



Cite this: *RSC Adv.*, 2017, 7, 55680

## Synthesis of silver nanoplate based two-dimension plasmonic platform from 25 nm to 40 $\mu\text{m}$ : growth mechanism and optical characteristic investigation *in situ*<sup>†</sup>

Xiao-Yang Zhang,<sup>abc</sup> Huan-Li Zhou,<sup>ac</sup> Feng Shan,<sup>ac</sup> Xiao-Mei Xue,<sup>bc</sup> Dan Su,<sup>bc</sup> Yi-Ran Liu,<sup>a</sup> Yu-Zhang Chen,<sup>a</sup> Jing-Yuan Wu<sup>ac</sup> and Tong Zhang <sup>\*abc</sup>

Silver nanoplates (SNPs) with single-crystalline structures are highly expected building blocks for the construction of a two-dimensional (2D) plasmonic platform regarded as an advanced tool for the development of subwavelength light management technologies. One barrier for SNP fabrication is how to achieve structures with a lateral size large enough and a thickness thin enough for large-scale light manipulation. Here we propose a multistep chemical synthesis strategy which overcomes this bottleneck and greatly enriches the morphological features of SNPs by deliberately enhancing the behaviors of selective etching and self-assembly during a dynamic crystal growth process. Growth mechanisms were comprehensively studied with the help of real-time microscopic monitoring *in situ*. Three types of SNP with different morphology features including 100% yield small SNPs with a controllable surface plasmon resonance band, large SNPs decorated with hot spots and giant SNPs (up to  $\sim 40 \mu\text{m}$ ) with an ultra-high aspect ratio (over 1000 : 1) and ultrathin thickness were successfully achieved. Using such an engineered 2D platform combining waveguiding and nanoantenna effects, we demonstrated that plasmon enhanced optical information in deep subwavelength volumes can be remotely excited, transferred and scattered into free space directionally. Subwavelength light transmission with multiple excitation wavelengths and tunable one-dimensional (1D) and zero-dimensional (0D) light scattering as well as photoluminescence enhancement of light emitters was exhibited. It represents significant advances in *in situ* light manipulation which are useful in various applications for nanophotonics and remote sensing.

Received 4th October 2017  
Accepted 22nd November 2017

DOI: 10.1039/c7ra10952k

rsc.li/rsc-advances

### 1. Introduction

Chemical synthesizing and morphology engineering are essential technologies for the achievement of plasmonic nanocrystals with high-quality crystal structures in a rapid and high-yield way,<sup>1</sup> meeting the practical requirements of nanophotonic devices and system fabrication.<sup>2</sup> Compared with lithographically fabricated polycrystalline nanostructures,<sup>3</sup> chemically synthesized nanocrystals having single-crystalline structures with few (even zero) crystallographic defects are superior in the achievement of plasmonic devices with much lower ohmic

damping and better performance.<sup>4,5</sup> Such chemically synthesized nanocrystals serving as surface plasmon polariton (SPP) waveguides initiated innovative applications in nanophotonics, such as optical processing,<sup>2,4-7</sup> logic photonic circuitry,<sup>8</sup> remote excitation of surface enhanced Raman scattering<sup>9,10</sup> and plasmon enhanced subwavelength light sources.<sup>11</sup> However, previously studies on chemically synthesized SPP waveguides were primarily focused on silver nanowires.<sup>2,4-12</sup> The SPP mode supported by these one-dimensional (1D) waveguides only propagates along one direction and scatters into free-space light at their two ends. In contrast, two-dimensional (2D) silver nanoplates (SNPs) with large surface area provide an alternative waveguide structure supporting SPP modes with more diverse modal characteristics and spatial freedom, opening up a new research direction of high performance plasmonic platform construction.<sup>13</sup> As a new research field, only basic plasmonic characteristics of SNPs such as polarization dependent mode propagation and light scattering at edges were demonstrated.<sup>13,14</sup> Actually, the diversity of 2D SPP modal characteristics is dependent on both the size and thickness of SNPs. As there is still a lack of effective synthesis methods of SNPs with

<sup>a</sup>Joint International Research Laboratory of Information Display and Visualization, School of Electronic Science and Engineering, Southeast University, Nanjing, 210096, People's Republic of China. E-mail: tzhang@seu.edu.cn

<sup>b</sup>Key Laboratory of Micro-Inertial Instrument and Advanced Navigation Technology, Ministry of Education, School of Instrument Science and Engineering, Southeast University, Nanjing, 210096, People's Republic of China

<sup>c</sup>Suzhou Key Laboratory of Metal Nano-Optoelectronic Technology, Suzhou Research Institute of Southeast University, Suzhou, 215123, People's Republic of China

<sup>†</sup> Electronic supplementary information (ESI) available: Experimental details and supplementary figures and movies. See DOI: 10.1039/c7ra10952k



an ultra-high aspect ratio (large size and ultrathin thickness), the advantages of an SNP based 2D plasmonic platform have not been comprehensively explored in experiments.

Many efforts have been applied to the synthesis of SNPs with single-crystalline structure using wet-chemical synthesis,<sup>14–24</sup> especially SNPs with large size.<sup>14–21</sup> Among those reported methods, the multi-step synthesis of SNPs in aqueous environment<sup>22–24</sup> is superior in structural uniformity, size and thickness control. Large SNPs with a high aspect ratio were expected to be obtained by the continual increment of reaction cycles. However, the maximum size of SNPs synthesized using such a strategy was less than 5  $\mu\text{m}$  (ref. 22 and 23) which did not meet the practical size requirement of 2D SPP mode manipulation. To overcome this bottleneck, it is of great importance to explore the growth kinetics comprehensively during the crystal growth process. This will motivate the development of SNP synthesis strategies with enhanced morphological controllability.

Here we first reveal the decisive factors limiting the size and yield of SNPs in a dynamic reaction assisted by real-time visualization of the crystal growth process under optical microscopy *in situ*. As the anisotropic growth mechanism and simultaneous existing  $\text{H}^+$  etching behavior during the reaction were clarified, we proposed the addition of an etching agent of hydrogen peroxide ( $\text{H}_2\text{O}_2$ ) and a protection agent of chloride ions ( $\text{Cl}^-$ ) in different stages of the reaction which greatly improved the effectivity of crystal type self-etching and self-screening behaviors during the growth process. With these improvements, the size and thickness limitations of SNP synthesis in aqueous solution were broken through and three types of SNP with well-engineered morphological features were achieved, as illustrated in Scheme 1. They are nano-antennas with controllable localized surface plasmon resonance (LSPR) bands and huge 2D crystal structures having atomically flat surfaces or decorated with hot spots. Based on these morphological engineered huge SNPs, fundamental optical properties of 2D plasmonic platforms were comprehensively explored under optical microscopy *in situ*. Thickness dependent SPP modal distribution, propagation and sub-diffraction limit scattering in far-field, multiplexed broadband optical transmission as well as plasmon enhanced

remote excitation of light emitters of the 2D SNP platforms were investigated. Our achievements indicated that huge SNPs fabricated by low-cost and simple chemical routes can serve as advanced plasmonic platforms and are useful in various fields such as nanoscale light manipulation,<sup>10–14,25–27</sup> nonlinear optical enhancement,<sup>28,29</sup> optoelectronic multiplex<sup>30,31</sup> and quantum information transmission.<sup>32–34</sup>

