

Cite this: *Chem. Sci.*, 2022, 13, 8355

All publication charges for this article have been paid for by the Royal Society of Chemistry

# An atomically precise silver nanocluster for artificial light-harvesting system through supramolecular functionalization†

Anish Kumar Das,<sup>‡a</sup> Sourav Biswas,<sup>‡a</sup> Surya Sekhar Manna,<sup>Ⓜb</sup> Biswarup Pathak<sup>Ⓜb</sup> and Sukhendu Mandal<sup>Ⓜ\*a</sup>

Designing an artificial light-harvesting system (LHS) with high energy transfer efficiency has been a challenging task. Herein, we report an atom-precise silver nanocluster (Ag NC) as a unique platform to fabricate the artificial LHS. A facile one-pot synthesis of  $[\text{Cl}@\text{Ag}_{16}\text{S}(\text{S-Adm})_8(\text{CF}_3\text{COO})_5(\text{DMF})_3(\text{H}_2\text{O})_2] \cdot \text{DMF}$  ( $\text{Ag}_{16}$ ) NC by using a bulky adamantanethiolate ligand is portrayed here which, in turn, alleviates the issues related to the smaller NC core designed from a highly steric environment. The surface molecular motion of this NC extends the non-radiative relaxation rate which is strategically restricted by a recognition site-specific supramolecular adduct with  $\beta$ -cyclodextrin ( $\beta$ -CD) that results in the generation of a blue emission. This emission property is further controlled by the number of attached  $\beta$ -CD which eventually imposes more rigidity. The higher emission quantum yield and the larger emission lifetime relative to the lesser numbered  $\beta$ -CD conjugation signify  $\text{Ag}_{16} \cap \beta\text{-CD}_2$  as a good LHS donor component. In the presence of an organic dye ( $\beta$ -carotene) as an energy acceptor, an LHS is fabricated here *via* the Förster resonance energy transfer pathway. The opposite charges on the surfaces and the matched electronic energy distribution result in a 93% energy transfer efficiency with a great antenna effect from the UV-to-visible region. Finally, the harvested energy is utilized successfully for efficient photocurrent generation with much-enhanced yields compared to the individual components. This fundamental investigation into highly-efficient energy transfer through atom-precise NC-based systems will inspire additional opportunities for designing new LHSs in the near future.

Received 18th May 2022  
Accepted 20th June 2022

DOI: 10.1039/d2sc02786k

rsc.li/chemical-science

## Introduction

Light-harvesting systems (LHSs) have a crucial role in natural photosynthesis for transforming photonic energy into chemical energy, that covers an outlook for the utilization of renewable energy sources.<sup>1,2</sup> In natural photosynthesis, the system-specific organization of chlorophylls advances strong photon absorptivity and constituent emission energy; simultaneously conquering energy-dissipating excited-states decay processes. These unique features permit excited-states energy transfer to the next neighbour of LHS and, ultimately, to the reaction centre with impressive efficiency.<sup>3-6</sup> For the fabrication of artificial LHSs, two important prerequisites are crucial for efficient energy transfer between the donor-acceptor through the Förster

Resonance Energy Transfer (FRET) pathway. Firstly, the packing of donors should not be dense and intramolecular self-quenching should be avoided.<sup>7-11</sup> Secondly, the multiple donors should correspond to one acceptor. In a natural LHS, proteins and chlorophylls interacted through non-covalent interaction. Several artificial LHSs were evolved to mimic the outstanding structure and functionality of the natural photosynthesis system. However, the efficient charge transfer property of such a system is the main concern for practical applicability.<sup>12,13</sup> To solve the issues of traditional chromophores, an encouraging strategy depends on host-guest interaction to keep up miniature rift chromophores during the suppression of exciting energy quenching.<sup>1,2</sup> The use of large hosts like cucurbit[*n*]uril and pillar[*n*]arene have the advantage of boosting the fluorescence intensity through encapsulation of the guest group of the chromophores while impeding chromophores from stacking.<sup>14-19</sup> So, it is clear that the design of the donor molecules is the most essential factor for determining the activity of the artificial LHS. In this current circumstance, to continue the development of the artificial LHSs with the efficient FRET process, atom-precise metal NCs would be the appropriate alternative candidate.

<sup>a</sup>School of Chemistry, Indian Institute of Science Education and Research Thiruvananthapuram, Kerala 69551, India. E-mail: [sukhendu@iisertvm.ac.in](mailto:sukhendu@iisertvm.ac.in)

<sup>b</sup>Department of Chemistry, Indian Institute of Technology, Indore, Madhya Pradesh 453552, India

† Electronic supplementary information (ESI) available. CCDC 2169435. For ESI and crystallographic data in CIF or other electronic format see <https://doi.org/10.1039/d2sc02786k>

‡ Shows the equal contribution.



Ligand-stabilized silver nanoclusters (Ag NCs) have gained tremendous attention due to their fascinating structural architectures with molecular precision and exotic photophysical properties.<sup>20–28</sup> Many recent studies have identified that the geometry, size, surface ligands, inorganic–organic interfaces, and chemical functionalization implement an important role in controlling their distinct physical and chemical properties.<sup>20,21,29–31</sup> Although the high-nuclear Ag NCs related to the core–shell arrangement are well studied, the anion templated mono-layered core containing the  $d^{10}$  configuration of  $\text{Ag}(I)$  ions is relatively less researched. It has been reported that templating species were utilized to control the formation of the  $\text{Ag}(I)$  core based on electrostatic interactions that regulate the size and geometry of the core.<sup>32–34</sup> However, most of the reported anion template silver NCs rarely exhibit photoluminescence (PL) properties at room temperature and it is the most perplexing issue which needs to be resolved.<sup>35–38</sup> Progressing research activities suggest that imposing bulkiness on the surface protecting groups can restrict the intramolecular motion of the NC in solution that can directly decrease the non-radiative relaxation pathway and enhance the emission properties.<sup>39–48</sup> Therefore, it will be very interesting to encapsulate the ligand-protected Ag NC with a bulky host to build the host–guest adduct at the recognition site of the NC to improve the emission properties by introducing surface rigidity through bulkiness to counteract the thermally activated non-radiative relaxation which will be beneficial for the artificial LHS fabrication.

Herein, we have designed an atom-precise Ag NC-based novel artificial LHS with a very high energy transfer efficiency. We have strategically synthesized bulky adamantanethiolate ligand-protected,  $[\text{Cl}@\text{Ag}_{16}\text{S}(\text{S-Adm})_8(\text{CF}_3\text{COO})_5(\text{DMF})_3(\text{H}_2\text{O})_2]\cdot\text{DMF}$ ,  $\text{Ag}_{16}$  NC. Although a bulky ligand system is utilized to restrict the surface molecular vibrations in solution, it is not sufficient for generating the emission properties of this as-synthesized NC at room temperature. To alleviate this issue, we have applied a host–guest approach to rigidify the outer surface of the cluster by anchoring a specific hydrophobic guest molecule  $\beta$ -cyclodextrin ( $\beta$ -CD) to alter the relaxation mechanism. The successful entrapment of the adamantane moieties inside the cavity of the  $\beta$ -CD is fully characterized by various spectroscopic tools and confirms the number of the supramolecular adduct,  $\text{Ag}_{16} \cap \beta\text{-CD}_n$  ( $n = 1, 2$ ), while the facile position is identified by the binding energy calculation. We have then utilized this system for generating blue emission energy to construct an LHS *via* the FRET process with an organic dye  $\beta$ -carotene. A 93% energy transfer efficiency and good antenna effect from the UV-to-visible region are detected, which is further utilized for efficient photocurrent generation.

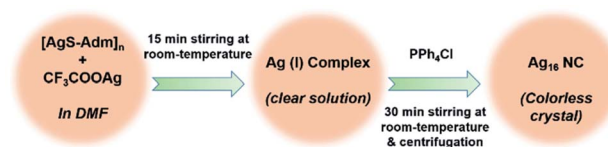
## Results and discussion

To begin our investigation, a facile one-pot chemical reaction pathway was followed to synthesize  $\text{Ag}_{16}$  NC. At first, the reaction took place between the  $[\text{AgS-Adm}]_n$  complex and  $\text{CF}_3\text{-COOAg}$  in dimethylformamide (DMF) solvent (Scheme 1). Afterward,  $\text{PPh}_4\text{Cl}$  was added into the mixture as a source of

a templating agent of chloride ion. The final mixture was kept in dark for 3 days and transparent colorless block-shaped crystals were acquired, yielding 58% (Ag-based). In general, synthesizing smaller NCs by using the bulky thiolate groups is a challenging task as the bulkiness of the ligand provides a higher oxidation resistance to larger Ag NCs.<sup>49</sup> In this report, we successfully achieve the smaller core-sized Ag NC utilizing the bulky adamantanethiolate ligand which will surely throw light on the synthesis procedure with the controlled size of the NCs.

