}) ILs revealed that asymmetric NTf\textsubscript{2} heat more efficiently than symmetric BF\textsubscript{4} anions,\textsuperscript{48} because the dipole relaxation takes place only in asymmetric anions. Hydrophobic anions such as NTf\textsubscript{2} result in lower moisture content of the IL than for hydrophilic anions\textsuperscript{48} and such reduce the number of OH\textsuperscript{−} defects in fluoride products. In the present paper, we investigate the influence of ILs with diallyldimethylammonium cations and several different counter anions on the synthesis of \( \beta \)-NaGdF\textsubscript{4}:Yb\textsuperscript{3+},Er\textsuperscript{3+} nanoparticles with respect to phase purity and particle size. The synthetic parameters, as well as the core, core/shell, and ligand design, were optimized for UC luminescence intensity.

2. Experimental

2.1. Sample preparation

2.1.1. Ionic liquid synthesis and purification. The ILs for our investigations were prepared by a metathesis reaction from diallyldimethylammonium chloride (DADMA Cl, Sigma Aldrich, 65% w/w in water) and the respective sodium, silver, or lithium salt in saturated aqueous solution with a 2% stoichiometric excess of the salt.\textsuperscript{49} DADMA OTf (trifluoromethanesulfonate) and DADMA BF\textsubscript{4} (tetrafluoroborate) were obtained from the respective sodium salt (NaOTf, Sigma Aldrich, 98%; NaBF\textsubscript{4}, Sigma Aldrich, 98%), following an established synthesis route for DADMA BF\textsubscript{4}.\textsuperscript{45} DADMA TFA (trifluoroacetate) was prepared from a solution of AgTFA (Merck, 98%+) with DADMA Cl results in the precipitation of AgCl, which was filtered off. DADMA NTf\textsubscript{2} (bis(trifluoromethanesulfonyl)amide) was synthesized from LiNTf\textsubscript{2} (Alfa Aesar, 98+%), as reported in the literature.\textsuperscript{46} The products were decolorized over an active carbon column,\textsuperscript{49} and finally dried at 80 °C under dynamic vacuum overnight. DADMA ILs were obtained as clear, transparent, colorless liquids, and stored under nitrogen atmosphere.

2.1.2. Ethylene glycol purification. EG (Dr Grogg Chemie AG, Switzerland) was purified and dried as described for the ILs above. This equivalent treatment avoids any bias in the comparison between the different solvents due to the purification process.

2.1.3. Core nanoparticle synthesis. \( \beta \)-NaGdF\textsubscript{4}:18% Yb\textsuperscript{3+},2% Er\textsuperscript{3+} nanoparticles were synthesized from rare earth acetates, NaCl, and a fluoride source in various solvents. The rare earth acetates were prepared from the oxides by dissolving Gd\textsubscript{2}O\textsubscript{3}, Er\textsubscript{2}O\textsubscript{3}, and Yb\textsubscript{2}O\textsubscript{3} (Metal Rare Earth Limited, 5 N) in a mixture of glacial acetic acid (Merck, 99.9%) and bidistilled water at 120 °C. After a transparent solution was obtained, the liquid was dried up on a heating plate. The dry powder was converted into the anhydrous acetates at 180 °C under vacuum. The reaction was monitored by thermogravimetric analysis and powder X-ray diffraction, see Fig. S1 and S2 (ESI).

In a typical reaction, 30 mg (0.083 mmol) anhydrous rare earth acetate RE(ACO)\textsubscript{3} (RE = Gd, Er, Yb) was ground with 10 mg (0.17 mmol) NaCl (Reagent grade, Merck) under inert atmosphere and then mixed with 0.5 ml solvent (EG or IL). The molar ratios were fixed to 80/2/18 for Gd/Er/Yb and 2/1 for Na/RE in all samples. These ratios were optimized for bulk samples before.\textsuperscript{67} The acetate solution was heated to 60 °C for 5 minutes with a Biotage Initiator+ Robot Eight microwave synthesis system (Biotage AB, Uppsala, Sweden). Fluoride-containing ILs or NH\textsubscript{4}F (Alfa Aesar, 96%) were used as fluoride sources; the effects of the reactant (BF\textsubscript{4}− or NH\textsubscript{4}F) and the F− concentration (60–180 mg, 1.62–4.86 mmol) were evaluated. In a microwave glass vial, the fluoride source was added to 1.5 ml of an EG/IL solution or the IL alone and stirred up to homogeneity. The acetate suspension was added dropwise to the microwave vial and stirred overnight before the microwave heating. The temperature in the reactor was controlled by dry compressed air (4 bar), and microwave powers of 400 W or 50 W were used for EG/IL solutions or pure IL, respectively. For separation, the nanoparticles were sequentially washed several times with methanol and ethanol. They were collected by centrifugation at 3000 rpm for 5 min and dried at 80 °C overnight. Alternatively, the nanoparticles were kept in solution for subsequent growth of an undoped shell. Experimental conditions for all samples are summarized in Table 1.

