

REVIEW

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2, 7620Imprinting chirality in inorganic nanomaterials for
optoelectronic and bio-applications: strategies,
challenges, and opportunitiesPrakash Chandra Mondal,^a Deepak Asthana,^b Ranjeev Kumar Parashar^a and
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Chirality, ubiquitous in biological systems, is the property of the non-superimposable structures found in both macroscopic and nanoscopic objects. In the last ten years, the research on chiral inorganic nanomaterials has witnessed tremendous growth due to their shape, size, crystal facets-controlled unparalleled physical, chemical, optoelectronic, and bio-compatibility features. Due to the quantum confinement, the chiral nanomaterials exhibit high anisotropy factors and, therefore, there has been an upsurge in research activities around the chiroptical and (bio)recognition properties of optically active inorganic nanomaterials. Advancements in synthesis and characterization technologies have made it possible to prepare complex multi-functional materials. Inorganic nanoparticles are known to exhibit fascinating semiconducting or metallic behavior and when combined with chirality, their usability in various devices is enhanced drastically due to an additional degree of freedom over achiral systems. In this review, we discuss different chemical routes to transfer chirality from intrinsic chiral systems into achiral nanomaterials for optoelectronics and bio-applications, in the light of recent experimental findings. Despite the progress in the research field of nanomaterials, imprinting chirality, device fabrication, scalability, long-term stability, and reproducibility remain challenging tasks that need to be addressed. Since the synthesis and growth mechanisms of chiral inorganic systems can be found in literature, here, we shed light only on the complimentary metal-oxides semiconductors (CMOS), and bio-compatibility of either intrinsic or imprinted chiral nanomaterials and their potential applications in next-generation chiroptical devices. We also discuss feasible strategies to resolve existing issues related to the integration into devices and their operational aspects.

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Key learning points

- (1) Origin of homochirality and chirality transfer from intrinsic chiral systems to achiral inorganic nanomaterials.
- (2) Various routes to functionalize chiral nanomaterials for governing size, shape, and optical properties.
- (3) Recent trends in optoelectronic devices and bio-applications.
- (4) Future scopes, challenges, and possible solutions.

1. Introduction

Chirality or handedness refers to materials, molecules, and architectures that lack, a plane, center, or axis of symmetry.

In other words, an object that has non-superimposable mirror images is called chiral. The concept of chirality in molecules was first demonstrated by Louis Pasteur in 1848,¹ while the phrase 'chirality' was first coined by Lord Kelvin in the year 1894.² This finding has paved a new avenue for discovering many hidden phenomena at the macroscopic and also nanoscopic levels. Chirality appears as a 'holy grail' that can help us understand why nature prefers homochirality, which is a 100% enantiomeric excess (ee).³ In the recent past, chiral molecules have attracted the immense attention of researchers, particularly for their diverse applications in chiral optical switches,

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bio-sensors, and optoelectronic devices.^{4–7} Chiral systems show a high degree of chiral anisotropy that stems from strong resonance upon interacting with the electromagnetic waves. Thus, the chiral systems can rotate the plane-polarized electromagnetic waves (light in general) in either the left or right direction, depending on the optical configuration of the system. Of late, through the lenses of experiments and theoretical studies, Ron Naaman and others have found that organic chiral molecules such as DNA, oligopeptides, proteins, and polymers are capable of filtering the spin when a spin-polarized current passes through them.^{8–11} However, chiral inorganic nano-materials are relatively less explored than (bio)organic molecules.^{12–16} Unlike the chiral organic materials, the exact understanding of the source of chirality in inorganic materials, including metallic or semiconductors, is still in the development stage.^{13,17,18} It has been observed that in many cases during the nanoparticle formation, the metal atoms in the cluster attained a lower symmetry and exhibited chiral effects.^{19–21} For instance,

the Whetten group developed a method to prepare an optically active thiourea-based gold cluster compound $\text{Au}_{28}(\text{SG})_{16}$ (where $\text{SG} = N\text{-}\gamma\text{-glutamyl-cysteinyglycine}$).²² The circular dichroism (CD) spectrum of this cluster exhibited a peak in the gold absorption band regions, which is an exciting feature observed with a chiral metal nanoparticle system. The Kelly group demonstrated that the microwave-assisted preparation of chiral luminescent CdS quantum dots (QDs) utilizing racemic and enantiomerically pure penicillamine.²³ The CD spectra of semiconductor chiral CdS QDs display distinct signatures in the UV region (200–390 nm). Dynamic monitoring of the CD signals and quantum mechanical modeling by density functional theory (DFT) calculation suggest that the nanoparticle (NPs) core remains achiral whereas the surface acquires chirality and contributes to new signals in the CD spectrum.