The obtained high-quality colorless crystals are suitable to elucidate the structural information exclusively *via* the single-crystal X-ray diffraction (SCXRD) technique. The block-shaped crystals are crystallized in a triclinic crystal system with space group  $P1$  (no. 2) (Table S1†). The crystallographic data implies that the overall molecular formula of the as-synthesized NC is  $[\text{Cl}@\text{Ag}_{16}\text{S}(\text{S-Adm})_8(\text{CF}_3\text{COO})_5(\text{DMF})_3(\text{H}_2\text{O})_2]\cdot\text{DMF}$ , ( $\text{Ag}_{16}$ ), depicted in Fig. 1 as top and side views. The chemical formula also reveals the neutral state of the  $\text{Ag}_{16}$  NC where all the Ag atoms are in the +1 state.

In the view of the structural aspects, the skeleton of the  $\text{Ag}_{16}$  core is constructed through metallophilic interactions among  $\text{Ag}(I)$  ions and adopts a geometry of square gyrobicupola ( $J_{29}$ ; Johnson solid) with a slight distortion (Fig. 2a). The crystallographic data reveals that the bond distances of  $\text{Ag}^I\cdots\text{Ag}^I$  in the  $\text{Ag}_{16}$  core fall in the range of 2.95–3.39 Å, which are lower compared to twice the van der Waals radius of Ag (3.44 Å).<sup>50,51</sup> The average distance indicates that some sort of long-range non-covalent interaction (NCI) is present between the two Ag atoms which must be an argentophilic interaction.<sup>52</sup> We have calculated the weak metallophilic interaction such as bond critical point (BCP), bond order, and NCI of two adjacent Ag atoms of the  $\text{Ag}_{16}$  core. The electron density [ $\nabla^2\rho(r_c) = 0.055$ ] between two Ag atoms with a positive Laplacian indicates the existence of bond critical point (BCP) (Fig. S1†). A low Mayer bond order (0.17) and an attractive NCI have been noticed as well between two Ag atoms of the  $\text{Ag}_{16}$  core (Fig. S2†) which indicates the reality of the nonbonding argentophilic interaction.<sup>53,54</sup> The geometry of the distorted  $\text{Ag}_{16}$  core comprises eight triangular facets and ten square facets (Fig. S3†). The  $\text{Ag}_{16}$  core can also be divided into three layers; one square (top, made by four Ag atoms), one octagon (middle, made by eight Ag atoms), and another square (bottom, made by four Ag atoms) (Fig. S4†). The middle octagon layer is connected with the two square layers from the opposite sides *via* eight square and eight triangular facets. In addition, one square layer is related to another square layer by 45° of rotation, and that classifies the gyro configuration of the core geometry instead of the *ortho*



Scheme 1 Synthetic outline of the adamantanethiolate protected  $\text{Ag}_{16}$  NC.



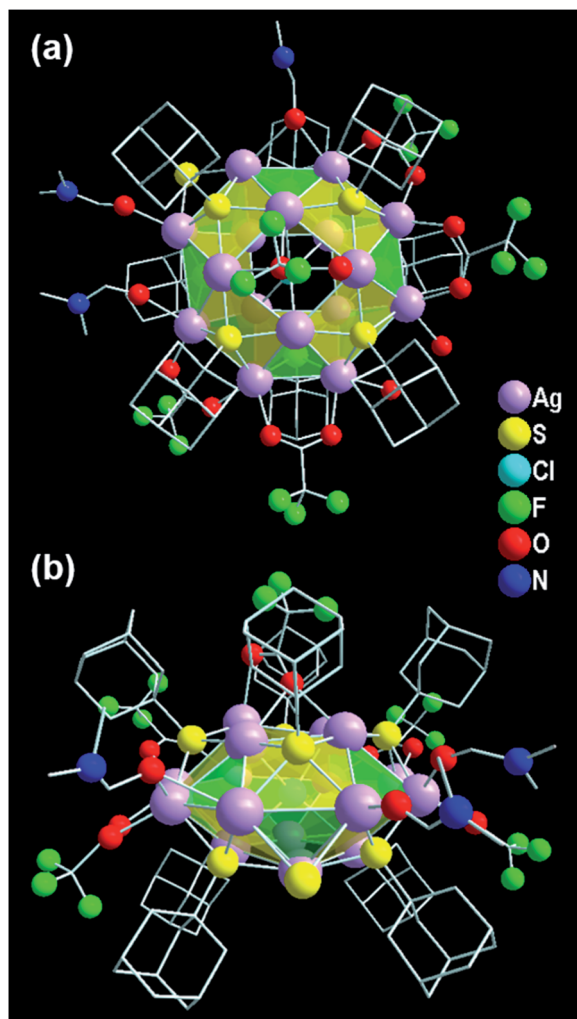


Fig. 1 Structural illustration of the overall  $\text{Ag}_{16}$  NC from (a) top and (b) side views.

(Fig. S5<sup>†</sup>). The chloride atom is enclosed at the center of the octagon layer, resulting in a  $\text{Cl}^-$  templated  $\text{Ag}_{16}$  core (Fig. 2b). We have observed that one  $\text{S}^{2-}$  ion is attached to the octagon layer by  $\mu_2$  coordination mode (Fig. 2c) which could be produced from the breakage of the C–S bond in DMF medium at room temperature.<sup>35</sup> The outer surface of the skeleton is protected by eight  $\text{Adm-S}^-$  ligands (Fig. 2d). Each of the thiolate ligands occupies each of the square facets by  $\mu_4$  ligation mode with Ag–S bond lengths ranging from 2.40 Å to 2.56 Å. The skeleton of the  $\text{Ag}_{16}$  is further capped by the five  $\text{CF}_3\text{COO}^-$  auxiliary ligands (Fig. 2e). Among the five  $\text{CF}_3\text{COO}^-$ , four are bonded with the Ag present in the octagon layer by  $\mu_2$  bridging mode with Ag–O bond lengths in the range of 2.28–2.38 Å, whereas the remaining one is attached with one of the square layers by  $\mu_1$  bridging with an Ag–O bond length of 2.49 Å. Additionally, three DMF molecules and two water molecules are coordinated with the octagon layer of the skeleton (Fig. 2f). Also, one non-coordinated DMF molecule is detected in the lattice by careful crystallographic measurement (Fig. S6<sup>†</sup>).

Further, to verify the chemical composition and identify the charge of the as-synthesized NC as well as its associated

fragments, electrospray ionization mass spectrometry (ESI-MS) is carried out (Fig. 3). In the positive mode ESI-MS spectrum, three prominent peaks are identified. The peak corresponding to the highest  $m/z$  value at 4026.30 is associated with  $[\{\text{Cl}@Ag_{16}\text{S}(\text{S-Adm})_8(\text{CF}_3\text{COO})_5(\text{DMF})_3(\text{H}_2\text{O})_2 \cdot (\text{DMF})\} + \text{H}]^+$ , which further justifies that the cluster itself is neutral. The experimental  $m/z$  value is well-matched with the simulated isotropic patterns, demonstrating the consistency of the assigned molecular formula of the  $\text{Ag}_{16}$  NC with the single-crystal data. The presence of the other two predominant peaks in the ESI-MS spectrum corroborates the associated fragments of  $[\{\text{Cl}@Ag_{15}\text{S}(\text{S-Adm})_8(\text{CF}_3\text{COO})_4(\text{DMF})_3(\text{H}_2\text{O})_2 \cdot (\text{DMF})\} + \text{H}]^+$  and  $[\{\text{Cl}@Ag_{15}\text{S}(\text{S-Adm})_8(\text{CF}_3\text{COO})_4(\text{DMF})_2(\text{H}_2\text{O}) \cdot (\text{DMF})\} + \text{H}]^+$  (Fig. S7<sup>†</sup>). Further, to verify the stability of the as-synthesized NC,  $^1\text{H}$ , and  $^{19}\text{F}$  nuclear magnetic resonance (NMR) studies are performed in  $\text{CDCl}_3$ . The shift of the adamantane proton in the NC compared to the free  $[\text{Ag-Adm}]_n$  complex confirms the presence of bonded Ag-Adm moieties in the structure of the  $\text{Ag}_{16}$  NC (Fig. S8<sup>†</sup>). Moreover, the presence of F atoms in the structure is also proved by  $^{19}\text{F}$  NMR (Fig. S9<sup>†</sup>).