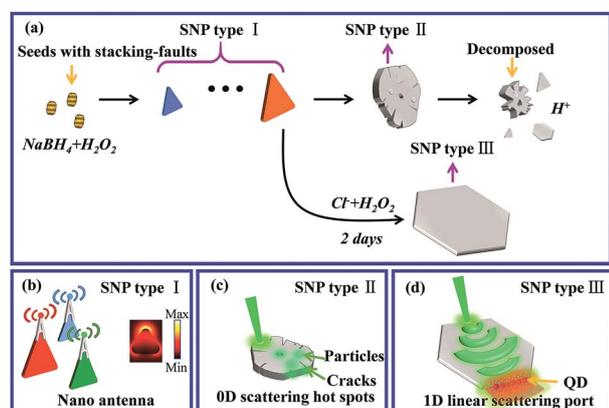
## 2. Experimental

### 2.1 Chemicals

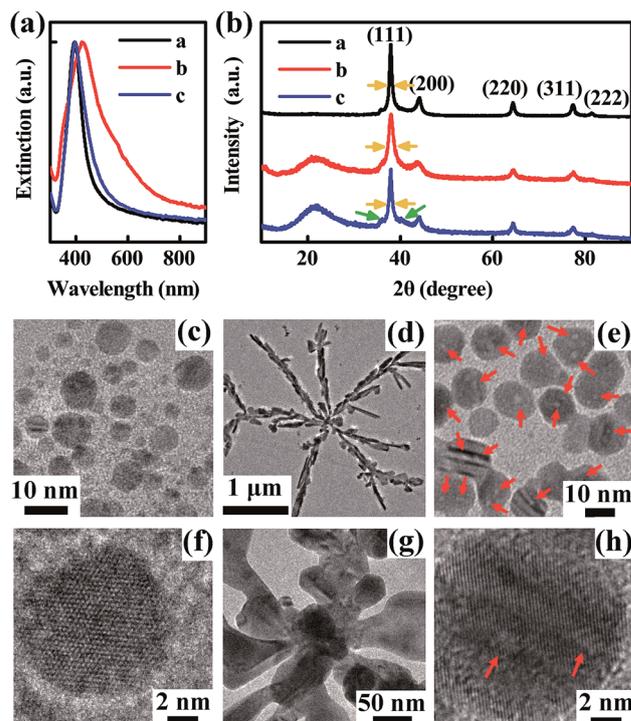
Silver nitrate ( $\text{AgNO}_3$ , 99.9%), sodium borohydride ( $\text{NaBH}_4$ , 98%), sodium citrate (99.9%) and sodium chloride ( $\text{NaCl}$ , 99.9%) were purchased from Sigma-Aldrich. Hydrogen peroxide solution ( $\text{H}_2\text{O}_2$ , 30% w/w) and hydrazine hydrate (99.5%) were purchased from Hujiang Co., Ltd. Ultrapure water ( $>18.2 \text{ M}\Omega \text{ cm}$ ) was used in all experiments.

### 2.2. Methods

**Preparation of silver seeds.** The optimal condition is as follows. 60  $\mu\text{L}$  of 0.5 M  $\text{AgNO}_3$  and 70  $\mu\text{L}$  of 0.5 M sodium citrate were added into 10 mL of water. Then, 180  $\mu\text{L}$  of  $\text{H}_2\text{O}_2$  solution and 500  $\mu\text{L}$  of 0.2 M  $\text{NaBH}_4$  were added into the solution successively under strong stirring. For comparison, different amounts of  $\text{H}_2\text{O}_2$  and  $\text{NaBH}_4$  were used to synthesise different types of seed solution, as demonstrated in Fig. 1. For seed sample a, the seeds were obtained using conventional



**Scheme 1** Schematic synthetic process and application prospect of SNPs with different morphologies and sizes.



**Fig. 1** (a) Extinction spectra of silver seeds synthesized under different conditions. Curves a to c correspond to seed samples a, b and c, respectively. (b) X-ray diffraction (XRD) spectra of the three seed samples. TEM and HRTEM images of seed samples a to c are shown in (c to e) and (f to h), respectively.



methods.<sup>24</sup> H<sub>2</sub>O<sub>2</sub> is absent and the concentration of NaBH<sub>4</sub> is 1/10 of the optimal condition (50 μL of 0.2 M). For seed sample b, 180 μL of H<sub>2</sub>O<sub>2</sub> solution and 50 μL of 0.2 M NaBH<sub>4</sub> were added into 10 mL solution. Seed sample c corresponds to the optimal condition described above.

**Preparation of solutions A and B.**<sup>24</sup> Solution A: 50 μL of 0.4 M hydrazine hydrate (N<sub>2</sub>H<sub>4</sub>) and 50 μL of 0.5 M sodium citrate, serving as the reducing agent and surfactant, were added into 18 mL of water. Solution B: 58 μL of 0.5 M AgNO<sub>3</sub> solution was added into 5 mL of water.

**Method 1: synthesis of SNP type I.** Different amounts of silver seed solution decreased from 3200 to 50 μL were added into solution A.<sup>24</sup> Then solution B was injected into the mixed solution dropwise by a syringe pump at a rate of 10 mL h<sup>-1</sup> to obtain a series of SNP solutions as shown in Fig. 2a.

**Method 2: synthesis of SNP type II.** A continual growth process using multisteps has been proposed.<sup>22,23</sup> Here, the seed solution in Method 1 was replaced by a 2 mL solution of SNP type I obtained from previous cycles. The other chemicals added are the same as those in Method 1. Furthermore, 50 μL of diluted H<sub>2</sub>O<sub>2</sub> solution (3% w/w) was added in each cycle to etch the self-nucleated spherical nanoparticles.

**Method 3: synthesis of SNP type III.** 2 mL of SNP solution with an average size of 1 μm synthesized using Method 2 with two cycles was added into solution A, followed by 50 μL of diluted H<sub>2</sub>O<sub>2</sub> solution (3% w/w) and 50 μL of 0.5 M NaCl. Finally, 5 mL of solution B was injected dropwise into the mixed solution at a rate of 5 mL h<sup>-1</sup>. The obtained solution was aged for 48 h at room temperature to complete the reaction. To

obtain the huge SNPs, the synthesis step was implemented twice.

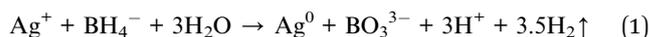
### 2.3 Characteristics

The structural features of the silver nanocrystals were characterized by scanning electron microscope (SEM, Zeiss Ultra Plus) operating at 20 kV under a high vacuum mode. A built-in energy dispersive X-ray spectrometer (EDS) was used to monitor elemental compositions of SNPs. Transmission electron microscopy (TEM) (Fei Tecnai G20) images were obtained with an acceleration voltage of 200 kV. The X-ray diffraction (XRD) spectra were taken using an X-ray diffractometer (Ultima IV) with Cu Kα irradiation (40 kV, 30 mA). The scanning rate was 10° min<sup>-1</sup> and the diffractograms were recorded in the 10–90° region with a 0.02° resolution. X-ray photoelectron spectra (XPS) were performed using a PHI 5000 VersaProbe (Ulvac-phi, Japan) with Al Kα X-rays (1486.6 eV) as the exciting source. Extinction spectra were measured using a fiber optic spectrometer (NOVA, Ideaoptics Technology Ltd. China). The *in situ* excitation, imaging and spectral analysis of SNPs were performed under an Olympus IX-71 optical microscope system.

## 3. Results and discussion

### 3.1 High yield seeds with stacking defects synthesized by the H<sub>2</sub>O<sub>2</sub>-assisted self-screening method

Multi-step aqueous synthesis of SNPs contains two steps.<sup>22–24</sup> The first step is the preparation of silver seed solution using NaBH<sub>4</sub> as a strong reducing agent following eqn (1) where the seeds rapidly grow up from the aggregation of reduced silver atoms under the protection of sodium citrate. In the second step, seeds are added into the SNP growth solution to trigger the reaction where N<sub>2</sub>H<sub>4</sub> is used to reduce Ag<sup>+</sup> and promote the anisotropic deposition of Ag<sup>0</sup> on to the seeds gradually. To increase SNP size, the second step can be repeated several times.<sup>22,23</sup> In this synthesis strategy, a high yield of anisotropic and defect enriched seeds is crucial for the successful growth of the monodispersed plate-like nanostructure.<sup>35</sup> As reviewed by Liz-Marzán,<sup>36</sup> it is a general rule that anisotropic seeds with defects play a critical role in the successful synthesis of high-purity anisotropic crystals. Once isotropic seeds<sup>37</sup> are mixed or randomly generated in the reaction solution, spherical by-products will grow up rapidly and greatly influence the yield and size of the SNPs.<sup>24</sup> Therefore, synthesis methods of high-purity anisotropic seeds are highly required in the research field of metal crystal design and engineering.<sup>36</sup>



Herein we created a self-screening environment to strictly control the type of seeds and obtained SNPs with high purity as shown in Fig. 1. Different from conventional seed synthesis methods, here an appropriate amount of H<sub>2</sub>O<sub>2</sub> and a remarkably excessive dose of NaBH<sub>4</sub> (10 times higher than the conventional dose<sup>24</sup>) are added in the seed synthesis step.

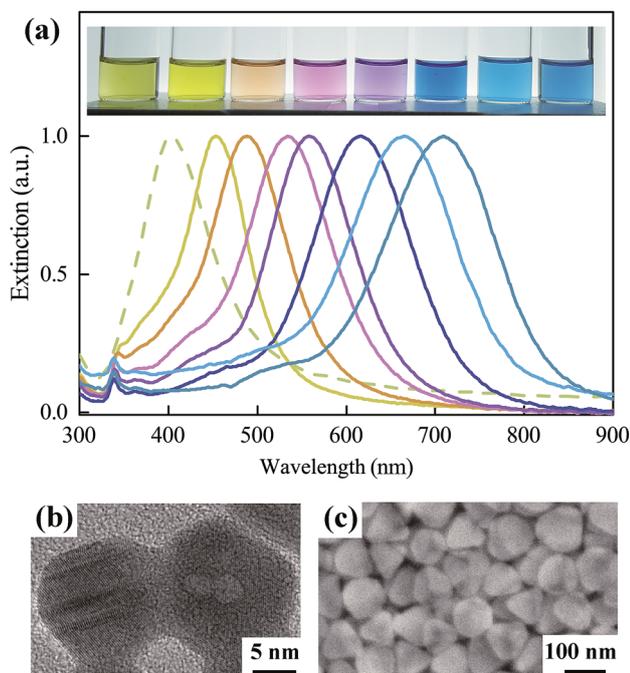


Fig. 2 (a) Photograph of diluted silver solutions and corresponding extinction spectra of silver seeds (dashed line) and SNP type I with different sizes (solid line). (b) HRTEM image of twin-crystal seeds with defects survived in an H<sub>2</sub>O<sub>2</sub> etching environment. (c) SEM image of monodispersed SNP type I with an LSPR band of ~710 nm.