A rather simpler approach to imprint chirality in an inorganic nanoparticle is to coat its surface with optically active moieties by performing surface engineering or ligand



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exchange. Surface engineering has found special applicability in the field of (chiral)nanoparticles (NPs).^{13,24–27} Two approaches are generally adopted for surface functionalization: one in which the nanoparticle core is coated with some inorganic shell such as ZnS or ZnSe to give core/shell or heterostructures, and the second in which the nanoparticle surface is protected with organic ligands.^{28,29} While core/shell type modification helps in removing surface defects, improves photo-stability and quantum efficiency, ligand exchange plays a vital function in controlling the overall usability of the induced chirality either in NPs or nanocrystals (NCs).^{30,31} Ligand exchange helps to tune the solubility or stability against aggregation, and further functionalization of nanoparticles.^{32,33} The nanoparticle surface can be modified with an intrinsic chiral ligand generally containing functional surface-active head groups such as $-\text{NH}_2$, $-\text{COOH}$, $-\text{SH}$, N_3 , *etc.*, which would allow the further attachment of any desired moiety around the NP's surface.^{34–40} In general, the choice of functionalizing reagents depends very much on the nature of the nanoparticles; for example, thiol derivatives for gold and silver nanoparticles, carboxylic or phosphoric derivatives for metal oxides, and silane-based compounds for silica nanoparticles. This approach has been widely used in the field of biosensing and imaging and has been reviewed elsewhere.^{26,27,41,42} Here, we mainly focus on the three sections, the first section deals with the chirality transfer and surface engineering, the second part talks about optoelectronic device integration, and the third section discusses bio-applications of the chiral inorganic nanomaterials. A schematic overview encompassing the subject matter of this review is illustrated in Fig. 1.

2. Origin of chirality and the surface engineering of chiral nanomaterials

The chirality in any object appears due to the non-zero numerical function of the dot product between two components, electric dipole moment transition μ , and magnetic dipole moment transition m , according to the Rosenfeld equation.⁴³

$$R = \text{Im}(\mu \cdot m) \quad (1)$$

R denotes the CD intensity at a given transition, and Im is the imaginary section of the scalar product. The concept of inducing chirality in the achiral inorganic NPs enriches the fundamental science domain and various applications. Inorganic chiral NPs originating from noble metals like gold and silver, or CdS/CdSe have emerged as appealing candidates for biocatalysts, sensing, optoelectronics, and spintronic applications, to name a few.^{44–48} Gold NPs have high Miller indices represented by $\{hkl\}$, $h \neq k \neq l \neq 0$, *e.g.*, (321) and (17 11 9), which exhibit either intrinsic or induced chirality. However, chirality in an achiral nanoparticle can also be induced by capping its surface with intrinsic chiral moieties such as amino acids, proteins, peptides, and DNA, *etc.*^{49–52} To exploit chiral materials in practical applications, in-depth knowledge about chiral morphology at the atomic level is highly desired.^{53–56}

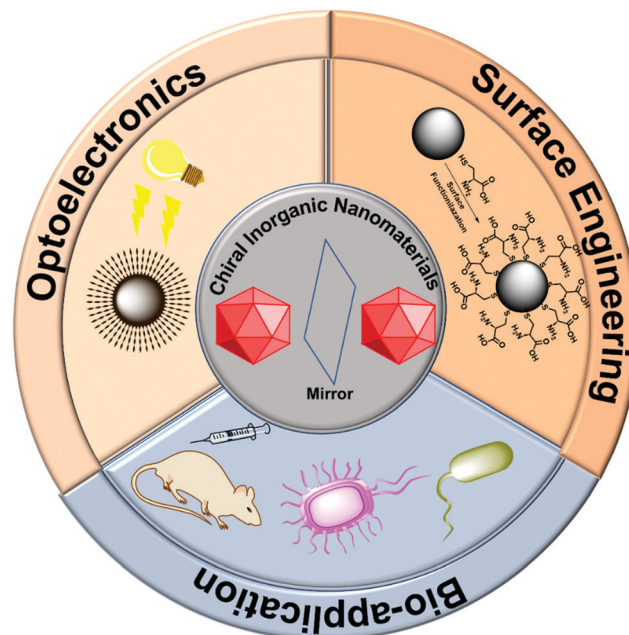


Fig. 1 A synopsis of the scope and applications of chiral inorganic nanomaterials, which is the key subject of the present review.

In this regard, a systematic theoretical description to comprehend the interaction between chiral/achiral ligands and metal surfaces can help to attain a rational design, preparation, and better understanding of the chirality imprint phenomena.^{57,58} Aliphatic or aromatic thiols or dithiols are known to be strongly adsorbed on Au surfaces, forming stable Au–S interfaces and, therefore, are one of the most widely used NP surface modifier ligands.^{59–63} Though many theoretical studies were made to predict the chirality transfer mechanisms and dynamics, the studies discussing enantioselectivity remain sparse.^{64–69} This void was nearly filled by the work from the Ortuño group who carried out computational simulations using density functional theory (DFT), providing a profound understanding of the enantioselective interaction at an atomic level. Their study included different forms of chiral amino acids such as L-cysteine (neutral, ionic, zwitterion) adsorbed on both the chiral enantiomeric Au facets, Au(321), Au(32 $\bar{1}$) and achiral Au(111), Au(110), Au(100) NPs surfaces.⁷⁰ In their theoretical framework, 4 atomic layers were considered for (111) and (110), 8 for (100), and 16 for (321) and (32 $\bar{1}$) facets. L-Cysteine growth adsorption energies on the multifacets Au NPs, E_{ads} were computed according eqn (2):

$$E_{\text{ads}} = E_{\text{S-L}} - E_{\text{S}} - nE_{\text{L}} \quad (2)$$

where $E_{\text{S-L}}$ denotes the energy of L-cysteine deposited on Au, E_{S} represents the energy of the Au NPs, nE_{L} signifies the energy of n no. of L-cysteine units. While the different forms of L-cysteine grafted on Au(111) are endothermic, the zwitterionic form A2, in which a negative charge resides on the sulfur linked to the Au(321) surface through S^- moieties, is the most stable form, having an adsorption energy of 0.58 eV (Fig. 2). From the quantum mechanical calculations, they concluded that the