The good agreement of the experimental powder X-ray diffraction patterns with the simulated data further dictates the high purity of the crystals of  $\text{Ag}_{16}$  NC (Fig. S10<sup>†</sup>). The high-resolution transmission electron microscopic (HR-TEM) image confirms that the diameter of the individual NC is  $\sim 0.8$  nm (Fig. 4a). The corresponding scanning electron microscopy (SEM) image and optical microscopic image of the bulk crystals corroborate the block shape of this NC (Fig. 4b). The X-ray photoelectron spectroscopy (XPS) survey spectrum illustrates the existence of the respective elements in this NC (Fig. 4c), that is matched with the energy dispersive X-ray analysis (Fig. S11<sup>†</sup>). The high-resolution binding-energy spectra of each element reveal their corresponding oxidation states. The binding energy spectrum of Ag 3d demonstrates the presence of Ag(I) by showing two peaks of  $3d_{5/2}$  and  $3d_{3/2}$  at 368.76 eV and 374.76 eV, respectively (Fig. 4d). The existence of S, Cl, F, O, N, and C is also confirmed (Fig. S12<sup>†</sup>).

To elucidate the photophysical properties of this newly synthesized adamantanethiol-protected  $\text{Ag}_{16}$  NC, UV-vis absorbance is measured in DCM solvent. A monotonic decrease in the absorbance spectrum with a small shoulder peak at 374 nm is observed (Fig. 5a). From the time-dependent density functional theory (TD-DFT) analysis, the calculated Kohn–Sham orbital population suggests that the occupied states have the major contribution of s- and p-like states of the ligand, whereas the unoccupied orbitals are mainly driven through the s-, p-, and d-like states of the core (Fig. 5b). Hence, the ligand to core transition is clearly understood by the Kohn–Sham orbital population analysis. Further, the computational study reveals that the shoulder peak at 369 nm originates due to the optical transition of  $\text{HOMO}-4 \rightarrow \text{LUMO}$  (Fig. 5c). The frontier molecular orbitals arrangement describes that the low intense peak of 369 nm is mainly constituted by the transition occurring from the p-like (2p) bonding state of adamantane carbon (ligand) to the antibonding states of Ag (4d) and S (3p). This transition is related to the small shoulder peak at 374 nm in the experimental absorbance spectrum. Thus, the observation of





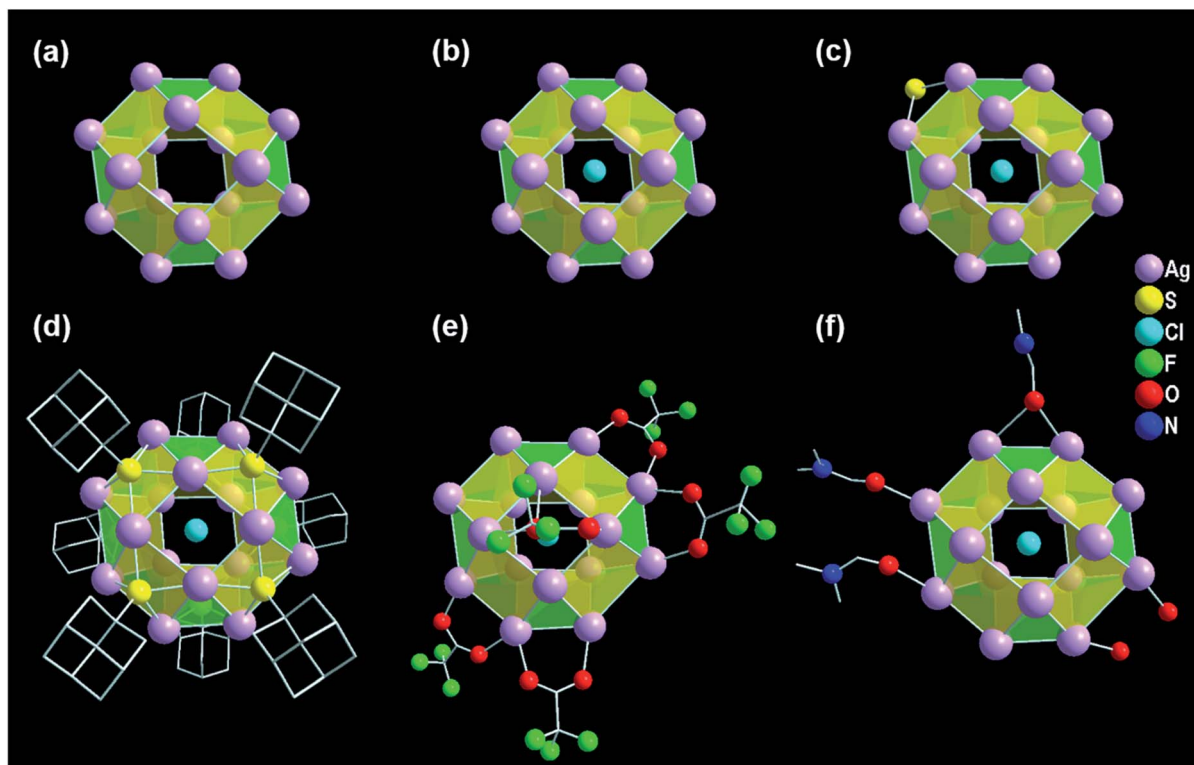


Fig. 2 Structural investigation of the  $\text{Ag}_{16}$  NC. (a) Square gyrobicupola geometry of the  $\text{Ag}_{16}$  skeleton, (b) chloride ion template, (c) attachment of  $\text{S}^{2-}$  with the middle octagon layer of the skeleton, (d) connection of eight S atoms of the thiolate groups with the square facets of the skeleton, and (e) and (f) linking of  $\text{CF}_3\text{COO}^-$  as auxiliary ligands and DMF and water as solvents with  $\text{Ag}_{16}$  skeleton, respectively.

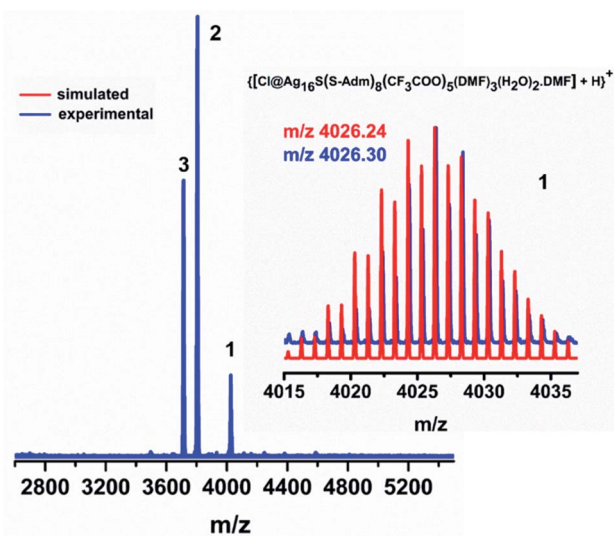


Fig. 3 Positive mode ESI-MS spectrum of the as-synthesized  $\text{Ag}_{16}$  NC over the  $m/z$  range from 2600 to 5600. Inset: experimental and simulated patterns of the parent  $\text{Ag}_{16}$  NC.

low intense peak is mainly derived due to the transition from the ligand to the core which justifies the simulated Kohn–Sham orbital population analysis of the  $\text{Ag}_{16}$  NC dissolved in DCM.