During this nucleation step,  $\text{H}_2\text{O}_2$  acts as an etchant<sup>20,21,37–39</sup> which can selectively etch the isotropic seeds following eqn (2),<sup>39</sup> while the excessive  $\text{NaBH}_4$  reduces the re-generated silver ions soon following eqn (1) again.

During this stage, the 'reduction–etching' process will repeat several times until high-yield anisotropic and defect enriched seeds having excellent etching resistance ability against  $\text{H}_2\text{O}_2$  are obtained. Movie 1 in the ESI† recorded this interesting dynamic reaction process vividly. The color of the reaction solution changed from dark brown to light yellow and back to dark brown several times. When the color changed to brown, the reaction described by eqn (1) occurred. Then, the reaction of  $\text{H}^+$  etching following eqn (2) occurred because the isotropic silver seeds generated can be etched by  $\text{H}_2\text{O}_2$ . In this stage, the color of the solution gradually became lighter. As the amount of  $\text{NaBH}_4$  is remarkably excessive, these two chemical reactions alternatively occurred several times. During this strictly self-screening step, initially generated isotropic seeds were eliminated and the surviving seeds were mainly planar twin-crystals. To show the critical roles of both  $\text{H}_2\text{O}_2$  and  $\text{NaBH}_4$ , we synthesized three types of seed under different conditions as described in the Experimental section. These samples show obviously different crystalline characteristics and morphologies. Although the extinction spectra of seed samples a and c (shown in Fig. 1a) are similar, their crystalline characteristics are different, as shown from the TEM images. Without  $\text{H}_2\text{O}_2$  (seed sample a), isotropic seeds possibly attributing to spherical by-products were easily observed, as shown in Fig. 1c and f. The HRTEM image in Fig. 1f clearly shows that the seeds are isotropic. In comparison, the seeds that survived the strict self-screening process are rich in defects and nano-sized hollow cavities (illustrated by the red arrows) as shown in Fig. 1e and h. The HRTEM image in Fig. 1h shows that the seeds had a planar-twin structure which is beneficial for anisotropic SNP growth. XRD analysis further confirmed this crystalline difference from a macroscopic aspect. The full width at half maximum (FWHM) of the Ag (111) peak (illustrated by the yellow arrows) of seed sample c is about two times wider than that of seed sample a which indicates that the defects and cavities in seed sample c are much more abundant than those in seed sample a, consistent with the results observed in the TEM images. These hollow cavities are unique structural features of the silver seeds synthesized with the addition of  $\text{H}_2\text{O}_2$ . We speculated that they are formed by local area etching of  $\text{H}_2\text{O}_2$ . Meanwhile, two sub-peaks observed (illustrated by the green arrows) in the curve of seed sample c further indicate that the seeds have stacking faults which is critical for SNP growth.<sup>40</sup> Note that the seed-selection process by the addition of  $\text{H}_2\text{O}_2$  that is helpful for the yield improvement of SNPs was first reported by Mirkin's group,<sup>21</sup> improved by Yin's group<sup>37,38</sup> and then extensively used hereafter.<sup>20</sup> However, the seed self-screening phenomenon (Movie 1, ESI†) and crystalline optimization method demonstrated currently have not yet been shown before. Here we found that simply adding  $\text{H}_2\text{O}_2$  is not sufficient to obtain such high-yield anisotropic seeds. As shown in the TEM image of seed sample b (Fig. 1d and g), the typical synthesized structures are mainly silver branches with micrometer sizes when the amount

of  $\text{NaBH}_4$  added is insufficient. The broadening and red-shifting of the absorption band shown in the extinction spectrum of seed sample b further showed their morphological difference. The main reason for achieving such irregular structures is because less  $\text{Ag}^0$  was reduced by  $\text{NaBH}_4$  (which is insufficient) after the  $\text{H}_2\text{O}_2$  etching process. In this condition, self-nucleation and self-screening of the seeds cannot continue. Instead,  $\text{Ag}^0$  slowly adsorbed onto the surface of the seeds already generated and then grew into micro-sized crystals. The optimal amount of  $\text{H}_2\text{O}_2$  was discussed in Fig. S1, ESI.†

The high-resolution transmission electron microscopy (HRTEM) images in Fig. 1h and 2b exhibited the unique anisotropic and defect enriched feature of the seeds that survived after strict  $\text{H}_2\text{O}_2$  etching. Using such seeds, ~100% yield monodispersed SNPs (regarded as SNP type I, the smallest samples with a mean size of 25 nm and a thickness of ~9 nm are shown in Fig. S2†) were obtained, as confirmed by the extinction spectra in Fig. 2a and scanning electron microscopy (SEM) image in Fig. 2c. The extinction spectra in Fig. 2a reflect the monodispersity of the obtained silver solution samples. The dashed line shows the prepared silver seeds with a LSPR band at ~410 nm. In the following step of SNP growth, the red shift of the main LSPR band corresponds to the growth of anisotropic SNPs.<sup>24</sup> Once more silver seeds are added in the reaction solution, the size of the final obtained SNP will be smaller. The yellow solid line in the extinction spectra corresponds to the obtained SNP sample with the smallest size (~25 nm). The SEM images are shown in Fig. S2.† With the decrease of the amount of silver seeds added, the size of the obtained SNPs increased gradually. The main LSPR band of the extinction spectra red shifted from 450 nm to 700 nm, accordingly. Note that, as shown from the extinction spectra of SNP solutions, the usually observed LSPR band at ~420 nm corresponding to spherical by-products<sup>24,41</sup> is completely absent. SNP solutions in Fig. 2a appeared as diverse colors and single LSPR bands corresponding to different particle sizes. It is also observed that pure SNP solution with an LSPR band longer than 600 nm appears blue. Interestingly, a green color which is the usual color of SNP solution<sup>18,19,24</sup> was absent. Actually, the green color is due to a color mixture effect of blue color representing large SNPs and yellow color representing spherical by-products which was difficult to avoid, reflecting their impurity. Fig. S3† further provided a direct comparison between the color difference of these SNP solutions with high-purity and impurity.

Using this seed screening strategy, a long-standing problem of spherical by-products in the aqueous synthesis of SNPs was completely solved. It enables this method to be very competitive for the mass production of high-purity SNPs. As much as 0.32 g of monodispersed SNPs with high-density (0.012 M) were successfully synthesized by one growth cycle (shown in Fig. S4, ESI†) which is over 100-fold larger than that in previous reports.<sup>22–24</sup> The as-prepared monodispersed SNP type I sized from tens to a few hundred nanometers with single LSPR bands ranging from visible to near-infrared range is no doubt a critical building block meeting the requirement of various rapidly developing fields including nanoantenna,<sup>42,43</sup> true color

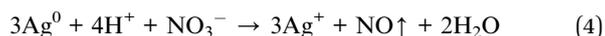


nanoink printing,<sup>44</sup> optical processing,<sup>45–47</sup> electronic packaging<sup>18</sup> and circuit fabrication.<sup>39</sup>

### 3.2 Bottleneck of conventional multistep synthesis of SNPs with high aspect ratio

To obtain SNPs with a higher aspect ratio and controllable thickness, a continual growth process using multisteps has been proposed.<sup>22,23</sup> However, there is a lateral growth limitation which has not been revealed before. Based on this usually used continual growth strategy,<sup>22,23</sup> we used Method 2 (described in Methods in the Experimental section) to explore the size increasing limitation of SNPs. The mean sizes of the SNPs synthesized at different cycles taken from SEM images are plotted in Fig. 3a. The size of the SNPs increased rapidly at the earlier stage. However, it increased gently and finally decreased after 8 cycles. This result indicated that simply increasing the growth cycles cannot obtain SNPs with a very large size. Due to this, the SNPs became unstable after multiple-cycle growth. They suffered etching, cracked from the edges and finally dissolved, rather than continuously grew up. The SEM images shown in Fig. 3b–d reveal this morphological change. At 6 cycles, SNPs with a mean size of 3  $\mu\text{m}$  and smooth surfaces were observed. However, when the cycles further increased, defects and small particles attached on the surface of the SNPs increased. After 9 cycles, defects and cracks enlarged and the size of the SNPs decreased accordingly. Eqn (4) provides a possible explanation about the mechanism of this continual etching phenomenon. It may be attributed to the continuous surface attacking by the accumulated  $\text{H}^+$  generated from the reduction reaction (described by eqn (3)). Once the defects were generated, fresh exposure surfaces without the protection of sodium citrate increased accordingly. This led to an acceleration of the  $\text{H}^+$  etching process from the areas where defects generated. Meanwhile, with a continual size increase, the Ag

atom absorption ability of these large SNPs decreased because of the surface energy decrease. In this stage, the ‘dissolution’ process described by eqn (4) becomes dominant, replacing the ‘growth’ process. More Ag atoms are dissolved by  $\text{H}^+$  which finally leads to a complete dissolution of the SNPs. This etching phenomenon was clearly exhibited by a comparison experiment under a real-time visualization of the reaction process *in situ*, as recorded in Movie 2, ESI†



This  $\text{H}^+$  etching effect during the SNP growth process also leads to the serious size variation of SNPs with the increase of the re-growth cycles, as is clearly shown in Fig. 3a. This is because more and more large SNPs will be etched from the lateral edges. This is a problem that cannot be easily avoided by this synthesis strategy. Although decreasing the re-growth cycles will be beneficial for improving the size variations of SNPs, their size cannot be increased easily to over ten microns.