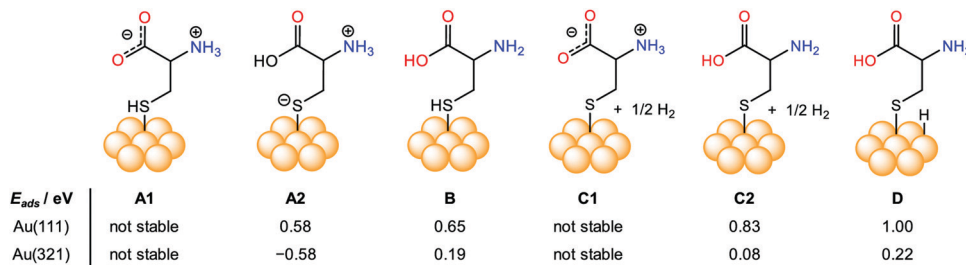


Fig. 2 Pictorial representation and comparison of the simulated surface adsorption energy, E_{ads} (eV), of different forms of intrinsic L-cysteine amino acid (neutral, ionic, zwitterion) adsorbed on various crystal facets, including achiral Au(111), and chiral Au(321). Reprinted with permission.⁷⁰ Copyright 2019, American Chemical Society.

adsorption energy of the chiral amino acid, L-cysteine on the Au(321) facet increased as compared to the case when adsorbed on the Au(321) at a relatively lower surface coverage. This observation can help to rationalize the design of enantioselective experiments at the nanoscale. However, in their investigation, the computational studies on D-cysteine were not performed. The authors concluded that the computational tool might have immense potential to elucidate many complexities associated with the origin of chirality.

Recently, Yang and co-workers employed a new strategy for chirality transfer in achiral Au NPs superlattices.⁷¹ The group utilized the chiral porphyrin-based supramolecular assembly and exploited the van der Waals interactions to induce chirality. It was also claimed to be the first report of its kind

on chirality transfer employing van der Waals interactions. Remarkably, it was observed that the chiral porphyrin supramolecular structure adopted J-type stacking and, upon metallization with zinc metal ion, changed to H-type stacking while maintaining its chirality. To induce the chirality in achiral Au NPs, first, the metallic NPs were functionalized with achiral *n*-octadecanethiol, resulting in core/shell heterostructures. Second, the chirality was introduced to the core/shell heterostructures by adding alkyl-terminated porphyrin or Zn-porphyrin solution. The presence of a new absorption band in the wavelength range of 500–800 nm in the UV-visible spectra (Fig. 3(a and c)) indicated the formation of two enantiomers such as (R)/(S)-PP-Au and (R)/(S)-ZnPP-Au assemblies. The additional spectral features appeared due to the extinction of