This as-synthesized NC is not emissive at room temperature. However, the strategical modification of the cluster surface

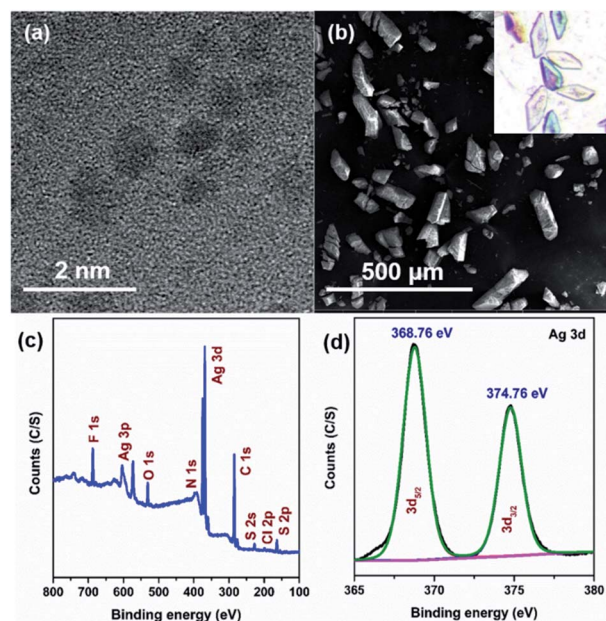


Fig. 4 (a) HR-TEM and (b) SEM images of the  $\text{Ag}_{16}$  NC, (c) XPS survey spectrum, and (d) binding energy spectrum of Ag of the  $\text{Ag}_{16}$  NC.

bulkiness using  $\beta$ -CD through host–guest interaction is able to generate the luminescence properties of this  $\text{Ag}_{16}$  NC at room temperature upon excitation at 380 nm (Fig. S13†). Owing to the



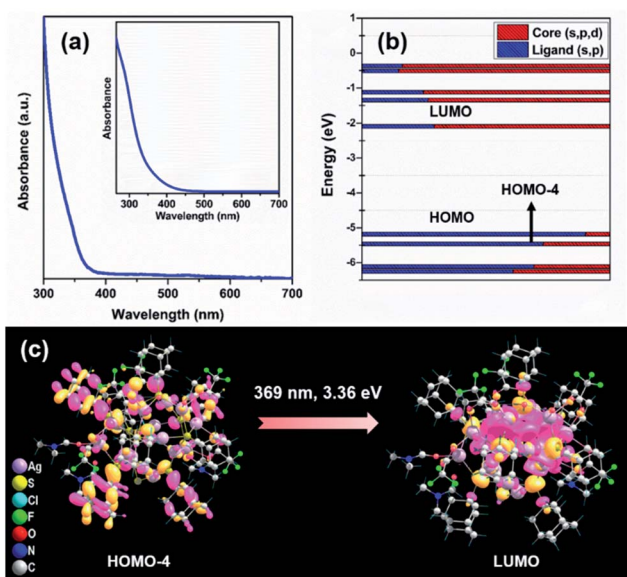


Fig. 5 (a) UV-vis absorbance spectrum of the  $\text{Ag}_{16}$  NC dissolved in DCM, inset: simulated absorbance spectrum of the  $\text{Ag}_{16}$  NC, (b) Kohn-Sham molecular energy diagram of the NC, and (c) corresponding transitions and related frontier-molecular orbitals diagram.

specific size of the inner hydrophobic cavity, the guest  $\beta$ -CD molecule can easily encapsulate the hydrophobic surface of the host NC *i.e.*, the adamantane group which can act as a recognition site for constructing the  $\beta$ -CD-functionalized stable host-guest hybrid assemblies.<sup>44,55</sup> To verify the intermolecular host-guest interaction between the adamantane moiety and the  $\beta$ -CD molecule, the  $^1\text{H}$  NMR study shows more splitting at the adamantane region which confirms the insertion of the ligand in the  $\beta$ -CD cavity (Fig. S14<sup>†</sup>). However, no certain change is witnessed from the UV-vis absorbance spectra of  $\text{Ag}_{16}$  NC after the supramolecular adduct formation with  $\beta$ -CD, suggesting the unaffected ground-state electronic structure (Fig. S15<sup>†</sup>). Further, the electrochemical properties of the parent  $\text{Ag}_{16}$  NC *vs.*  $\beta$ -CD functionalized  $\text{Ag}_{16}$  NC dissolved in DCM are examined. In cyclic voltammetry (CV) and differential pulse voltammetry (DPV) measurements, the redox peaks disappear in the case of  $\beta$ -CD functionalized  $\text{Ag}_{16}$  NC, which suggests that the host-guest complex is resistant to the redox reaction (Fig. S16<sup>†</sup>). Finally, the number of interacted  $\beta$ -CD molecules with the surface of the NC *via* non-covalent interactions is identified in the solution medium by ESI-MS measurements. The positive mode spectrum of a 1 : 1 mixture of  $\text{Ag}_{16}$  NC with  $\beta$ -CD reveals that the predominate  $m/z$  peak at 2565.15 corresponds to  $\text{Ag}_{16} \cap \beta\text{-CD}_1^{2+}$  ( $\cap$  symbolizes the supramolecular adducts) along with a small amount of  $\text{Ag}_{16}^{2+}$  species and two other associated fragments represented in Fig. S17.<sup>†</sup> The +2 charge originates after the removal of  $\text{S}^{2-}$  from the parent NC which is further validated by the characteristic isotropic distribution with the peak separation of  $m/z \sim 0.50$ . The well-matched experimental and simulated mass spectrum data indicates that only one of the adamantane moieties is entrapped in the  $\beta$ -CD cavity. Interestingly, the mass data of 1 : 2 or excess (1 : 3, 1 : 4, 1 : 5,

and 1 : 8) mixture of  $\text{Ag}_{16}$  NC with  $\beta$ -CD reveals that a maximum of 2 molecules of  $\beta$ -CD can be attached with the NC yielding  $\text{Ag}_{16} \cap \beta\text{-CD}_2^{2+}$  at an  $m/z$  value of 3132.65 along with a lesser amount of  $\text{Ag}_{16} \cap \beta\text{-CD}_1^{2+}$  and  $\text{Ag}_{16}^{2+}$  species (Fig. S18<sup>†</sup>). The packing fashion of the two  $\beta$ -CD molecules anchored on the cluster surface induces steric hindrance in such a way that the approach of any further CD molecules towards the cluster surface is restricted. However, the peak maximum of the adduct is shifted toward the higher  $m/z$  value from the one  $\beta$ -CD attachment to the two  $\beta$ -CD attachments in solution signifying the complexation of  $\beta$ -CD on the NC.

To understand the stability of  $\text{Ag}_{16} \cap \beta\text{-CD}_2$ , we have modelled the stepwise encapsulation process of  $\beta$ -CD over the adamantane group. Initially, we have considered the three different possible attachment positions of  $\beta$ -CD on the  $\text{Ag}_{16}$  NC (S2, S4, S8) depending on the surrounding chemical environment of each adamantane group (Fig. S19<sup>†</sup>). Here, we have considered both the head and tail conformation of  $\beta$ -CD. Our calculated binding energy suggested that the  $\beta$ -CD encapsulation in its head conformation on the S4 position is more favorable compared to other positions due to the lowest binding energy of  $-3.3$  eV (Fig. S20, Table S2 and eqn (S1)<sup>†</sup>). We have then identified the preferred position of the next attachment of  $\beta$ -CD on the  $\text{Ag}_{16}$  NC. For this, seven possible attachment positions (S4-S1, S4-S2, S4-S3, S4-S5, S4-S6, S4-S7, and S4-S8) are considered with respect to the encapsulated  $\beta$ -CD on the S4 position of  $\text{Ag}_{16} \cap \beta\text{-CD}_1$ . Among these, the S4-S6 attachment position is found to be more stable because of the lowest binding energy of  $-2.66$  eV compared to other positions which are experiencing high steric repulsion (Table S3<sup>†</sup>). So, the experimental observation and computational findings suggest that  $\beta$ -CD molecules would always prefer the oppositely-placed adamantane groups over the adjacent for host-guest adduct formation, yielding  $\text{Ag}_{16} \cap \beta\text{-CD}_2$  (Fig. 6a).