Meanwhile, as the defects increased gradually during the repeated re-growth steps, small SNPs generated by self-nucleation were apt to be decorated on the surfaces of these large SNPs randomly as shown in Fig. 3c and S5 (ESI†). The maximum size of such SNPs synthesized by 8 growth cycles (regarded as SNP type II) reached  $\sim 8 \mu\text{m}$  which is large enough for the construction of a 2D plasmonic platform.

### 3.3 Breaking the size limitation of SNPs using self-screening

From the analysis above, one can see that the generation of defects on the surface of SNPs is the main factor limiting the size increase of the SNPs. Here we improved the conventional multiple re-growth strategy from the following three aspects: (1) an appropriate amount of  $\text{Cl}^-$  was added to protect the Ag {111} facets of SNPs during the crystal growth process; (2) the reaction time was prolonged from 15–20 min to 2 days per cycle to create a self-screening condition for SNPs; (3) decreasing the repeated regrowth cycles.

The main contribution of  $\text{Cl}^-$  is that it can easily adsorb onto Ag {111} facets<sup>20</sup> to avoid the SNPs being etched by newly generated  $\text{H}^+$  in the reaction. *In situ* observations of the SNP regrowth step with and without the addition of  $\text{Cl}^-$  were compared in Movie 2, ESI† where the effectiveness of  $\text{Cl}^-$  protection was clearly confirmed. The aim of prolonging the reaction time is to enhance the self-screening behaviour of the SNPs in the regrowth step. Once  $\text{Cl}^-$  was added, the SNPs already having defects can be etched by various mild reaction processes, such as  $\text{H}^+$  etching and  $\text{Cl}^-/\text{O}_2$  selective etching of Ag {110} facets of SNPs. Newly generated  $\text{Ag}^+$  was then reduced by  $\text{N}_2\text{H}_4$  again into  $\text{Ag}^0$  following eqn (3). The  $\text{Ag}^0$  selectively deposited onto the lateral sides (Ag {110} facets) of the SNPs without defects very slowly as the surface energy of the large SNPs had decreased accordingly. Those SNPs well protected by  $\text{Cl}^-$  with defect-free surfaces eventually became the ultimate beneficiaries from such a ‘survival of the fittest’ rule, growing

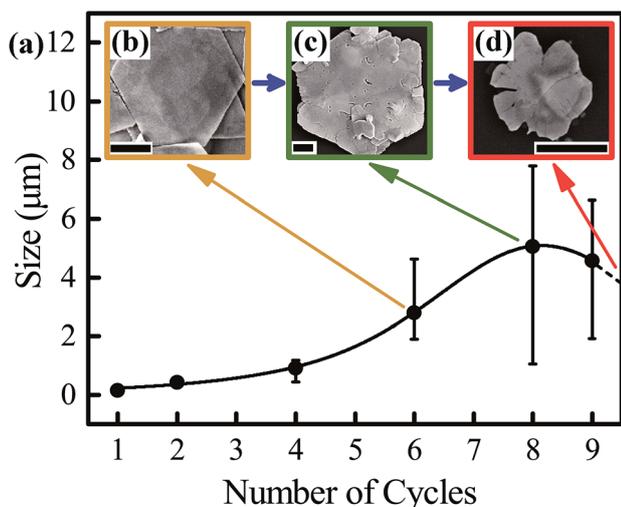


Fig. 3 (a) Mean size of SNPs synthesized by the conventional multiple re-growth process as a function of the number of growth cycles. SEM images in b, c and d show typical morphologies of SNPs synthesized after 6, 8, and 9 cycles, respectively. Scale bar in SEM images is 1  $\mu\text{m}$ .



into astounding huge structures with an ultra-high aspect ratio. In addition, decreasing the number of regrowth cycles is beneficial for the achievement of SNPs with a thinner thickness.

Morphological features of the synthesized huge SNPs, regarded as SNP type III including atomically smooth surfaces and edges, as well as ultrathin thicknesses, are clearly shown by the SEM and TEM images in Fig. 4a–e. The single-crystalline structure with fringe spacing between adjacent silver atoms of 2.5 Å corresponding with {111} was observed as shown in Fig. 4b, similar to that of Fig. S5b.† A serendipitously observed triangular crystallographic cracking inside the SNP shown in Fig. 4e further indicated the regularity of Ag atom arrangement. SEM images in Fig. S6† show that the thickness of SNP type III is between 10–60 nm (mean 30 nm) which is much thinner than that of previous reports.<sup>18–20,22</sup> SNPs with a size larger than 10 μm can be easily observed. A SNP with a size of ~40 μm and an ultra-high aspect ratio of over 1000 : 1 was observed, as shown in Fig. S7.† Note that when 25 nm sized SNPs with a thickness of several nanometers grew up to 40 μm, the size of the crystal increased over 1600-fold. However, their thickness was only increased less than 10-fold. This fully showed the robustness of the current synthesis strategy in thickness confinement. The current achievement with the high-aspect-ratio 2D silver nanostructure with perfect crystal structures and bigger specific surface area will promote the development of a 2D plasmonic platform and high-efficiency photocatalysis applications.<sup>25,29,48,49</sup> It is also a highly expected raw material to fabricate nano-devices with single-crystalline structures.<sup>43,50</sup>

To fully understand the functionality of Cl<sup>−</sup> during the SNP synthesis process, we compared the morphologies of SNP type II (without adding Cl<sup>−</sup>) and SNP type III using HRTEM as shown in Fig. 5a and b. For SNP type II, an organic shell with a width of ~5 nm was observed which was formed by sodium citrate. As discussed above, such an organic shell is not dense enough to protect SNPs against etching. Therefore cracks can be found at the edge of SNP type II in the HRTEM image (Fig. 5a). In comparison, those organic shells were removed from the surface of SNPs once Cl<sup>−</sup> was added by ligand-exchanging, as

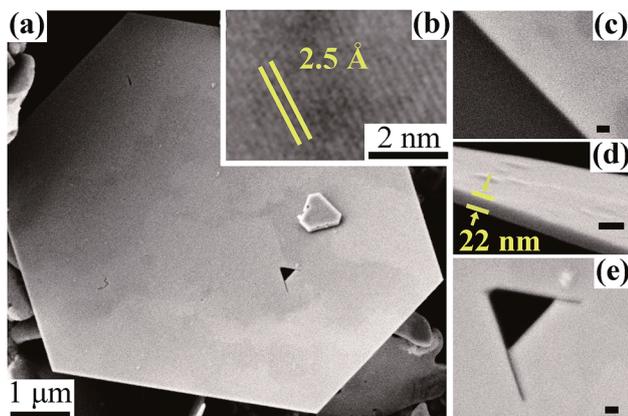


Fig. 4 (a) SEM image of SNP type III. (b) HRTEM image of SNP. (c–e) SEM images of the detailed morphology of the SNP in (a). (Scale bar in (c–e) is 50 nm).