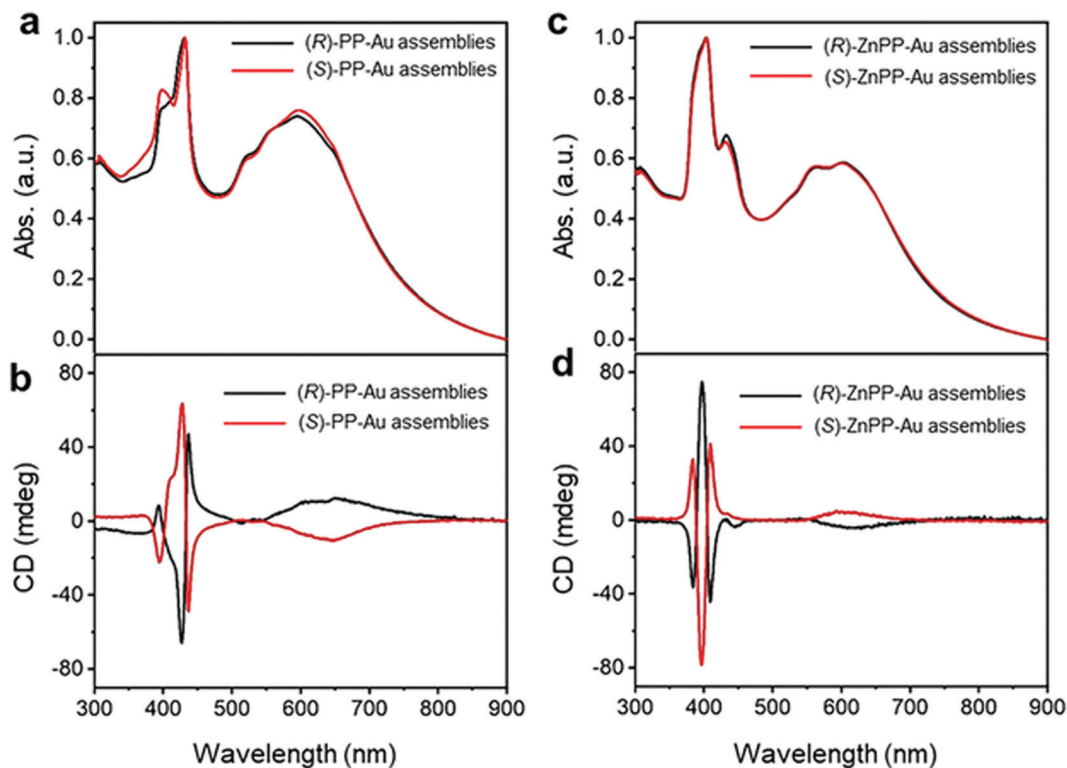


Fig. 3 (a) UV-Vis spectra recorded on (R)/(S)-PP-Au NPs, (c) (R)/(S)-ZnPP-Au NPs illustrating the appearance of new spectral features in the range of 500 to 800 nm. (b) CD spectra of (R)/(S)-PP-Au NPs, (d) (R)/(S)-ZnPP-Au NPs confirming the chirality transfer into metallic Au NPs. Reprinted with permission.⁷¹ Copyright 2021, Wiley-VCH.



Nam and co-workers have made an outstanding contribution towards imprinting the chirality in achiral metallic nanoparticles by embedding biomolecules and exploring optoelectronics,

Among the biologically important systems, dipeptides are one of the most relevant systems having intrinsic chirality suitable for use in chirality transfer reactions. The Nam group has employed this impressive methodology and considered two different dipeptides, gamma-glutamyl-cysteine (γ -Glu-Cys) and cysteinyl-glycine (Cys-Gly), to induce chirality in 3-dimensional (3D) metallic Au NPs.⁸⁷ The strategy is schematically illustrated in Fig. 6. As a result of chirality induction after the introduction of γ -Glu-Cys the Au NPs attained a cube-like morphology, while the Cys-Gly enforced a rhombic dodecahedron shape with elliptical cavities in each face. The SEM images show the reflection of surface-engineered Au NPs encoded with biomolecules and the resulting crystal facets (Fig. 7). To probe the origin of crystallographic chiral surfaces in Au NPs, 3D geometric models were employed, which entail the formation of chiral intermediates that drive the ultimate chirality. The present study demonstrates the dipeptide-mediated induction of the chirality in 3D Au NPs that can diversify their

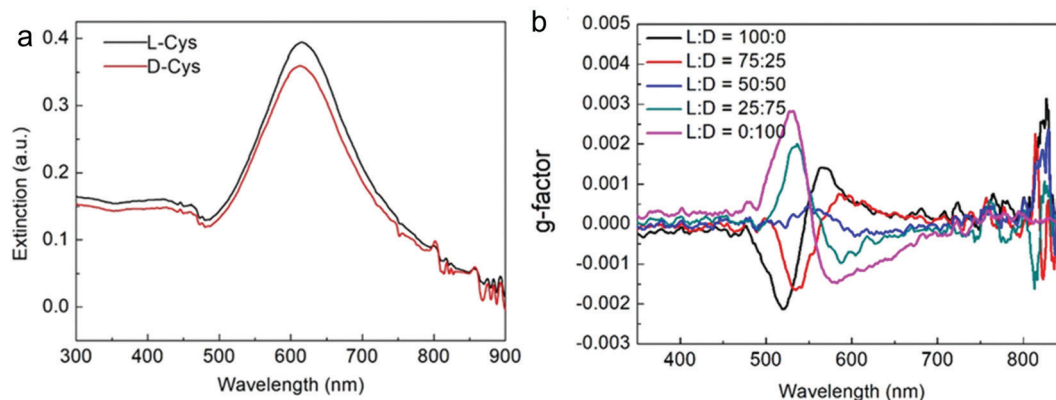


Fig. 4 (a) Absorbance spectra of L-Cys-Au NPs (solid black line) and D-Cys-Au NPs (solid red line). (b) The spectra affecting the g-factor of Au NPs recorded in different concentration ratios of L-Cys and D-Cys. Reprinted with permission.⁸⁸ Copyright 2020, Springer Nature.

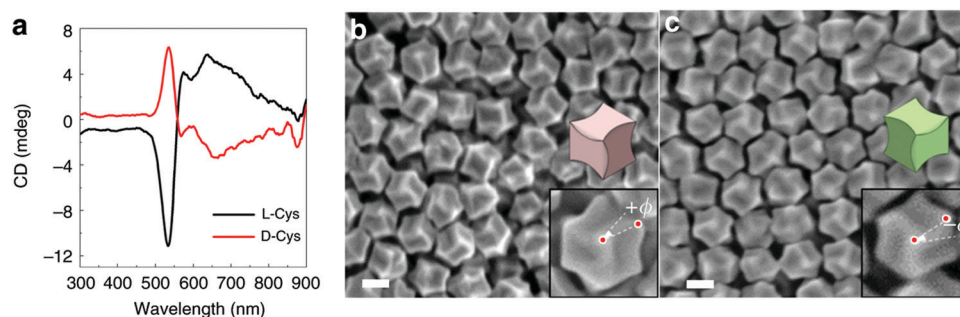


Fig. 5 (a) Visible-region CD spectra recorded on the L/D-cysteine-induced chiral metallic Au NPs. (b) SEM images of L-cysteine, and (c) D-cysteine induced chiral Au NPs. Reprinted with permission.⁸⁸ Copyright 2020, Springer Nature.