So, we have strategically designed the adamantanethiolate-protected  $\text{Ag}_{16}$  NC with inspiration for using their molecular recognition properties to originate the photoluminescence through host-guest chemistry. The precise supramolecular functionalization of the NC with the  $\beta$ -CD restricts the intramolecular motion of this NC in the solution phase. Interestingly, we have observed that the luminescence intensity and emission lifetimes ( $0.48 \mu\text{s}$  to  $1.03 \mu\text{s}$ ) increase with the number of attached  $\beta$ -CD molecules (Fig. 6b-d). The physicochemical interactions arising at the ligand surface may alter the efficacy of the relaxation process, which may eventually result in the improvement of their emission properties. So, the origin of the pronounced enhancement in the PL emission QY (relative QY from 0.12 to 0.26) lies in the formation of the host-guest supramolecular adduct which certainly suppresses the surface ligand motions. The theoretical investigation revealed that at this excitation, charge transition is majorly contributed by the p-like (2p) bonding state of the adamantane moiety. However, a similar UV-vis absorbance study excludes the possibilities of any ground-state overlapping, so the rigidification will clearly affect the relaxation pathway. After calculating the radiative and nonradiative relaxation rates, we have identified that the suppression of surface ligand vibrations is closely correlated





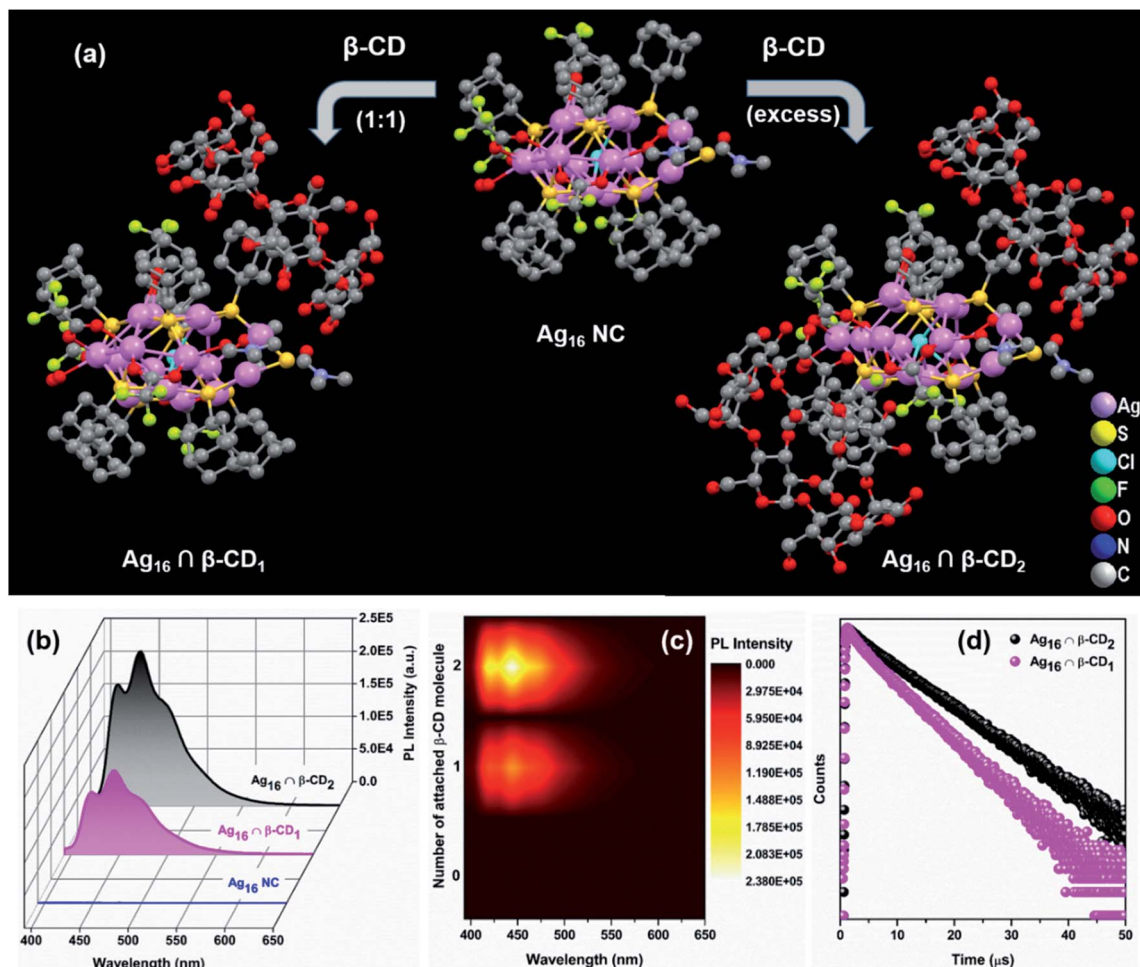


Fig. 6 (a) Reaction scheme of the formation of supramolecular adducts,  $\text{Ag}_{16} \cap \beta\text{-CD}_1$ , and  $\text{Ag}_{16} \cap \beta\text{-CD}_2$  from the parent NC, (b) emission spectra of the  $\text{Ag}_{16}$  NC,  $\text{Ag}_{16} \cap \beta\text{-CD}_1$ , and  $\text{Ag}_{16} \cap \beta\text{-CD}_2$  dissolved in DCM upon excitation at 380 nm, (c) corresponding 2D-contour plot for visualizing the difference in PL intensities with the number of the attached  $\beta\text{-CD}$  molecules, and (d) emission lifetimes data upon excitation at 375 nm.

with the nonradiative relaxation pathway (eqn (S2), (S3) and Table S4†). So, the enhancement in the luminescence behaviour is observed here due to the effective suppression of the non-radiative relaxation rate (from  $1.83 \times 10^6$  to  $7.18 \times 10^5$ ), while the radiative relaxation rate is almost intact after the supramolecular adduct formation. Hence, interfacing individual NCs with functional supramolecular arrangements is an intriguing outlook that delivers new potential.

For utilizing the enhanced emission properties of the  $\beta\text{-CD}$  functionalized supramolecular adduct, *i.e.*,  $\text{Ag}_{16} \cap \beta\text{-CD}_2$  in an artificial light-harvesting application through efficient energy transfer by FRET, an acceptor dye molecule is required whose absorption considerably overlaps with the emission of  $\text{Ag}_{16} \cap \beta\text{-CD}_2$ .<sup>7</sup> Here, we have strategically chosen  $\beta\text{-carotene}$  as a dye that has a broad absorptivity with an absorbance maximum at 462 nm in DCM (Fig. 7a). The energy transfer process is examined by the fluorescence titration of  $\beta\text{-carotene}$  ( $2 \times 10^{-6}$  M) with the  $\text{Ag}_{16} \cap \beta\text{-CD}_2$  ( $10^{-5}$  M) in DCM upon excitation at 380 nm (Fig. 7b). With the gradual addition of 25  $\mu\text{L}$  of  $\beta\text{-carotene}$  in the  $\text{Ag}_{16} \cap \beta\text{-CD}_2$  containing solution, a drop in the

PL intensity is detected at 444 nm, corresponding to the emission maximum of  $\text{Ag}_{16} \cap \beta\text{-CD}_2$  (donor), while a secondary emission is generated at the  $\sim 540$  nm region which corresponds to the emission of  $\beta\text{-carotene}$  (Fig. 7c, d and S21†). So, this observation clearly suggests the energy transfer from donor  $\text{Ag}_{16} \cap \beta\text{-CD}_2$  to acceptor  $\beta\text{-carotene}$  is facilitated. The CIE chromaticity diagram further verifies the FRET process between acceptor and donor molecules by a change in fluorescence color from blue (co-ordinate of CIE (0.15, 0.09)) to green (co-ordinate of CIE (0.36, 0.53)) *via* an intermediate color of the donor-acceptor system of light blue (co-ordinate of CIE (0.24, 0.32)) (Fig. 7e). The considerable overlap between the emission and absorption band of the donor and acceptor molecules is the primary factor for the energy transfer process, while the two opposite surface zeta potential values ( $\zeta$  of  $\text{Ag}_{16} \cap \beta\text{-CD}_2 = 60$  mV, and  $\zeta$  of  $\beta\text{-carotene} = -29$  mV) support the formation of the adducts for charge transfer feasibility (Fig. S22† and Scheme 2). However, the unchanged absorbance spectra with continuous addition of the similar amount of  $\beta\text{-carotene}$  on the  $\text{Ag}_{16} \cap \beta\text{-CD}_2$  solution eliminates the possibility of the ground