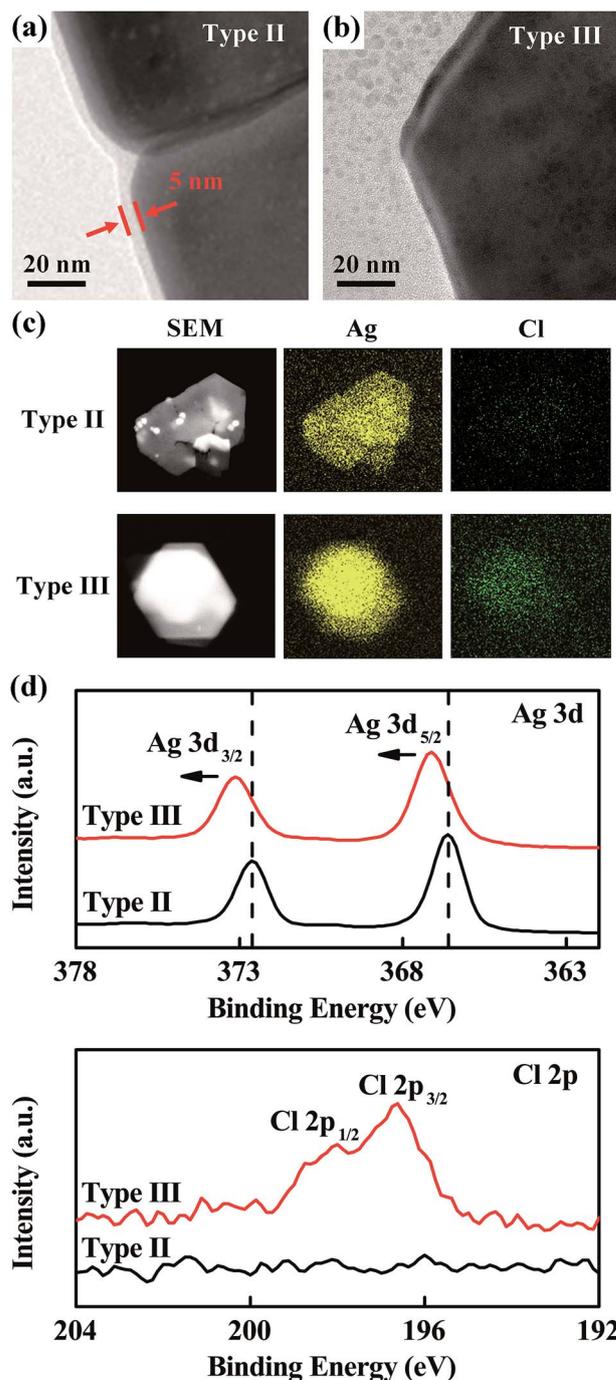


Fig. 5 HRTEM images in (a and b) compared the shells of SNP type II and SNP type III at their edges, respectively. (c) Comparison of elemental distributions taken from EDS mapping. (d) Compared XPS analysis results.

shown in the HRTEM image of SNP type III. During the crystal growth process, a dense passivation layer<sup>20</sup> formed because the adsorption of Cl<sup>−</sup> on the Ag {111} facets of the SNPs provided a better protection of the surfaces of SNPs against H<sup>+</sup> etching. This leads to the successful growth of SNP type III with huge sizes and better crystalline quality. For SNP type III, cracks cannot be found anymore as seen from the HRTEM image in



Fig. 5b. This ligand exchanging mechanism of  $\text{Cl}^-$  to replace organic shells had been investigated by Hui *et al.* using silver nanowires,<sup>51</sup> in accordance with current observation. Compared element mapping between single SNP type II and SNP type III by EDS as shown in Fig. 5c confirmed that Cl adsorbed on the surface of SNP type III. XPS analysis results shown in Fig. 5d confirmed that the passivation layer on the surface of the SNPs mainly constitutes AgCl. This is because the Cl 2p peaks appear in the XPS of SNP type III. Meanwhile, the Ag 3d peaks moved to a higher binding energy which is consistent with the XPS characteristic of AgCl.<sup>52</sup> XPS in the full spectrum range can be seen in Fig. S8, ESI.†

The optimal concentration of  $\text{Cl}^-$  was investigated. Fig. 6 shows the morphologies of the SNPs obtained by adding different amounts of  $\text{Cl}^-$ . Note that to show the intermediate states of the SNPs during the regrowth process, these SNPs experienced only one-circle growth. Without  $\text{Cl}^-$  protection, the SNPs cracked (Fig. 6a). Once  $\text{Cl}^-$  was added, the morphology of the SNPs (Fig. 6b) became different. Some of them already having defects cracked more obviously and would eventually decompose because of etching. Some of them well protected by  $\text{Cl}^-$  showed less cracks and better crystal quality. When the amount of  $\text{Cl}^-$  increased to an optimal value, SNPs with excellent structures were obtained (Fig. 6c). However, when the amount of  $\text{Cl}^-$  was excessive, the SNPs became thicker (Fig. 6d) and the by-products increased obviously (Fig. 6e). This is because an excessive amount of  $\text{Cl}^-$  will also adsorb on the Ag {100} facets and promote the vertical growth of the SNPs, as described in ref. 47.

### 3.4 Thickness dependent 2D SPP modal characteristic of SNPs

Previous reports mainly focused on the demonstration of boundary SPP modes supported by silver–air interfaces of SNPs because the SNPs were thick.<sup>13,14</sup> Here we explained why thinner SNPs are needed for 2D SPP mode transmission and revealed

that thickness plays an important role in determining the modal characteristics of 2D SPP waveguides. The diversity of 2D modes supported by SNPs can be achieved only when their thickness decreases to a sub-100 nm level. For 2D SPP waveguides,<sup>53</sup> once the evanescent fields supported by both surfaces of the SNP couple with each other, the proportion of light intensity distributed inside silver will be greatly suppressed. Such coupled SPP modes suffer much smaller ohmic damping of metal and their propagation lengths increase accordingly. Here the 2D finite element method (FEM) was employed to investigate thickness dependent mode characteristics of SNPs by numerically solving the wave equation.<sup>50</sup> By considering the influence of surrounded dielectrics, we investigated the modal characteristics of 2D plasmonic waveguide configurations where SNPs embedded in asymmetric or symmetric dielectrics. The simulation results in Fig. 7 show that SPP modal characteristics change obviously in both cases when the thickness decreases down to a threshold value (illustrated by the dashed lines in Fig. 7a).

For the asymmetric case where the SNP is placed on the glass substrate ( $n_1 = 1.52$ ) and covered by air ( $n_2 = 1.0$ ), light can be tightly confined at metal/dielectric interfaces. For all the wavelengths calculated, the propagation lengths of the SPP modes greatly increase with the decrease of thickness, especially at the excitation wavelength of 473 nm. Interestingly, the simulated mode intensity distribution shown in Fig. S9 and S10† further exhibited that the capacity of mode confinement of thinner SNPs did not decline obviously, especially for SPP modes with short wavelengths. Therefore this provides an effective method to prolong the propagation length of short-wavelength SPP modes and maintain their mode confinement ability by decreasing the thickness of SNPs. Such ultrathin SNPs can be used as a robust platform for remote excitation and manipulation of subwavelength signals.<sup>9–12,54</sup>

For the symmetric case, the advantage induced by the thickness decrease of SNPs is more obvious as shown in Fig. 7b. This is because thin SNPs surrounded by symmetric dielectric layers support coupling modes where the power ratio

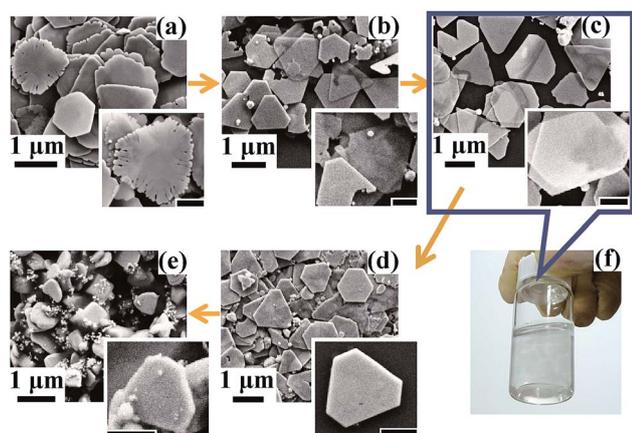


Fig. 6 SEM images represent the morphology difference of SNP type III dependent on  $\text{Cl}^-$  concentration.  $\text{Cl}^-$  added in the synthesis solutions is 0  $\mu\text{L}$ , 20  $\mu\text{L}$ , 50  $\mu\text{L}$ , 100  $\mu\text{L}$  and 250  $\mu\text{L}$ , in (a) to (e), respectively. (f) Shows the photograph of the SNP solution corresponding to (c).