2.1.4. Core-shell nanoparticle synthesis. After the core synthesis, another 1/3 of the NH\textsubscript{4}F mass used in the first step was added to the nanoparticle solution and stirred. A solution of 30 mg gadolinium acetate and 10 mg NaCl in 0.5 ml of the respective solvent was prepared and heated to 60 °C for 5 min by microwave irradiation. The solution was added dropwise to the core nanoparticle solution and stirred for 10 minutes. Afterward, the solution was heated to 200 °C for 5 min with 50 W...
microwave power. The core–shell nanoparticles were separated as described above.

2.2. Experimental methods

X-ray diffraction (XRD) patterns were measured on a StadiP powder diffractometer (STOE, Darmstadt, Germany) in transmission geometry at room temperature. Cu K\(_{\alpha1}\) radiation (\(\lambda = 1.540 598\) Å) was used from a 40 kV 40 mA X-ray source and a focusing Ge (111) monochromator. Layers of nanoparticles were prepared on thin plastic foil. Data were recorded by a Mythen 1K linear position sensitive detector (Dectris, Baden, Switzerland) with 400 m pixels resolution in 2-theta.

Transmission electron microscopy (TEM) analyses were performed on a G2 Spirit BioTWIN TEM (Tecnai) using a ZrO\(_2\)/W (100) Schottky field emitter gun. A Falcon II direct electron detector (FEI) with 4096 × 4096 pixels and an Orius SC200 CCD camera (Gatan Inc.) with 2048 × 2048 pixels were used. The specimens were prepared from diluted ethanol solutions on carbon-coated copper grids (300 mesh).

The pictures were analyzed by the ImageJ 1.49v software (National Institute of Health, USA).

The thermal decomposition of rare earth acetates was investigated under a nitrogen flow of 20 ml min\(^{-1}\) and a heating rate of 5 K min\(^{-1}\) by thermogravimetric analysis using a TGA/SDTA 851e (Mettler-Toledo, Schwerzenbach, Switzerland).

3. Results and discussion

3.1. \(\beta\)-NaGdF\(_4\):Yb\(^{3+}\),Er\(^{3+}\) nanoparticles from EG/IL mixtures

X-ray diffraction patterns of cubic \(\alpha\) and hexagonal \(\beta\)-NaGdF\(_4\):18% Yb\(^{3+}\),2% Er\(^{3+}\) nanoparticles are shown in Fig. 1. Samples were obtained from EG/DADMA BF\(_4\) (EG1) and EG/DADMA NTf\(_2\) (EG2) solutions with BF\(_4\)\(^{-}\) and NH\(_4\)F as fluoride sources, respectively, see Table 1. Both samples were heated for 30 min at 200 °C. Tetrafluoroborate anions decompose at 160 °C and release F\(^{-}\) ions into the solution. Sample EG1 crystallizes in the cubic \(\alpha\)-NaGdF\(_4\) phase. The calculated peak positions are listed in Table 1. The strongest diffraction peak (111) is located at 28° 2-theta.
The pure hexagonal β-NaGdF₄ phase was obtained from an NH₄F solution for sample EG2, see the bottom trace in Fig. 1. The calculated peak positions are shown as blue ticks. β-NaGdF₄ crystallizes in the β-NaNdF₃ structure with space group P6 and lattice parameters of a = 6.0304 (9) Å and c = 3.6111 (7) Å at room temperature.⁸ A F⁻ excess favors the formation of the hexagonal β-NaGdF₄ phase.⁹ BF₄⁻ as fluorine source yields preferentially the cubic α phase in a microwave synthesis, since the slow BF₄⁻ decomposition during heating provides low fluoride concentration, only. The UC luminescence from cubic α-NaGdF₄ bulk material is about an order of magnitude less intense compared to the hexagonal β phase.⁹ Accordingly, NH₄F was chosen as fluorine source in order to obtain hexagonal nanoparticles with strong UC emission.