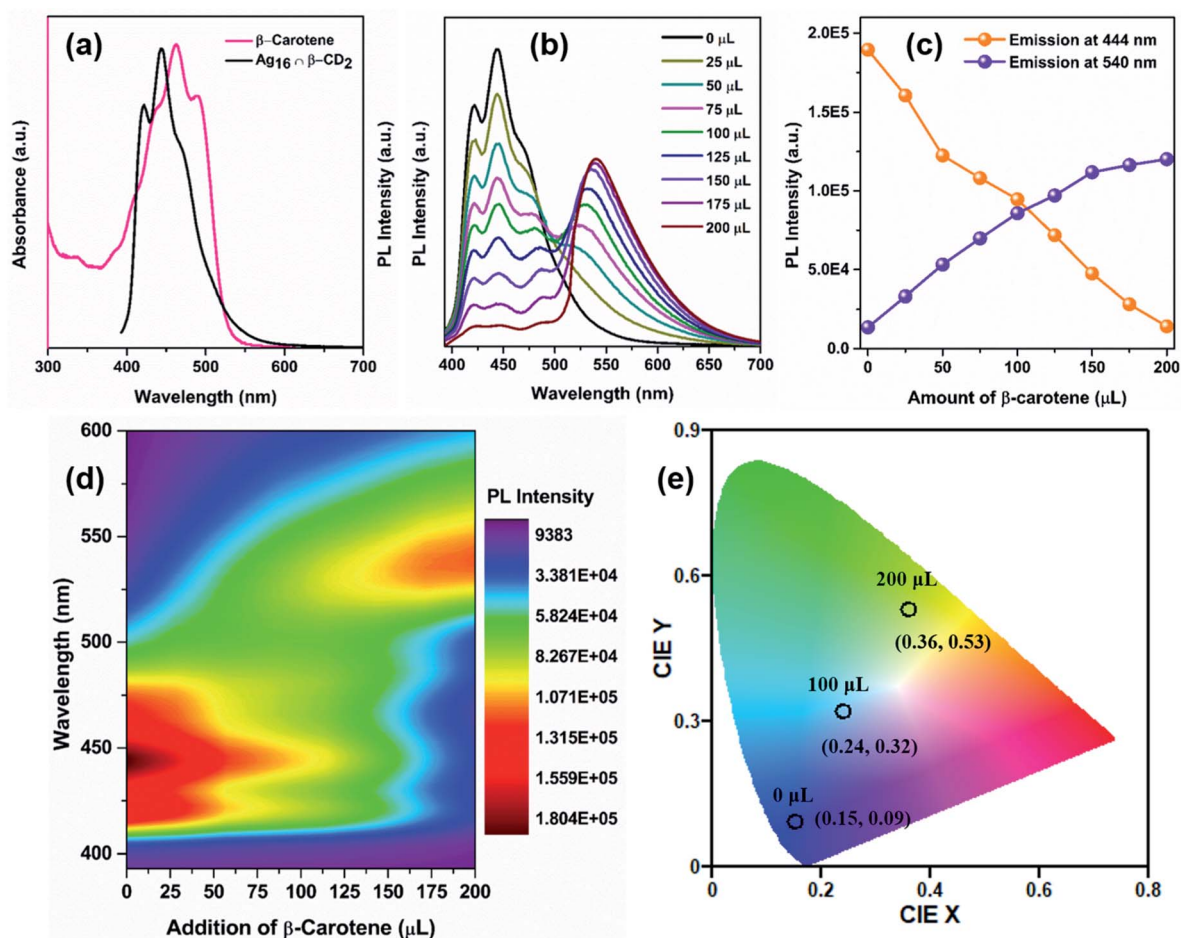
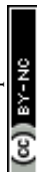


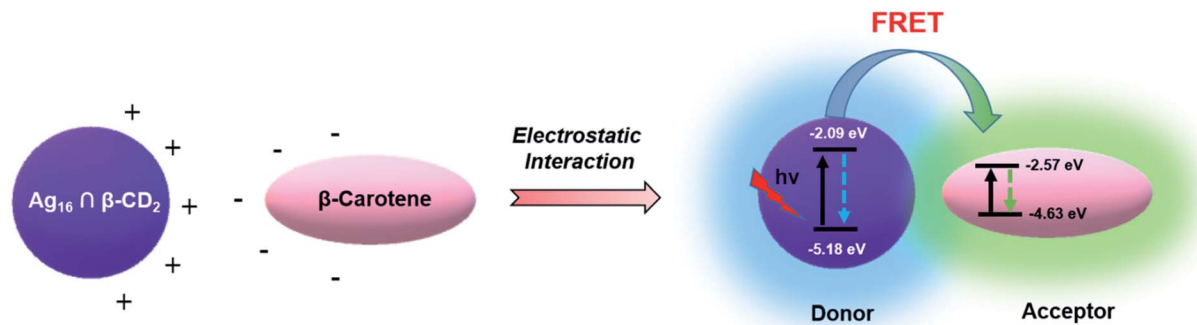
Fig. 7 (a) Absorbance spectrum of acceptor  $\beta$ -carotene and emission spectrum of donor  $\text{Ag}_{16} \cap \beta\text{-CD}_2$ , (b) emission spectra of the donor molecule with the addition of the measured amount of  $\beta$ -carotene ( $\lambda_{\text{ex}} = 380 \text{ nm}$ ) in DCM, (c) variation in emission intensity for  $\text{Ag}_{16} \cap \beta\text{-CD}_2$  during the addition of the same amount of  $\beta$ -carotene at 444 nm and 540 nm, (d) associated 2D-contour plot of the FRET process for the visualization of the PL intensity transferring process, and (e) change of 1931 CIE chromaticity coordinates of  $\text{Ag}_{16} \cap \beta\text{-CD}_2$  upon titration with  $\beta$ -carotene.

state interaction (Fig. S23<sup>†</sup>). This observation is consistent with the  $^1\text{H}$  NMR spectrum of the  $\text{Ag}_{16} \cap \beta\text{-CD}_2$  and  $\beta$ -carotene mixture which displays the essentially non-shifted proton peaks compared to the individual components (Fig. S24<sup>†</sup>). So, the energy transfer exclusively originates from the excited state of the  $\text{Ag}_{16} \cap \beta\text{-CD}_2$  by forming an effective sphere with the acceptor that leads to a fast quenching of the excited donor before they diffuse apart. The continuous enhancement in the emission intensity at a higher wavelength region gets saturated after the addition of 200  $\mu\text{L}$  of the  $\beta$ -carotene solution to 2 mL of  $\text{Ag}_{16} \cap \beta\text{-CD}_2$ , suggesting the detectable maximum energy transfer limit when the maximum number of acceptor molecules is present in the effective sphere. The theoretically optimized structure of this mixture suggests that both molecules are maintaining an average  $\sim 2 \text{ nm}$  gap at the ground state (Fig. S25<sup>†</sup>). However, the Förster radius of the effective sphere ( $\sim 3.7 \text{ nm}$ ) is calculated experimentally by eqn (S4)<sup>†</sup> from the average emission lifetime and the energy transfer rate per donor–acceptor pair.<sup>56–60</sup> As per the size of each donor molecule, it is obvious that during the excitation, both the molecules will

be in a favorable position for the FRET. The calculated radiative rate of the donor ( $2.52 \times 10^5 \text{ s}^{-1}$ ) is much slower than the radiative energy transfer rate between the donor and acceptor ( $2.6 \times 10^7 \text{ s}^{-1}$ ). Thus, we can assume that during the excitation of a donor, acceptor molecules are present in a favorable position and the excited donor molecule can promptly transfer the energy to the adjacent acceptor molecules which results in a quenching of the excited state of the donor (Scheme 2).<sup>8</sup> The energy transfer efficiency between the donor and acceptor is calculated from the emission intensities to be 93% by following eqn (S5).<sup>†</sup> The additional evidence for the energy transfer is envisioned by obtaining a 17 times shorter average emission lifetime of the conjugated materials than the parent  $\text{Ag}_{16} \cap \beta\text{-CD}_2$  (Fig. S26<sup>†</sup>). In a controlled experiment, the conjugated material and  $\beta$ -carotene were excited at the original excitation of the  $\beta$ -carotene, showing an impressive antenna effect of 34 (Fig. S27 and eqn (S6)<sup>†</sup>).

To explore the practical usability of this fabricated LHS, we explore its photocurrent generation ability under irradiation of sunlight at a 10 V bias on a patterned gold electrode. As sunlight





Scheme 2 Proposed mechanism of the FRET process between  $\text{Ag}_{16} \cap \beta\text{-CD}_2$  (donor) and  $\beta$ -carotene (acceptor).

is a combination of UV and visible light, the  $\beta$ -carotene has low absorption in the UV region whereas the synthesized  $\text{Ag}_{16} \cap \beta\text{-CD}_2$  exhibits strong absorption in the UV region. So, as a combination, the conjugation serves as an antenna source to efficiently transfer the energy from the UV-to-visible region to generate the maximum photocurrent (472 nA) on the gold electrode (Fig. S28<sup>†</sup>), whereas the individual components show a much lesser photocurrent generation ability which is depicted in a 2D contour plot (Fig. S29<sup>†</sup>).