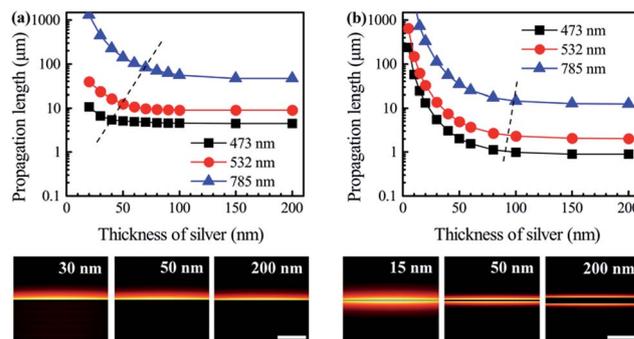


Fig. 7 Numerical simulation results of the relationship between the mode propagation length and SNP thickness for asymmetric (a) and symmetric (b) 2D SNP waveguides at different excitation wavelengths. Insets in (a) and (b) illustrate the cross-sectional view of the mode distribution at  $\lambda = 532$  nm with different thicknesses (scale bar is 500 nm).



distributed in metal dramatically decreases (discussed in Fig. S9 and S10, ESI†). Therefore, thin SNPs are also favourable for the construction of long-range SPP waveguides and devices.<sup>53</sup> In addition, as the mode intensity penetrates deeper into the dielectric layers for ultrathin SNPs (illustrated in insets of Fig. 7), such thin SNPs can serve as platforms with plasmon enhanced light-matter interactions, especially suitable for nonlinear optical applications.<sup>55</sup>

### 3.5 Optical characterization of SNP based 2D plasmonic platform

On the basis of the breakthrough in the synthesis of SNPs with a thinner thickness and much larger size, we explored their diverse optical modal and spectral characteristics experimentally using *in situ* optical microscope imaging together with a scattering spectrum measurement system taken from a micro-domain (illustrated in Fig. S11, ESI†). Washed SNPs were placed onto a glass substrate ( $n_1 = 1.52$ ) and surrounded by air ( $n_2 = 1.0$ ), regarded as an asymmetric 2D SPP waveguide. Focused laser beams with different wavelengths were used as the excitation source.

Here we showed that SNP type II and SNP type III with different morphological features are both suitable for the construction of a 2D plasmonic platform, however, with distinguishing features. SNP type III having smooth surfaces supports the 2D planar propagation SPP mode which uniformly scattered at the output edges, as shown in Fig. 8. As observed from the optical excitation images in Fig. 8a and the SEM image in Fig. 8b, one can see that excited SPP mode with wavelength  $\lambda = 473$  nm propagated more than  $10 \mu\text{m}$  and uniformly coupled to optical signals at the other edge. The scattered light appeared as a 1D linear intensity distribution which can carry deep subwavelength information. Numerical simulation employing 3D FEM modeling<sup>56</sup> (Fig. 8c) exhibited the mode intensity distribution of such SPP modes from different cross-section views. A greatly enhanced light scattering effect can be observed at the edges of the SNP (indicated by the blue arrows)

where the profile area of the scattered output signals reached  $\sim 1/50 \lambda$  orthogonal to the direction of the output edge. This indicated that light information enhanced by plasmons with resolution breaking of the optical diffraction limitation can be effectively coupled into free-space by edge scattering of SNPs. Optical spectra of the scattered signals at the output positions were taken using *in situ* micro-domain measurement modules designed by Ideaoptics Technology Ltd. China, with a high resolution as discussed in Fig. S12 and S13.†

For applications such as *in situ* photocatalysis and optical sensing at the nanoscale,<sup>9,10</sup> the magnitude of scattering intensity needs to be controlled. Previous works demonstrated a tuning strategy based on changing the polarization of incident light. However, the electromagnetic field distribution of the modes was also changed accordingly.<sup>13</sup> Here we present an alternative strategy to control the excitation of SPP modes without obviously changing their electromagnetic field distribution. A beam splitter prism (BSP) was added into the excitation light path as illustrated in Fig. S11, ESI.† Here  $\theta$  corresponds to the angle between the incident light beam and the reference axis without the BSP. Once the BSP was added in the light path,  $\theta$  can be continuously changed by rotating the BSP. We defined a parameter  $\delta\theta$  to describe the direction change of incident light. As shown in Fig. 8a and d, the scattering signals can be controlled by simply rotating the angle of the BSP. The inset in Fig. 8d recorded the relationship between the scattered intensity and the relative rotation angle  $\delta\theta$  at  $\lambda = 473$  nm. The 'on' and 'off' states of scattering signals can be switched at  $\sim 12^\circ$ . Besides, the BSP can also be used to enhance the imaging resolution as discussed in Fig. S12.† Excitation experiments using  $\lambda = 532$  nm and 785 nm are further shown in Fig. S13, ESI.† The realization of SNP type III based 2D plasmonic platforms in a wide wavelength range provides a possibility for transmitting and scattering subwavelength information of uniformly distributed molecules<sup>57</sup> or large area covered 2D materials<sup>58</sup> attached on the surface of SNPs with high resolution.

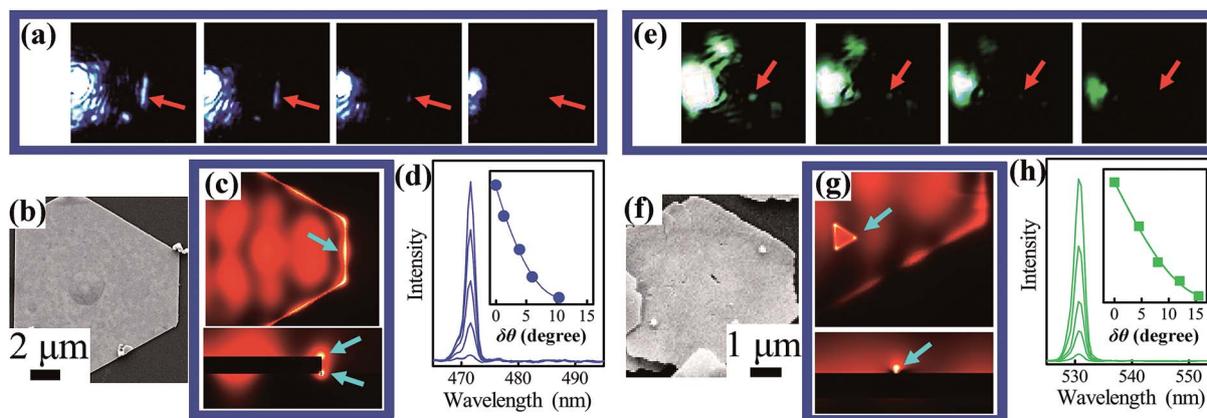


Fig. 8 (a) Excitation of SPP modes using SNP type III with different  $\delta\theta$  of BSP. (b) SEM image of SNP type III used for SPP excitation. (c) Top and lateral view of the light intensity distributions of SNP type III taken from 3D simulation. (d) Light scattering spectra taken from the output edge illustrated by the red arrow in (a) as a function of  $\delta\theta$ . The inset in (d) shows light intensity variation as a function of  $\delta\theta$ . Analogous results for SNP type II are shown in (e–h).



To further extend the manipulability and enhance the light confinement volume of SPP waveguides, subwavelength scattering centre fabricated by top-down technologies such as focused ion beam was demonstrated.<sup>13,14</sup> Here we show that the chemically synthesized nanostructure without post-processing (shown in Fig. 8e–h) can also serve as a similar platform with an excellent light scattering enhancement ability. As described above, the morphological feature of SNP type II is unique as there are randomly distributed nanoparticles and nanogaps decorated on the surfaces. Such defects serving as 0D nano antennas greatly improved the light coupling efficiencies and promoted significant light scattering enhancement at nano-scale volumes, known as ‘hot spots’.<sup>59,60</sup> The optical images in Fig. 8e exhibit the switchable 0D hot spots with tunable scattering intensity at  $\lambda = 532$  nm. Fig. 8g shows the simulation result describing this deep subwavelength scattering enhancement effect which is induced by the attached small nanoparticles. The intensities of the scattered signal from those spots are also adjustable utilizing the BSP. Micro-domain optical spectrum measurements of the single hot spot in Fig. 8h monitored the intensity response according to the variation of  $\delta\theta$ . Such SNPs with plenty of hot spots are excellent candidates for the development of single molecular detection and super-resolution bio-imaging.<sup>61</sup>

### 3.6 Simultaneous multi-wavelength transmission at subwavelength range

Using the 2D plasmonic platform, SPP modes with multiple wavelengths can either propagate along the smooth surface of the SNPs or be guided and scatter into a single hot spot by the antenna effect as shown in Fig. S14 (ESI<sup>†</sup>) and 9. A super-continuum laser source (SC-PRO, 6W) with an acousto-optic tunable filter module (AOTF-Pro, Yangtze Soton Laser Co., Ltd., China) was used to generate collinear laser beams with variable wavelengths. The central peak of the laser beam with a FWHM of  $\sim 10$  nm can be tuned ranging from 450 nm to 700 nm by the acousto-optic effect. The optical images in Fig. 9 demonstrate that SPP modes can be excited by laser beams with either a single wavelength or mixed wavelengths together in a broadband range. Fig. 9h showed that light beams with seven

mixed wavelengths together have been successfully converged into a single hot spot with high spatial resolution. The measured scattering spectrum of the hot spot is shown in Fig. 9i. The successful realization of such directional excitation of a white color hot spot is due to the effective removal of the chromatic aberration. By setting the angle of the BSP to an appropriate value, the dispersion error of laser beams with multiple wavelengths was eliminated. Meanwhile, acousto-optic tunable filters can be easily used to control the excitation intensity of laser beams with different wavelengths, separately. Therefore, an arbitrary combination of multiple SPP modes can be guided into a deep subwavelength volume easily which provides an advanced platform for nonlinear optical effect study. More details about the multiple wavelength excitation and characterization are discussed in Fig. S14.<sup>†</sup>