The influence of DADMA ILs on the microwave synthesis was investigated. The thermal stability of the IL depends on the anion and increases along the series TFA < BF₄ < OTf < NTf₂. The stability determines the maximum reaction temperature, ranging from 120 °C (TFA) to 200 °C (BF₄, OTf, and NTf₂) for EG/IL solutions and 400 W microwave power. Higher temperatures result in partial decomposition of the IL, and the colorless solutions turn brown. Samples were synthesized under identical conditions in different ILs. Their phase purity and UC luminescence are compared in Fig. 2. All products show a pure hexagonal phase in powder XRD, see Fig. 2 (ESI†). The anions strongly influence the β-NaGdF₄:18% Yb³⁺,2% Er³⁺ UC luminescence which increases with the thermal stability of the ILs along the series BF₄ < OTf < NTf₂, see Fig. 2 (ESI†). Asymmetric anions such as TFA, OTf, and NTf₂ have a higher microwave absorption, which results in more effective heating compared to the symmetric BF₄⁻. The luminescence from the DADMA TFA sample was too weak to be compared with the others. Also, the hygroscopicity of the IL is relevant. Samples from hydrophobic DADMA NTf₂ show a stronger luminescence than those from hygroscopic ILs. Lower water contamination of the IL decreases the number of OH⁻ impurities in the nanoparticles and non-radiative losses from these defects. The crucial importance of strictly dry reaction conditions is further supported by samples obtained from anhydrous rare earth acetates versus those from acetate hydrates, see Fig. S5 (ESI†).

The UC luminescence from nanoparticles synthesized from dry precursors is much stronger than from hydrates. Fig. 2 shows the influence of fluoride concentration, temperature, and rare earth concentration on the UC luminescence. The NH₄F content was varied from 60 mg to 120 mg for samples EG3 to EG6, see Table 1. The stability of the hexagonal phase increases with the F⁻ concentration, as shown in Fig. S6† (ESI). The weak (111) diffraction peak of the cubic phase around 28° 2-theta disappears by raising the NH₄F content from 60 to 100 mg. For higher F⁻ concentrations, only the hexagonal phase is obtained. As an example, the pure hexagonal phase was obtained for sample EG2, see the bottom trace in Fig. 1. The calculated peak positions for cubic α-NaGdF₄ (Fm3m) and hexagonal β-NaGdF₄ (P6) are shown as red and blue ticks below the traces, respectively.

Fig. 1. Powder XRD patterns of nanocrystalline α-NaGdF₄:18% Yb³⁺,2% Er³⁺ (EG1, top trace) and β-NaGdF₄:18% Yb³⁺,2% Er³⁺ (EG2, bottom trace). Samples were synthesized in EG/IL solutions with DADMA BF₄ and NH₄F, respectively, see Table 1. The calculated peak positions for cubic α-NaGdF₄ (Fm3m) and hexagonal β-NaGdF₄ (P6) are shown as red and blue ticks below the traces, respectively.
impurity, NaF was identified, which does not influence the UC luminescence. The NaF diffraction peaks have a narrow line width and are marked with asterisks for the top trace of Fig. S6.† The crystallite size was evaluated from the XRD patterns in Fig. S6,† ranging from 6 to 14 nm.

For the UC luminescence spectra in Fig. 2, Yb$^{3+}$ ions are excited at 970 nm. Excited Yb$^{3+}$ ions can transfer energy resonantly to neighboring Er$^{3+}$ ions in several steps. The energy transfer upconversion (ETU) first populates the $^4I_{11/2}$ state and in a second step the $^4F_{7/2}$ state of Er$^{3+}$, which decays via multphonon relaxation to the $^4I_{15/2}$ and $^4I_{13/2}$ states. These states are in thermal equilibrium, and their emission to the $^4I_{15/2}$ ground state yields the typical green Er$^{3+}$ UC luminescence.5,7 The $^4F_{9/2}$ state is populated via cross relaxation.8 Its emission to the ground state adds a red component to the luminescence. The UC luminescence intensity increases 2-fold from sample EG3 to EG6 for higher F/RE ratio and a high reaction temperature strongly increases the aggregate size and increases the UC luminescence.