## Conclusions

In summary, a unique artificial LHS has been strategically designed with an outstanding energy transfer efficiency based on atom-precise Ag(I) NC. Initially, we successfully synthesized bulky ligand-protected  $\text{Ag}_{16}$  NC with the small core *via* controlling the reaction condition. Owing to the hydrophobic nature of the cluster surface, we have implemented a versatile approach of specific host-guest supramolecular adduct formation between  $\beta$ -CD and the adamantane ligand anchored on the cluster. Due to this host-guest interaction, we have effectively generated luminescence properties in the  $\text{Ag}_{16}$  NC at room temperature by altering the relaxation mechanism. This adduct is further utilized to construct the artificial LHS through an efficient energy transfer process through FRET occurring from the donor ( $\text{Ag}_{16} \cap \beta\text{-CD}_2$ ) to the acceptor ( $\beta$ -carotene) with an energy transfer efficiency of 93% and a good antenna effect. The energy transfer process is well justified by the decrease in emission lifetime and emission color change from blue to green which is depicted in the CIE chromaticity diagram. Finally, the designed LHS has been successfully utilized as a source of unconventional energy generation with much-enhanced yields compared to the individual components. So, the unique design strategy of the atom-precise nanocluster based-LHS may surely open up a new direction for nanocluster research in sustainable energy generation.

## Experimental section

### Synthesis of $\text{Ag}_{16}$ NC

An equimolar (0.1 mmol) mixture of  $[\text{AgS-Adm}]_n$  complex and  $\text{CF}_3\text{COOAg}$  was treated in 3 mL DMF at room temperature for

15 min. Then, 18 mg  $\text{PPh}_4\text{Cl}$  was added to the clear mixture and continued stirring for a further 30 min. A white color precipitate was formed which was separated by centrifugation. The clear solution mixture was kept in a dark environment for crystallization. After 3 days, colorless crystals were obtained at room temperature.

The  $[\text{AgS-Adm}]_n$  complex was prepared by a reported method where an equivalent amount of  $\text{AgNO}_3$  was treated with  $\text{AdmSH}$  and  $\text{Et}_3\text{N}$  in methanol.

### Synthesis of $\text{Ag}_{16} \cap \beta\text{-CD}_n$ (where $n = 1, 2$ )

After dissolving  $\text{Ag}_{16}$  NC (3 mg) in 4 mL DCM, a 2 mL aqueous solution of  $\beta$ -CD was added in such a way that the desired  $\text{Ag}_{16}/\beta\text{-CD}$  ratio was preserved in the mixture (1 : 1 for  $\text{Ag}_{16} \cap \beta\text{-CD}_1$  and 1 : 2, 1 : 3, 1 : 4, 1 : 5, and 1 : 8 for  $\text{Ag}_{16} \cap \beta\text{-CD}_2$ ). Then, the above solution mixture was carefully kept under sonication for 20 min. Further, the solution mixture was stirred for 40 min with intermittent sonication for 3 min at 10 min intervals. After separating the aqueous phase, the organic phase was dried by a rotatory evaporator.

## Data availability

The ESI<sup>†</sup> contains experimental details, crystal structure refinement parameters, binding energies and emission properties of the host-guest inclusion complexes, QTAIM molecular plot of the  $\text{Ag}_{16}$  NC, RDG isosurface of argentophilic interaction, different facets of the  $\text{Ag}_{16}$  skeleton, different layers of the  $\text{Ag}_{16}$  skeleton, relation between the top and bottom square facets, presence of the lattice DMF molecule in the lattice of the  $\text{Ag}_{16}$  NC, positive mode ESI-MS data of the  $\text{Ag}_{16}$  NC,  $^1\text{H}$  NMR spectrum in  $\text{CDCl}_3$  of  $\text{Ag}_{16}$  NC,  $^{19}\text{F}$  NMR spectrum in  $\text{CDCl}_3$  of the  $\text{Ag}_{16}$  NC, PXRD, EDS analysis of the  $\text{Ag}_{16}$  NC, deconvoluted XPS spectra of each element of the  $\text{Ag}_{16}$  NC, UV-vis spectrum of the  $\text{Ag}_{16}$  NC,  $^1\text{H}$  NMR spectra in  $\text{CDCl}_3$  of the  $\beta$ -CD attached  $\text{Ag}_{16}$  NC and pure  $\beta$ -CD, UV-vis absorbance spectra of the  $\text{Ag}_{16}$  NC and the  $\beta$ -CD functionalized  $\text{Ag}_{16}$  NC, CV, DPV of the  $\text{Ag}_{16}$  NC and the  $\beta$ -CD functionalized  $\text{Ag}_{16}$  NC, partial positive mode ESI-MS data of the  $\text{Ag}_{16} \cap \beta\text{-CD}_1^{2+}$ , partial positive mode ESI-MS data of the  $\text{Ag}_{16} \cap \beta\text{-CD}_2^{2+}$ , numbering of the possible  $\beta$ -CD attachment sites on the  $\text{Ag}_{16}$  NC, binding energy calculation





of the tail and head configuration of  $\beta$ -CD attachment on the stable S4 position, PL emission of the organic dye  $\beta$ -carotene, zeta potential data of the donor and acceptor, UV-visible spectra of  $\text{Ag}_{16} \cap \beta\text{-CD}_2$  with the gradual addition of  $\beta$ -carotene,  $^1\text{H}$  NMR spectra in  $\text{CDCl}_3$  of the  $\beta$ -CD attached  $\text{Ag}_{16}$  NC with  $\beta$ -carotene and pure  $\beta$ -carotene, optimized distance between the donor and acceptor, emission lifetime data of  $\text{Ag}_{16} \cap \beta\text{-CD}_2$  and  $\text{Ag}_{16} \cap \beta\text{-CD}_2 + \beta$ -carotene, fluorescence intensity of the donor-acceptor system to calculate the antenna effect, photocurrent from the donor-acceptor light-harvesting system, 2-D contour plot for differentiating the photocurrent generation, and references.

## Author contributions

A. K. D. and S. B. performed the synthesis, characterization and data interpretation of the nanocluster and light harvesting system. S. S. M. and B. P. performed the theoretical calculation. S. K. M. was involved in manuscript preparation. All authors discussed the results and were involved in manuscript writing.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