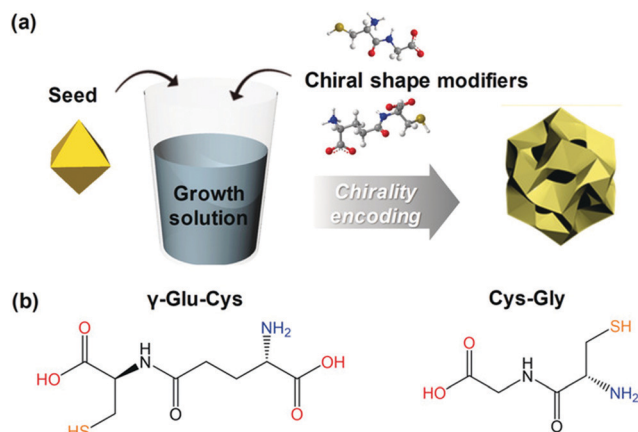


Fig. 6 (a) A schematic diagram indicating the preparation of chiral shape modifier-induced Au nanoparticles formation. (b) Chemical structures of the dipeptides are used as chirality inducers. Reprinted with permission.⁸⁷ Copyright 2020, Wiley-VCH.

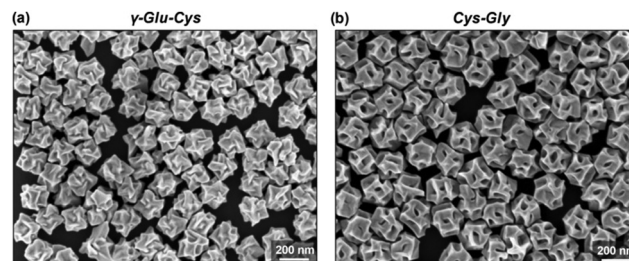


Fig. 7 (a) SEM images reveal crystal facets of chiral Au NPs generated by γ-Glu-Cys, and (b) Cys-Gly, respectively. Reprinted with permission.⁸⁷ Copyright 2020, Wiley-VCH.

applications in several ways. The examples discussed so far fall in the category that discusses the induction of chirality in noble metallic NPs. However, such induction of chirality can also be used with inorganic semiconductor NPs. There are only limited

examples of Chiral CdSe nanocrystals (NCs) that exhibit chiral-ligand-induced morphology and chiroptical behavior.^{89–94} He and co-workers could successfully modify the surface of CdSe-dot/CdS-rod NCs with the chiral amino acids such as L- and D-cysteine.⁹⁵ The chirality in semiconductors was induced *via* a ligand-exchange method where L- and D-cysteine produced L-/D-Cys-CdSe/CdS chiral quantum nanocrystals that display exceptional CD and CPL characteristics. The study showed a remarkable anisotropic factor (g_{lum}) equal to 4.66×10^{-4} in CPL spectra with positive and negative line shapes obtained for D- and L-Cys-CdSe/CdS NCs, respectively. The anisotropic factor



(g_{lum}) can be expressed as eqn 3:⁹⁶

$$g_{\text{lum}} = \frac{2(I_L - I_R)}{(I_L + I_R)} \quad (3)$$

I_L , I_R signify the corresponding intensities of the left- and right-handed CPL. Both the CD and CPL characteristics are significantly dependent on the concentration of chiral L/D cysteine, as at higher concentrations CD and CPL signals get suppressed. They further performed the DFT study and concluded that the orbital hybridization of either L- or D-cysteine with CdSe/CdS is the key factor of chirogenesis.

Cheng and co-workers utilized surface functionalization to prepare CdSe/CdS nanocrystals (NCs) of various morphologies including nanoflowers, tadpoles, *etc.* with precise shape control (Fig. 8).⁹¹ The detailed study provided a correlation between the shape and chirality induced by the ligand. In a typical procedure, CdSe/CdS NCs were first prepared followed by a post-synthesis ligand exchange reaction in aqueous media, NCs were modified with L-/D-cysteine. The UV-vis and photoluminescence (PL) spectral analysis of CdSe/CdS nanostructures were performed before and after ligand exchange. The formation of the CdS shell caused a red-shift in the absorption peak from 539 nm to 606 nm depending upon the morphology. Transmission electron microscopy (TEM) confirmed that the shape-directing agents successfully fine-tuned the morphology of the CdSe/CdS nanocrystals (Fig. 9). As can be seen from images (a–f), changing the concentration and nature of the ligand, nano-shapes such as nanoflowers, tadpoles, and dot/rods could be obtained. The obtained CdS shell phase was in agreement with the previous studies that suggest a wurtzite phase CdS growth when wurtzite CdSe seeds are used. It was observed that when the concentration was low or alkylphosphonic acid ligands were absent, the formation of spherical nanocrystals was favored

(Fig. 9a–c). To explain the chiroptical changes in various morphologies, a quantitative comparison of CD responses was performed. The CdSe core used to prepare different morphologies of NCs was from the same batch and has been subjected to identical ligand exchange reactions. This implies that any change in CD spectra could be linked to the change in the nanostructure's geometry. As shown in Fig. 10, nanoflowers, tadpole and long dot/rod (DR) shaped structures, which have relatively thicker shells (>1 nm) and large CdS volumes, exhibited CD signals corresponding to CdS only. Conversely, the CD signal for the CdSe core was still present in the short DR sample. It was also observed that the anisotropy factor is enhanced as the morphology changes from the nanoflower shape to the tadpole shape.