### 3.7 Remote excitation of subwavelength scattering light source

Directional controlling of far-field emission from a subwavelength emitter is a highly expected function of plasmonic platforms which is useful in molecular sensing and sub-diffraction limit light source integration.<sup>11</sup> In Fig. 10 we demonstrated the remote excitation and switching of a subwavelength light source using quantum dot (QD) adsorbed 2D SNPs. To prepare the sample, 5 mL SNP type III solution was centrifuged twice and dropped on a glass substrate. Then a mixture of polystyrene/CdS@ZnS QDs film was spin coated on the substrate and then dried. Fig. 10b shows that focused laser beam with  $\lambda = 532$  nm successfully excited the 2D SPP mode which propagated along the metal–dielectric interfaces of the SNP and eventually scattered into free space at the other edge. In this circumstance, as QDs that adsorbed on the surface of the SNP were effectively stimulated by the guided SPP modes, photoluminescence of QDs was observed at the output scattering edges, as shown in Fig. 10d. It was observed that the plasmon enhanced subwavelength light emitter presented a linear intensity distribution along the edge of the SNP and scattered in front. Meanwhile, it can be either turned on or turned off remotely by simply changing  $\delta\theta$  of the BSP, as compared in Fig. 10c and d.

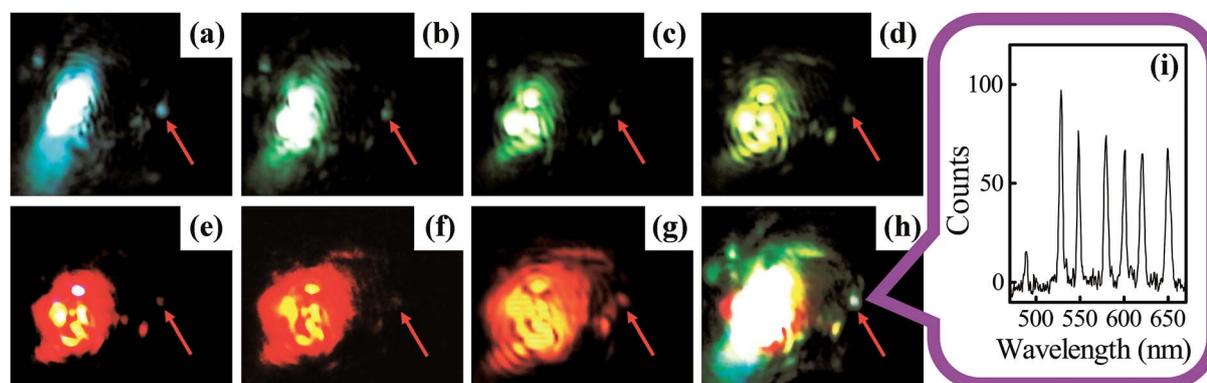
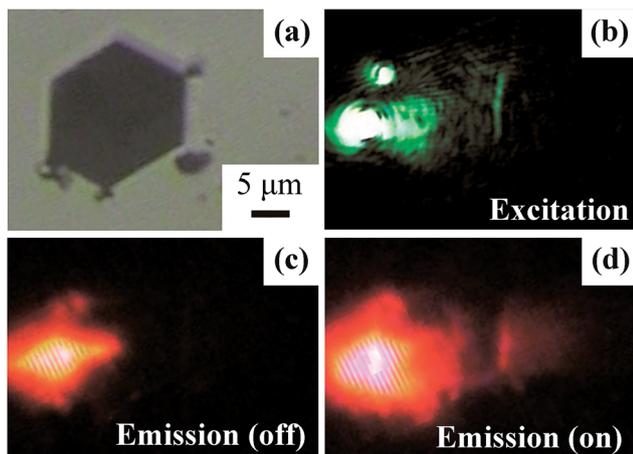


Fig. 9 Excitation of SNP type II using a focused laser beam with wavelengths ranging from 490 nm (a) to 530 nm (b), 550 nm (c), 580 nm (d), 600 nm (e), 620 nm (f), and 650 nm (g), and their mixture in (h). (i) Scattering spectrum taken from a single white color spot in (h).





**Fig. 10** Remote excitation and switching of QDs by a guided 2D SPP mode supported by SNP type III. (a) Optical image of SNP covered with quantum dots/polystyrene mixture. (b) Excitation by focused laser spot with  $\lambda = 532$  nm. (c and d) show the photoluminescence images taken from the 'off' and 'on' states by tuning the rotation angle of the BSP. Here a long-pass filter was used to cut off the excitation light.

## 4. Conclusions

In summary, the growth kinetics of 2D anisotropic SNP synthesis were comprehensively studied with a series of comparison experiments assisted by real-time reaction monitoring under optical microscope observation. The important roles of seed type screening and  $H^+$  etching behaviours during the crystal growth process were clarified. Based on these mechanism elucidations, we successfully realized the fabrication of huge sized single-crystalline SNPs with atomically flat surfaces, broke the size limitation of ultrathin SNPs (with an aspect ratio over 1000 : 1), and also obtained SNPs with various morphological features suitable for nanophotonic applications. The proposed chemical synthesis strategy with strengthened self-screening and self-assembly behaviours provides a promising nanofabrication technology with low cost and short time consumption alternative to the lithographic process for the building of preconceived crystal structures. An optical characteristics investigation examined that a SNP based 2D platform can effectively manipulate light with plasmons in subwavelength volumes with a high degree of freedom. Especially, when the thickness of the SNPs was decreased, the mode propagation properties of 2D SPP modes in a broadband wavelength range were improved accordingly. Besides, particular functionalities of 1D and 0D light scattering enhancement with controllable intensities, as well as remote excitation of gain medium, make such 2D plasmonic platforms advanced tools for optical processing, remote molecular sensing, nano-imaging and active nanophotonic device integration applications.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

This work is supported by MOST under Grant Number 2017YFA0205800, NSFC under grant numbers 61307066, 11734005 and 61450110442, the Natural Science Foundation of Jiangsu Province under grant number BK20130630, the Doctoral Fund of Ministry of Education of China under grant number 20130092120024, and the Innovation fund of School of Electronic Science and Engineering, Southeast University under grant number 2242015KD006.

## Notes and references

- M. Rycenga, C. M. Cobley, J. Zeng, W. Li, C. H. Moran, Q. Zhang, D. Qin and Y. N. Xia, *Chem. Rev.*, 2011, **111**, 3669–3712.
- X. Guo, Y. G. Ma, Y. P. Wang and L. M. Tong, *Laser Photonics Rev.*, 2013, **7**, 855–881.
- H. Duan, H. Hu, K. Kumar, Z. Shen and J. K. W. Yang, *ACS Nano*, 2011, **5**, 7593–7600.
- H. Ditlbacher, A. Hohenau, D. Wagner, U. Kreibig, M. Rogers and F. Hofer, *Phys. Rev. Lett.*, 2005, **95**, 257403.
- P. Kusar, C. Gruber, A. Hohenau and J. R. Krenn, *Nano Lett.*, 2012, **12**, 661–665.
- J. Takahara, S. Yamagishi, H. Taki, A. Morimoto and T. Kobayashi, *Opt. Lett.*, 1997, **22**, 475–477.
- H. Wei, S. Zhang, X. Tian and H. Xu, *Proc. Natl. Acad. Sci. U. S. A.*, 2013, **110**, 4494–4499.
- H. Wei, Z. P. Li, X. R. Tian, Z. X. Wang, F. Z. Cong, N. Liu, S. P. Zhang, P. Nordlander, N. J. Halas and H. X. Xu, *Nano Lett.*, 2011, **11**, 471–475.
- Y. Fang, H. Wei, F. Hao, P. Nordlander and H. X. Xu, *Nano Lett.*, 2009, **9**, 2049–2053.
- Y. Z. Huang, Y. R. Fang, Z. L. Zhang, L. Zhu and M. T. Sun, *Light: Sci. Appl.*, 2014, **3**, e199.
- H. S. Lee, D. H. Luong, M. S. Kim, Y. Jin, H. Kim, S. Yun and Y. H. Lee, *Nat. Commun.*, 2016, **7**, 13663.
- C. Yang, D. Pan, L. Tong and H. X. Xu, *Nanoscale*, 2016, **8**, 19195–19199.
- H. Lee, K.-Y. Jeong, T. Kang, M.-K. Seo and B. Kim, *Nanoscale*, 2014, **6**, 514–520.
- C. W. Chang, F. C. Lin, C. Y. Chiu, C. Y. Su, J. S. Huang, T. P. Perng and T. J. Yen, *ACS Appl. Mater. Interfaces*, 2014, **6**, 11791–11798.
- X. Cai and A. Zhai, *Rare Met.*, 2010, **29**, 407–412.
- S. Kumar, Y. W. Lu, A. Huck and U. L. Andersen, *Opt. Express*, 2012, **20**, 24614–24622.
- X. Liu, L. Li, Y. Yang, Y. Yin and C. Gao, *Nanoscale*, 2014, **6**, 4513–4516.
- R. Z. Li, A. Hu, D. Bridges, T. Zhang, K. D. Oakes, R. Peng, U. Tumuluri, Z. L. Wu and Z. L. Feng, *Nanoscale*, 2015, **7**, 7368–7377.
- H. M. Ren, Y. Guo, S. Y. Huang, K. Zhang, M. M. Yuen, X. Z. Fu, S. H. Yu, R. Sun and C. P. Wong, *ACS Appl. Mater. Interfaces*, 2015, **7**, 13685–13692.
- H. Gatemala, P. Pienpinijtham, C. Thammacharoen and S. Ekgasit, *CrystEngComm*, 2015, **17**, 5530–5537.