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From the literature, it is known that higher NH$_4$F concentrations increase the aggregate size of β-NaGdF$_4$:Eu$^{3+}$ and Er$^{3+}$ nanoparticles synthesized at room temperature in EG.5,45 Aggregation reduces the exposed particle surface; concomitantly, the non-radiative relaxation decreases and UC luminescence is not quenched. The same was reported for nanoparticles from octadecene/oleic acid solutions.51 Since nanoparticles from EG solutions are often affected by agglomeration,5,45 reactions in EG/IL mixtures and pure ILs, in particular the hydrophobic DADMA NTf$_2$, were investigated in order to reduce the aggregate size and increase the UC luminescence intensity.

UC luminescence spectra of β-NaGdF$_4$:18% Yb$^{3+}$,2% Er$^{3+}$ nanoparticles from EG/IL mixtures with DADMA OTf and DADMA NTf$_2$ are shown in Fig. 3. The UC emission intensity increases strongly when the EG content is reduced. The by far best results were obtained from syntheses in pure ILs, see the green traces in Fig. 3. Among the pure ILs, the product from hydrophobic DADMA NTf$_2$ (sample IL2) is significantly better than from DADMA OTf (sample IL1). The UC luminescence intensity of sample IL2 is 70-fold higher than for EG11. As discussed above, strictly anhydrous reaction conditions reduce the number of OH$^-$ defects in fluorides and restrict non-radiative losses of the luminescence intensity. Avoiding solvents with OH groups, such as EG, is equally important. These solvents may introduce traces of water into the reaction mixture or act as surface ligands with high energy vibration acceptor modes. Another bonus of the synthesis in pure ILs is an increased microwave absorption; compared to EG/IL mixtures, the microwave power could be reduced from 400 W to 50 W.

3.2. β-NaGdF$_4$:Yb$^{3+}$,Er$^{3+}$ nanoparticles from IL solutions

Reactions in pure ILs yield material with a better UC luminescence compared to EG or EG/IL syntheses, cf. Fig. 2 and 3, but

![Fig. 3](image-url) Upconversion luminescence of β-NaGdF$_4$:18% Yb$^{3+}$,2% Er$^{3+}$ nanoparticles from EG/IL syntheses with DADMA OTf (A) and DADMA NTf$_2$ (B) for various EG/IL ratios. The sample names refer to Table 1. The UC luminescence was excited at 970 nm with 580 mW (unfocused) laser power. Microwave absorption: compared to EG/IL mixtures, the microwave power could be reduced from 400 W to 50 W.

![Fig. 4](image-url) Upconversion luminescence of β-NaGdF$_4$:18% Yb$^{3+}$,2% Er$^{3+}$ nanoparticles from IL syntheses in dependence on fluoride concentration (bottom), reaction temperature (middle), and time (top). The sample names refer to Table 1. All samples were excited at 970 nm with 330 mW (unfocused) laser power.
a different solvent requires the adjustment of reaction parameters. Accordingly, the fluoride concentration, temperature, and reaction time were optimized for syntheses in DADMA NTf2 solutions, see Fig. 4. The ionic liquid has a vapor pressure close to zero and superb thermal stability, which allows higher reaction temperatures. No hints for any decomposition of the IL were detected up to 250 °C, the highest accessible temperature for the microwave system. Due to the stronger microwave absorption of the IL, the reaction time was reduced to 5 min compared to 30 min for EG/IL mixtures.

Increasing the fluoride concentration gives rise to a stronger UC luminescence, as demonstrated for samples IL3 to IL6 in Fig. 4, bottom, and in good agreement with Fig. 2. However, a comparison of Fig. S7 and S6 (ESI†) reveals the presence of α-phase in the XRD diagrams of several IL samples. The α-phase content is reduced from IL3 to IL5 and IL6 shows a pure β-phase. A higher fluoride concentration is required in the IL in order to obtain the pure β-phase. For identical fluoride concentration, e.g. EG vs. IL3, an IL sample shows a much stronger UC luminescence because of a reduced number of defects due to the anhydrous reaction conditions, despite an α/β phase mixture. Higher fluoride concentrations along the IL3 to IL6 series increase the nanoparticle size as evidenced by the narrower diffraction peaks.