3. Optoelectronic applications

Optoelectronic devices are used to either modulate or detect light under an external electric field. Such devices enable photons to be used in the same fashion as electrons in an electronic device.^{97,98} Compared to conventional bulk systems, optical properties of nanomaterials benefit from a broad absorption window, higher quantum yields, narrow emission wavelengths, and fewer photo-bleaching risks, thanks to the strong light–matter interactions.^{99,100} Unlike chiral (bio)-organic materials, which usually contain a narrow absorption cross-section, display weak optical responses, and exhibit optical responses in the UV-region only, chiral plasmonic nanostructures are associated with stronger chiroptical responses and can interact with visible or near-infrared (NIR) light. These appealing features of chiral NPs significantly enhance their applicability in sensing, recognition, and optoelectronic devices. Avoiding any need for delicate (bio)molecular assembly, the Tsuma research group successfully constructed a system based on gold nanorods deposited on a TiO₂ surface in which handedness could be reversed *via* a photo-induced process.¹⁰¹ Interestingly, the chirality switch achieved by irradiating either with left or right-CPL could be maintained even in the absence of external stimuli. Sites with localized electric fields were then deposited with PbO₂ making use of the plasmon-induced redox process to yield chiral nanostructures. Precursor gold nanorods (Au NRs) were first synthesized and were then adsorbed onto a TiO₂ surface (film thickness ~60 nm) present on a glass plate coated with indium-tin-oxide. As shown in Fig. 11, characteristic transverse and longitudinal mode extinction peaks (550 and 815 nm, respectively) and the absence of CD signals was observed. An electrochemical cell composed of two compartments (Fig. 11c) was used to illuminate the photo-responsive ITO/TiO₂/AuNR-modified electrode in the presence of Pb(NO₃)₂ solution (50 mM) for 24 h. Illumination causes the formation of electron–hole (e–h) pairs at resonance sites on surface s of Au NRs. The holes formed around the resonance sites lead to the oxidation of Pb²⁺ and deposition of PbO₂. Concurrently, excited electrons from the electron–hole pair systems are transferred from the lower energy valence band (VB) to the higher energy

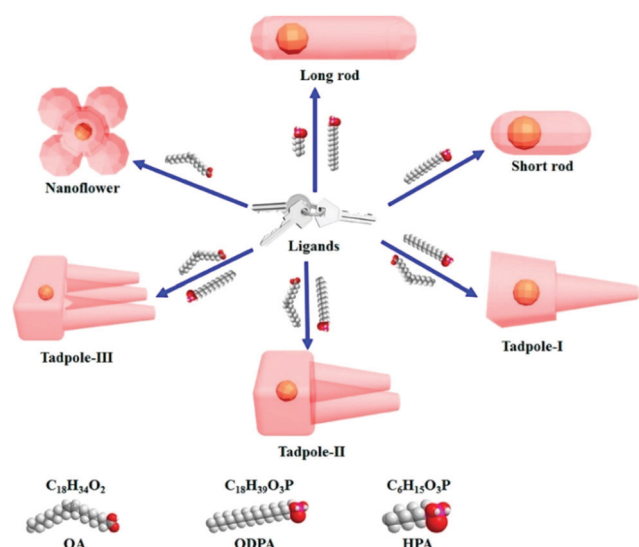


Fig. 8 A schematic diagram showing that chiral ligand-induced shape variations resulting in various anisotropic shapes can be achieved. Chemical structures of the chiral ligands are shown. Reprinted with permission.⁹¹ Copyright 2020, American Chemical Society.