- 21 G. S. Métraux and C. A. Mirkin, *Adv. Mater.*, 2005, **17**, 412–415.
- 22 J. Zeng, X. Xia, M. Rycenga, P. Henneghan, Q. Li and Y. N. Xia, *Angew. Chem., Int. Ed.*, 2011, **50**, 244–249.
- 23 Q. Zhang, Y. Hu, S. Guo, J. Goebel and Y. Yin, *Nano Lett.*, 2010, **10**, 5037–5042.
- 24 X. Y. Zhang, A. Hu, T. Zhang, W. Lei, X. J. Xue, Y. Zhou and W. W. Duley, *ACS Nano*, 2011, **5**, 9082–9092.
- 25 D. Erickson, X. Serey, Y. F. Chen and S. Mandal, *Lab Chip*, 2011, **11**, 995–1009.
- 26 H. Butt, A. K. Yetisen, R. Ahmed, S. H. Yun and Q. Dai, *Appl. Phys. Lett.*, 2015, **106**, 121108.
- 27 R. Ahmed, A. K. Yetisen and H. Butt, *ACS Nano*, 2017, **11**, 3155–3165.
- 28 J. Butet, P.-F. Brevet and O. J. F. Martin, *ACS Nano*, 2015, **9**, 10545–10562.
- 29 D. Wolf, T. Schumacher and M. Lippitz, *Nat. Commun.*, 2016, **7**, 10361.
- 30 Y. Salamin, W. Heni, C. Haffner, Y. Fedoryshyn, C. Hoessbacher, R. Bonjour and L. R. Dalton, *Nano Lett.*, 2015, **15**, 8342–8346.
- 31 A. Emboras, J. Niegemann, P. Ma, C. Haffner, A. Pedersen, M. Luisier, C. Hafner, T. Schimmel and J. Leuthold, *Nano Lett.*, 2016, **16**, 709–714.
- 32 K. J. Russell, T. L. Liu, S. Cui and E. L. Hu, *Nat. Photonics*, 2012, **6**, 459–462.
- 33 S. J. Kress, F. V. Antolinez, P. Richner, S. V. Jayanti, D. K. Kim, F. Prins, A. Riedinger, M. P. C. Fischer, S. Meyer, K. M. McPeak and D. Poulidakos, *Nano Lett.*, 2015, **15**, 6267–6275.
- 34 W. Zhu, R. Esteban, A. G. Borisov, J. J. Baumberg, P. Nordlander, H. J. Lezec, J. Aizpurua and K. B. Crozier, *Nat. Commun.*, 2016, **7**, 11495.
- 35 W. Niu, L. Zhang and G. Xu, *Nanoscale*, 2013, **5**, 3172–3181.
- 36 L. M. Liz-Marzán and M. Grzelczak, *Science*, 2017, **356**, 1120–1121.
- 37 Q. Zhang, N. Li, J. Goebel, Z. Lu and Y. Yin, *J. Am. Chem. Soc.*, 2011, **133**, 18931–18939.
- 38 N. Li, Q. Zhang, S. Quinlivan, J. Goebel, Y. Gan and Y. Yin, *ChemPhysChem*, 2012, **13**, 2526–2530.
- 39 X. Y. Zhang, J. J. Xu, J. Y. Wu, F. Shan, X. D. Ma, Y. Z. Chen and T. Zhang, *RSC Adv.*, 2017, **7**, 8–19.
- 40 T. C. Rocha and D. Zanchet, *J. Phys. Chem. C*, 2007, **111**, 6989–6993.
- 41 N. Bastús, F. Merkoçi, J. Piella and V. Puntes, *Chem. Mater.*, 2014, **26**, 2836–2846.
- 42 R. A. Jensen, I.-C. Huang, O. Chen, J. T. Choy, T. S. Bischof, M. Lončar and M. G. Bawendi, *ACS Photonics*, 2016, **3**, 423–427.
- 43 R. Méjard, A. Verdy, O. Demichel, M. Petit, L. Markey, F. Herbst, R. Chassagnon, G. Colas des Francs, B. Cluzel and A. Bouhelier, *Opt. Mater. Express*, 2017, **7**, 1157–1168.
- 44 X. Zhu, C. Vannahme, E. Højlund-Nielsen, N. A. Mortensen and A. Kristensen, *Nat. Nanotechnol.*, 2015, **11**, 325–329.
- 45 Y. Montelongo, A. K. Yetisen, H. Butt and S. H. Yun, *Nat. Commun.*, 2016, **7**, 12002.
- 46 H. J. Yang, S. Y. He, H. L. Chen and H. Y. Tuan, *Chem. Mater.*, 2014, **26**, 1785–1793.
- 47 M. H. Kim, S. K. Kwak, S. H. Im, J.-B. Lee, K.-Y. Choi and D.-J. Byun, *J. Mater. Chem. C*, 2014, **2**, 6165–6170.
- 48 Y. Yao, R. Shankar, M. A. Kats, Y. Song, J. Kong, M. Loncar and F. Capasso, *Nano Lett.*, 2014, **14**, 6526–6532.
- 49 Y. Liu, J. Lou, M. Ni, C. Song, J. Wu, N. P. Dasgupta, P. Tao, W. Shang and T. Deng, *ACS Appl. Mater. Interfaces*, 2016, **8**, 772–779.
- 50 X. Y. Zhang, A. Hu, J. Z. Wen, T. Zhang, X. J. Xue, Y. Zhou and W. W. Duley, *Opt. Express*, 2010, **18**, 18945–18959.
- 51 Z. Hui, Y. Liu, W. Guo, L. Li, N. Mu, C. Jin, Y. Zhu and P. Peng, *Nanotechnology*, 2017, **28**, 285703.
- 52 A. Zhang, L. Zhang, Q. Zhu, B. Dai, W. Sheng, S. Su and J. Xiang, *Fuel Sci. Technol.*, 2017, **159**, 222–231.
- 53 P. Berini, Long-Range Surface Plasmon Polaritons, *Adv. Opt. Photonics*, 2009, **1**, 484–588.
- 54 H. Xin, Y. Li, D. Xu, Y. Zhang, C. H. Chen and B. J. Li, *Small*, 2017, **13**, 1–10.
- 55 Y. Wu, X. Dai, Y. Xiang and D. Fan, *J. Appl. Phys.*, 2017, **121**, 103103.
- 56 X. Y. Zhang, T. Zhang, A. Hu, Y. J. Song and W. W. Duley, *Appl. Phys. Lett.*, 2012, **101**, 153118.
- 57 F. Benz, M. K. Schmidt, A. Dreismann, R. Chikkaraddy, Y. Zhang, A. Demetriadou, C. Carnegie, H. Ohadi, B. de Nijs, R. Esteban, J. Aizpurua and J. Aizpurua, *Science*, 2016, **354**, 726–729.
- 58 H. Butt, P. R. Kidambi, B. Dlubak, Y. Montelongo, A. Palani, G. A. Amaratunga, S. Hofmann and T. D. Wilkinson, *Adv. Opt. Mater.*, 2013, **1**, 869–874.
- 59 P. Peng, H. Huang, A. M. Hu, A. P. Gerlich and Y. N. Zhou, *J. Mater. Chem.*, 2012, **22**, 15495–15499.
- 60 R. Z. Li, A. Hu, T. Zhang and K. D. Oakes, *ACS Appl. Mater. Interfaces*, 2014, **6**, 21721–21729.
- 61 J. Y. Wu, X. Y. Zhang, X. D. Ma, Y. P. Qiu and T. Zhang, *RSC Adv.*, 2015, **5**, 95178–95182.