The reaction temperature has a stronger effect on the UC luminescence than the NH4F concentration, see Fig. 5A middle. At 225 °C (IL5) the UC luminescence intensity is 34-fold higher compared to 200 °C (IL7), but it drops again by a factor of 12 towards 250 °C (IL8). The increase from 200 °C to 225 °C is due to a higher crystallinity and bigger particle size of the β-phase. At 250 °C, a significant amount of the α-phase is observed which has a weaker UC emission. The higher temperature favors the formation of the metastable α-phase.

Doubling the reaction time from 5 to 10 min (IL8 vs. IL9) reduces the amount of α-phase, see Fig. S7 (ESI†). A longer reaction time supports completion of the α → β phase transition, which enhances the UC luminescence, see Fig. 4, top. Overall, sample IL6 shows the best UC emission. Optimized reaction parameters are a high fluoride concentration, 225 °C, and 10 min reaction time.

TEM micrographs of representative samples are shown in Fig. 5. Particle size and agglomeration increase with fluoride concentration, see Fig. 5A and B for samples IL3 and IL6. The crystallite size increases from 3–4 nm to 15–18 nm, and agglomerates form in the μm size range. Longer reaction times result in larger aggregates, see Fig. 5C and D. From higher magnification in Fig. 5E–G, an individual crystallite size in the order of 4.6 ± 1.1 nm becomes visible for sample IL4. It does not show significant aggregation and provides a good UC luminescence. This sample was further used for a core-shell synthesis, where an undoped β-NaGdF4 shell was grown on top of the β-NaGdF4:2% Er3+,18% Yb3+ particles. The core-shell particles (CS_IL4) have a crystallite size <20 nm and show rather small aggregates, see Fig. 5H. Sample IL4 contains a minor fraction of the α-phase, as evident from its XRD pattern, see Fig. S7 (ESI†). In the subsequent reaction step towards the core–shell sample CS_IL4, a complete conversion to the β-phase was achieved, see Fig. S8 (ESI†).

The UC luminescence of samples from EG (EG11) vs. core (IL4) and core–shell particles (CS_IL4) from IL syntheses is compared in Fig. 6 and 7. Samples from IL syntheses show strongly enhanced UC intensities over the EG sample, see Fig. 6.

Fig. 5 TEM and HR-TEM micrographs of representative samples: (A) IL3 and (B) IL6 from syntheses with 60 mg and 180 mg NH4F at 225 °C; (C) IL8 and (D) IL9 from syntheses with 140 mg NH4F at 250 °C for 5 min and 10 min; (E–G) IL4 from a synthesis with 100 mg NH4F at 225 °C for 5 min; (H) CS_IL4 core–shell nanoparticles. The sample names refer to Table 1.
The green-to-red ratio is representative for the level of multi-phonon relaxation losses in the sample. The $^4F_{9/2}$ state can be populated by cross-relaxations or multi-phonon relaxation from the $^4S_{3/2}$ state. In macroscopic samples, e.g. single crystals, the multi-phonon relaxation is negligible at room temperature for the $\beta$-NaGdF$_4$ host lattice due to its low effective phonon energy. Therefore, the red emission is indicative of impurities or lattice defects, which introduce acceptor modes of high-energy, depopulate the $^4S_{3/2}$ state, and strongly reduce the UC emission in total. The major enhancement of the UC luminescence from sample EG11 to IL4 is the result of a significant reduction of defects and acceptor modes due to an anhydrous synthesis and the absence of ligands with OH groups.

A further significant increase of the UC emission is achieved by growing an undoped $\beta$-NaGdF$_4$ shell around the $\beta$-NaGdF$_4$:18% Yb$^{3+}$:2% Er$^{3+}$ core, see the blue trace in Fig. 6 for sample CS_IL4. The shell saturates any partial coordination at the core surface and separates the particle surface from the optically active core. This approach reduces losses by multi-phonon relaxation via high energy vibration modes of ligand molecules and surface defects.