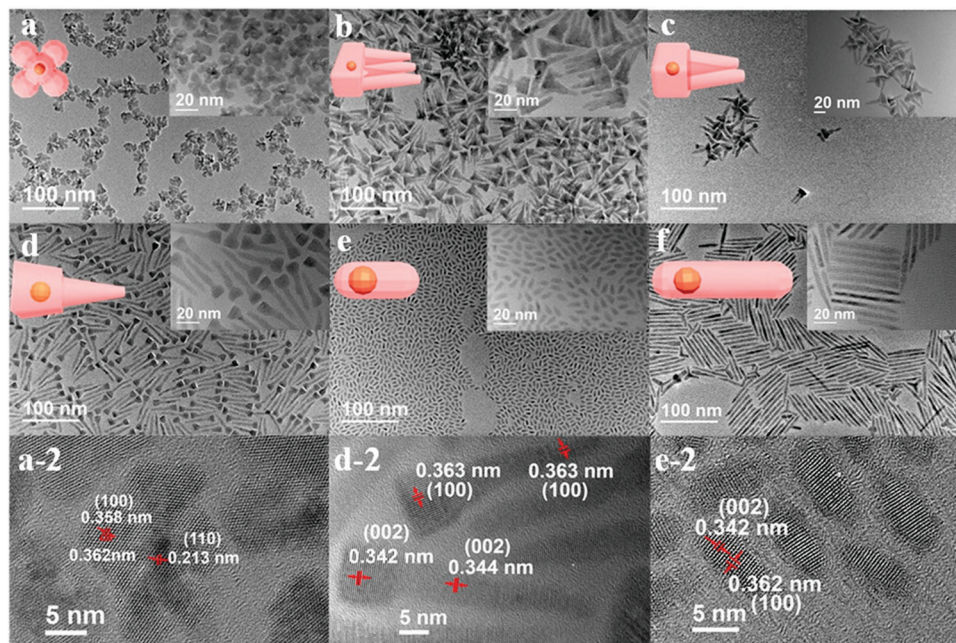


Fig. 9 (a–f) TEM images of CdSe/CdS NCs obtained in a variety of shapes including nanoflowers, Tadpole-III, Tadpole-II, Tadpole-I, short dot/rods, and long dot/rods. Reprinted with permission.⁹¹ Copyright 2020, American Chemical Society.

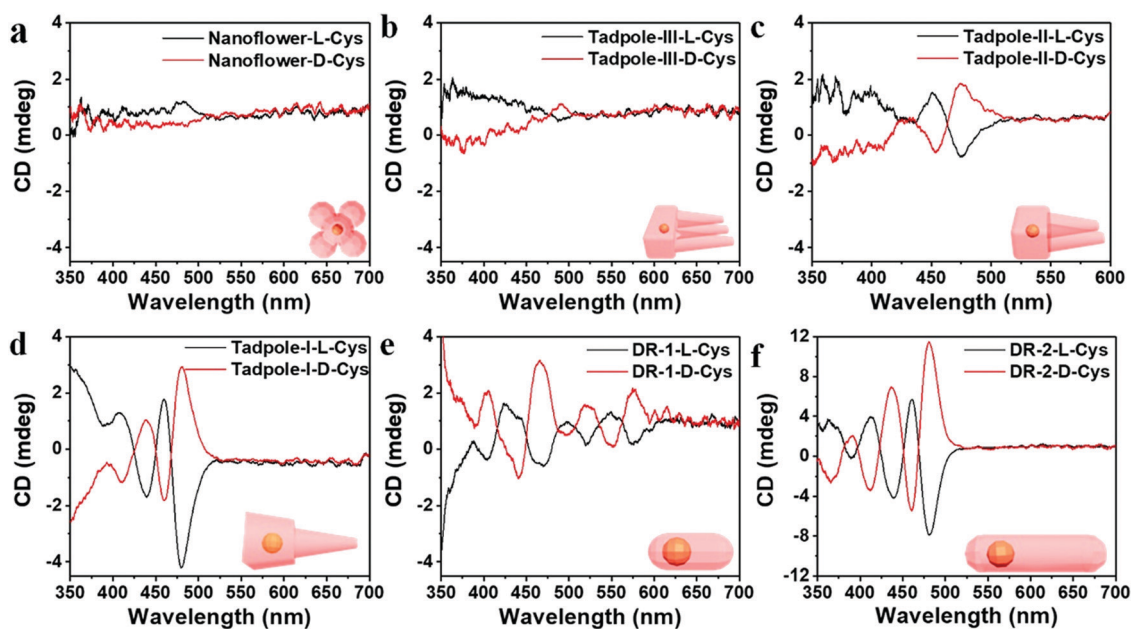


Fig. 10 (a–f) Mirror symmetry CD spectra of the L-/D-CdSe/CdS NCs having different shapes such as (a) nanoflowers, (b) Tadpole-III, (c) Tadpole-II, (d) Tadpole-I, (e) short dot/rods, and (f) long dot/rods. Reprinted with permission.⁹¹ Copyright 2020, American Chemical Society.

level conduction band (CB) of the TiO₂ substrate. Irradiating the system with left-CPL, a red shift in the extinction bands could be seen. CD spectra showed peaks at around 500–600 nm, 600–700 nm, and 750–900 nm. The CD spectral profile for the irradiation (L-CPL light) of AuNRs prepared using different aspect ratio exhibited similar CD signals. Scanning electron microscopic images (Fig. 11f and g) show the existence of one or two PbO₂

moieties on most AuNRs and the use of L-CPL and R-CPL produced mirror image geometries. The enantiomeric excess (ee) was found to be around ~50%. The authors also successfully performed the chiral switching in air medium. The device is highly reproducible and thus allows the reversible switching of chirality.