S. B. acknowledges the NPDF fund from SERB (PDF/2020/001085).

## References

- G. McDermott, S. Prince, A. Freer, A. Hawthornthwaite-Lawless, M. Papiz, R. Cogdell and N. Isaacs, *Nature*, 1995, **374**, 517–521.
- G. D. Scholes, G. R. Fleming, A. Olaya-Castro and R. Van Grondelle, *Nat. Chem.*, 2011, **3**, 763–774.
- P. D. Frischmann, K. Mahata and F. Würthner, *Chem. Soc. Rev.*, 2013, **42**, 1847–1870.
- A. Ajayaghosh, V. K. Praveen and C. Vijayakumar, *Chem. Soc. Rev.*, 2008, **37**, 109–122.
- H.-Q. Peng, L.-Y. Niu, Y.-Z. Chen, L.-Z. Wu, C.-H. Tung and Q.-Z. Yang, *Chem. Rev.*, 2015, **115**, 7502–7542.
- A. H. Proppe, Y. C. Li, A. Aspuru-Guzik, C. P. Berlinguette, C. J. Chang, R. Cogdell, A. G. Doyle, J. Flick, N. M. Gabor and R. van Grondelle, *Nat. Rev. Mater.*, 2020, **5**, 828–846.
- A. Kumar, R. Saha and P. S. Mukherjee, *Chem. Sci.*, 2021, **12**, 5319–5329.
- Y. Li, S. S. Rajasree, G. Y. Lee, J. Yu, J.-H. Tang, R. Ni, G. Li, K. N. Houk, P. Deria and P. J. Stang, *J. Am. Chem. Soc.*, 2021, **143**, 2908–2919.
- M. Hao, G. Sun, M. Zuo, Z. Xu, Y. Chen, X. Y. Hu and L. Wang, *Angew. Chem., Int. Ed.*, 2020, **59**, 10095–10100.
- J. J. Li, Y. Chen, J. Yu, N. Cheng and Y. Liu, *Adv. Mater.*, 2017, **29**, 1701905.
- D. Zhang, W. Yu, S. Li, Y. Xia, X. Li, Y. Li and T. Yi, *J. Am. Chem. Soc.*, 2021, **143**, 1313–1317.
- S. Kundu and A. Patra, *Chem. Rev.*, 2017, **117**, 712–757.
- M. R. Wasielewski, *Acc. Chem. Res.*, 2009, **42**, 1910–1921.
- Y. Zeng, Y. Li, M. Li, G. Yang and Y. Li, *J. Am. Chem. Soc.*, 2009, **131**, 9100–9106.
- F. Biedermann, E. Elmalem, I. Ghosh, W. M. Nau and O. A. Scherman, *Angew. Chem.*, 2012, **124**, 7859–7863.
- K. Liu, Y. Liu, Y. Yao, H. Yuan, S. Wang, Z. Wang and X. Zhang, *Angew. Chem., Int. Ed.*, 2013, **52**, 8285–8289.
- Y. Yao, X. Chi, Y. Zhou and F. Huang, *Chem. Sci.*, 2014, **5**, 2778–2782.
- Z. Xu, S. Peng, Y. Y. Wang, J. K. Zhang, A. I. Lazar and D. S. Guo, *Adv. Mater.*, 2016, **28**, 7666–7671.
- C.-L. Sun, H.-Q. Peng, L.-Y. Niu, Y.-Z. Chen, L.-Z. Wu, C.-H. Tung and Q.-Z. Yang, *Chem. Commun.*, 2018, **54**, 1117–1120.
- R. Jin, C. Zeng, M. Zhou and Y. Chen, *Chem. Rev.*, 2016, **116**, 10346–10413.
- I. Chakraborty and T. Pradeep, *Chem. Rev.*, 2017, **117**, 8208–8271.
- J. Yang and R. Jin, *ACS Mater. Lett.*, 2019, **1**, 482–489.
- Y. Jin, C. Zhang, X.-Y. Dong, S.-Q. Zang and T. C. Mak, *Chem. Soc. Rev.*, 2021, **50**, 2297–2319.
- X. Kang and M. Zhu, *Chem. Soc. Rev.*, 2019, **48**, 2422–2457.
- Y. Du, H. Sheng, D. Astruc and M. Zhu, *Chem. Rev.*, 2019, **120**, 526–622.
- K. D. M. Weerawardene, H. Häkkinen and C. M. Aikens, *Annu. Rev. Phys. Chem.*, 2018, **69**, 205–229.
- T. Kawawaki, A. Ebina, Y. Hosokawa, S. Ozaki, D. Suzuki, S. Hossain and Y. Negishi, *Small*, 2021, **17**, 2005328.
- H. Hirai, S. Ito, S. Takano, K. Koyasu and T. Tsukuda, *Chem. Sci.*, 2020, **11**, 12233–12248.
- Q. Yao, Z. Wu, Z. Liu, Y. Lin, X. Yuan and J. Xie, *Chem. Sci.*, 2021, **12**, 99–127.
- I. Chakraborty, W. Kurashige, K. Kanehira, L. Gell, H. Häkkinen, Y. Negishi and T. Pradeep, *J. Phys. Chem. Lett.*, 2013, **4**, 3351–3355.
- M. Ganguly, A. Pal, Y. Negishi and T. Pal, *Langmuir*, 2013, **29**, 2033–2043.
- Q.-M. Wang, Y.-M. Lin and K.-G. Liu, *Acc. Chem. Res.*, 2015, **48**, 1570–1579.
- Z. Wang, R. K. Gupta, G. G. Luo and D. Sun, *Chem. Rec.*, 2020, **20**, 389–402.
- S. Biswas, A. K. Das, A. C. Reber, S. Biswas, S. Bhandary, V. B. Kamble, S. N. Khanna and S. Mandal, *Nano Lett.*, 2022, **22**, 3721–3727.
- D. Sun, H. Wang, H.-F. Lu, S.-Y. Feng, Z.-W. Zhang, G.-X. Sun and D.-F. Sun, *Dalton Trans.*, 2013, **42**, 6281–6284.
- S. Yuan, Y.-K. Deng, X.-P. Wang and D. Sun, *New J. Chem.*, 2013, **37**, 2973–2977.
- K. Zhou, C. Qin, X.-L. Wang, K.-Z. Shao, L.-K. Yan and Z.-M. Su, *CrystEngComm*, 2014, **16**, 7860–7864.
- M. J. Alhilaly, R.-W. Huang, R. Naphade, B. Alamer, M. N. Hedhili, A.-H. Emwas, P. Maity, J. Yin, A. Shkurenko, O. F. Mohammed and O. M. Bakr, *J. Am. Chem. Soc.*, 2019, **141**, 9585–9592.



- 39 E. Khatun, A. Ghosh, P. Chakraborty, P. Singh, M. Bodiuzzaman, P. Ganesan, G. Natarajan, J. Ghosh, S. K. Pal and T. Pradeep, *Nanoscale*, 2018, **10**, 20033–20042.
- 40 X. Kang and M. Zhu, *Coord. Chem. Rev.*, 2019, **394**, 1–38.
- 41 A. Nag and T. Pradeep, *ACS Nanosci. Au*, 2022, **2**, 160–178.
- 42 M. A. Ganayee, C. Manju, W. A. Dar, B. Mondal and T. Pradeep, *Ind. Eng. Chem. Res.*, 2020, **59**, 12737–12744.
- 43 T. Jiang, G. Qu, J. Wang, X. Ma and H. Tian, *Chem. Sci.*, 2020, **11**, 3531–3537.
- 44 A. Mathew, G. Natarajan, L. Lehtovaara, H. Hakkinen, R. M. Kumar, V. Subramanian, A. Jaleel and T. Pradeep, *ACS Nano*, 2014, **8**, 139–152.
- 45 A. Nag, P. Chakraborty, A. Thacharon, G. Paramasivam, B. Mondal, M. Bodiuzzaman and T. Pradeep, *J. Phys. Chem. C*, 2020, **124**, 22298–22303.
- 46 C. Yan, C. Liu, H. Abroshan, Z. Li, R. Qiu and G. Li, *Phys. Chem. Chem. Phys.*, 2016, **18**, 23358–23364.
- 47 M. A. H. Muhammed, L. K. Cruz, A. H. Emwas, A. M. El-Zohry, B. Moosa, O. F. Mohammed and N. M. Khashab, *Angew. Chem.*, 2019, **131**, 15812–15817.
- 48 H.-H. Deng, X.-Q. Shi, F.-F. Wang, H.-P. Peng, A.-L. Liu, X.-H. Xia and W. Chen, *Chem. Mater.*, 2017, **29**, 1362–1369.
- 49 L. Ren, P. Yuan, H. Su, S. Malola, S. Lin, Z. Tang, B. K. Teo, H. Häkkinen, L. Zheng and N. Zheng, *J. Am. Chem. Soc.*, 2017, **139**, 13288–13291.
- 50 A. K. Das, S. Biswas, A. Thomas, S. Paul, A. S. Nair, B. Pathak, M. S. Singh and S. Mandal, *Mater. Chem. Front.*, 2021, **5**, 8380–8386.
- 51 H. Schmidbaur and A. Schier, *Angew. Chem., Int. Ed.*, 2015, **54**, 746–784.
- 52 T. Lu and F. Chen, *J. Comput. Chem.*, 2012, **33**, 580–592.
- 53 A. K. Das, S. Biswas, S. S. Manna, B. Pathak and S. Mandal, *Inorg. Chem.*, 2021, **60**, 18234–18241.
- 54 A. K. Das, S. Maity, T. Sengupta, D. Bista, A. C. Reber, A. Patra, S. N. Khanna and S. Mandal, *J. Phys. Chem. Lett.*, 2021, **12**, 2154–2159.
- 55 M. Mohamadhoseini and Z. Mohamadnia, *Coord. Chem. Rev.*, 2021, **432**, 213711.
- 56 T. Förster, *Ann. Phys.*, 1948, **437**, 55–75.
- 57 R. C. Powell and Z. G. Soos, *J. Lumin.*, 1975, **11**, 1–45.
- 58 S. Crooker, J. Hollingsworth, S. Tretiak and V. I. Klimov, *Phys. Rev. Lett.*, 2002, **89**, 186802.
- 59 S. Halivni, A. Sitt, I. Hadar and U. Banin, *ACS Nano*, 2012, **6**, 2758–2765.
- 60 A. J. Mork, M. C. Weidman, F. Prins and W. A. Tisdale, *J. Phys. Chem. C*, 2014, **118**, 13920–13928.