The UC luminescence from IL4 and CS_IL4 nanoparticles is further compared to $\mu$m-size $\beta$-NaYF$_4$:20% Yb$^{3+}$, 2% Er$^{3+}$ bulk material, see Fig. 8. The samples were excited at 970 nm by only 70 mW (unfocused) laser power. Despite a low excitation density of 2.6 W cm$^{-2}$, see Fig. S9 and Table S1 (ESI†) for excitation density calculation, the nanoparticle samples (red and blue traces) show a well detectable UC emission with a spectral distribution close to the bulk material (black trace). The spectra of the nanoparticle samples are scaled by a factor $\times 500$ in Fig. 8. The UC luminescence from the core (IL4) and core–shell nanoparticles (CS_IL4) are 0.067 (4)% and 0.122 (5)% of the bulk material. The green-to-red intensity ratio is 1.80 for the core vs. 1.66 and 1.52 for the core–shell and core nanomaterials, respectively. The slight decrease of the green-to-red intensity ratio along this series is due to increasing multi-phonon relaxation losses. These results are very good for sub 20 nm particles and compare well with literature.

A synthesis of <6 nm $\beta$-NaGdF$_4$:30% Yb$^{3+}$, 2% Er$^{3+}$ core–shell particles in oleic acid and octadecene at 240 $^\circ$C for 20 min was reported by Liu et al.$^{26}$ The UC luminescence of these particles was excited at 10 W cm$^{-2}$ 5.5 nm $\beta$-NaYF$_4$: Yb$^{3+}$, Er$^{3+}$ particles from an oleic acid, oleylamine, and octadecene solution at 310 $^\circ$C for 30 min were reported by Ostrowski et al.$^{24}$ Their UC
spectra were excited by 1 W. 5–20 nm β-NaYF₄: Yb³⁺, Ln³⁺ (Ln = Er, Tm or Ho) particles were obtained from a synthesis in trioctylphosphine oxide (TOPO), which allows higher reaction temperatures of 330 °C to 370 °C. Samples were synthesized within 30 to 90 min without impurities of the z-phase. 11 nm particles were excited by a focused laser with 1 W power. 8–10 nm β-NaYF₄: Yb³⁺, Er³⁺ particles were synthesized in oleylamine at 330 °C for 1 hour by Boyer et al. They reported quantum yields of 3% and 0.005% for bulk (>>100 nm) and 8–10 nm particles for 20 W cm⁻² and 150 W cm⁻² excitation density, respectively. The UC luminescence intensity of our nanoparticles, see Fig. 8, was achieved by 70 mW unfocused excitation power, corresponding to a power density of 2.6 W cm⁻². This value is more than an order of magnitude lower than most literature values cited above. Incomplete information on the excitation conditions often hampers a direct comparison of UC emission intensities from the literature. Due to the non-linearity of UC processes the measurement of absolute quantum yields and a comparison between different excitation densities is inherently difficult. We restrict our study to a relative comparison of nanoparticles with bulk material, whose high quality was demonstrated in previous studies.

4. Conclusions

The synthesis under anhydrous conditions in the ionic liquid DADMA NTf₂ is a promising path to β-NaGdF₄:18% Yb³⁺,2% Er³⁺ nanoparticles. A microwave reactor allows for efficient heating and short reaction times. Anhydrous conditions and appropriate solvents are essential for the synthesis of highly luminescent upconversion material. These conditions limit non-radiative losses due to defects and high energy acceptor modes of ligand vibrations. Another critical factor is the architecture of the nanoparticles. Core–shell particles show a more intense luminescence compared to the bare core particles. The growth of an isostructural undoped shell saturates coordination environments at the core surface and separates the core from the particle surface. Sub 20 nm core–shell nanoparticles were obtained in two reaction steps without intermediate separation. Very low excitation power was required to excite the upconversion luminescence, which renders these nanoparticles attractive for biological imaging. We proved that the proposed synthesis provides an effective route to minimize the presence of OH-defects, which have been demonstrated to represent the main source of luminescence quenching in fluorides. We believe that further optimization of this microwave-assisted synthesis will yield even more efficient and monodisperse nanomaterials, providing a fast and easily scalable synthesis route to brighter fluoride upconversion nanoparticles.

Conflicts of interest

There are no conflicts to declare.

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Notes and references