Circularly polarized luminescence and chirality are closely related properties. Materials that can detect CPL are of great

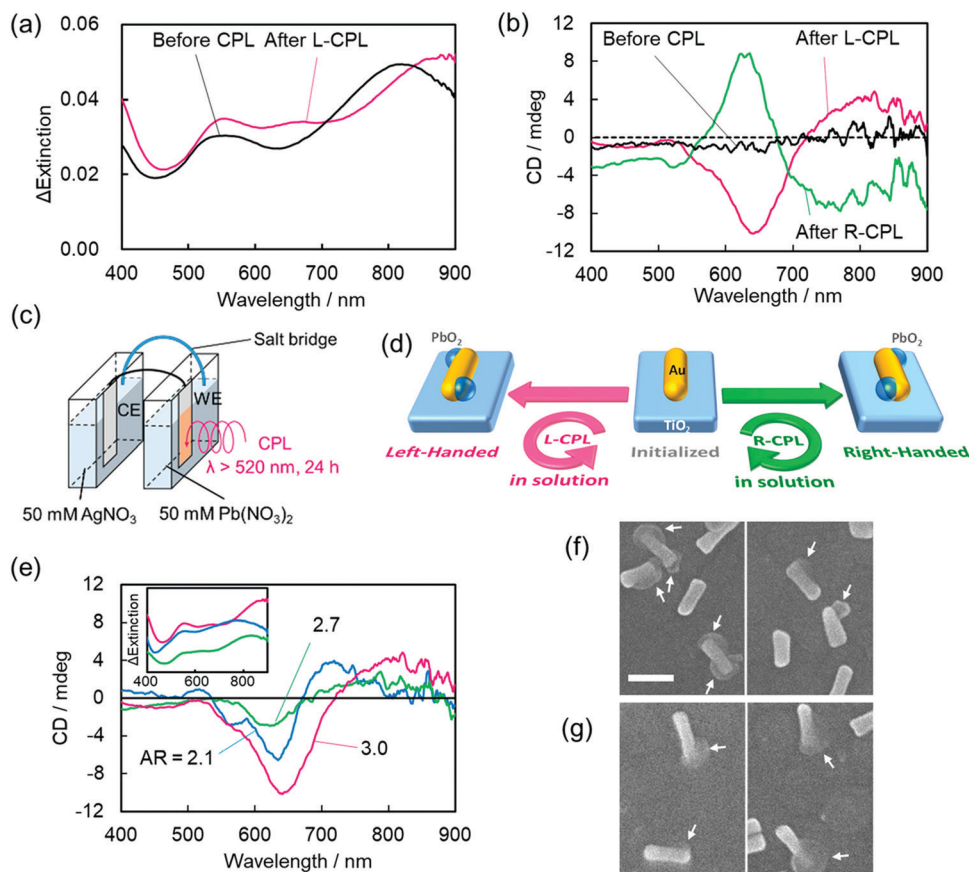


Fig. 11 (a) The change in extinction coefficient, (b) the visible region CD spectra of AuNRs deposited onto the TiO_2 substrate before and after the deposition of lead oxide by either right- or left-CPL irradiation. (c) An electrochemical cell used for PbO_2 deposition. (d) A schematic description of the device integration showing the formation of a chiral nanostructure. (e) CD signals of AuNRs after PbO_2 deposition, (f and g) the corresponding SEM images of AuNRs. Reprinted with permission.¹⁰¹ Copyright 2020, American Chemical Society.

importance for many applications at the nanoscale. Normally, the detector is coupled with the optical polarizer to achieve CPL detection. This requirement causes compromise with the sensitivity and resolution limits. Optically active systems interact differently with left or right-CPL and, therefore, display different absorption coefficients. Chiral metal halides are potentially important for the direct detection of CPL. Recently, a two-terminal device based on one-dimensional and quasi-two-dimensional perovskites has been demonstrated to be capable of direct photodetection of CPL.^{102,103} However these devices suffered from the need for high operational voltage and incapability of producing high output currents. One of the major issues associated with devices based on low-dimensional chiral metal halide semiconductors is low or sometimes even negligible conductivities/photoconductivity. Lu, Blackburn, and co-workers proposed a two-step strategy to circumvent this and prepared a device for the direct detection of CPL, which is comprised of chiral copper chloride and a carbon nanotube heterostructure.¹⁰⁴ In the first step, zero-dimensional (0D) chiral copper chloride hybrids (*R/S*-MBA)₂CuCl₄, where *R/S*-MBA stands for methylbenzylammonium, are prepared and characterized by CD spectroscopy. These hybrids are insulating in nature. Copper chloride hybrids thus formed are then

coupled with single-walled carbon nanotubes (SWCNTs). The heterostructure MBA₂CuCl₄/SWCNT helps in transducing the CPL signal into electrical output while maintaining high anisotropy. The functioning of this junction based on spin-selective interfacial charge transfer is hugely facilitated by the excellent carrier transport properties of the SWCNT. The complete process involves the absorption of CPL photons by MBA₂CuCl₄, followed by electron transfer to the SWCNT. This system could work at voltages as low as 0.01 V with an anisotropy factor as high as 0.21. To ensure an effective electric current response, it is important to have a highly conductive material interface. A 10 nm thick highly enriched (6,5) SWCNT was spin-coated with a chiral MBA₂CuCl₄ layer making a 50 nm thick heterojunction. Linear absorption spectra and CD measurements for SWCNT and chiral heterojunctions are shown in Fig. 12. Field-effect transistor (FET) measurement was performed to investigate the direction of charge transfer. When illuminated with 405 nm laser light a significant decrease in source-drain current and a negative shift in threshold voltage is observed (Fig. 13). For optoelectronic detection of left (L) and right (R) CPL, a set-up consisting of a laser diode source generating 405 nm continuous-wave L/R-CPL, a polariser, a neutral density filter and a quarter-wave plate was prepared



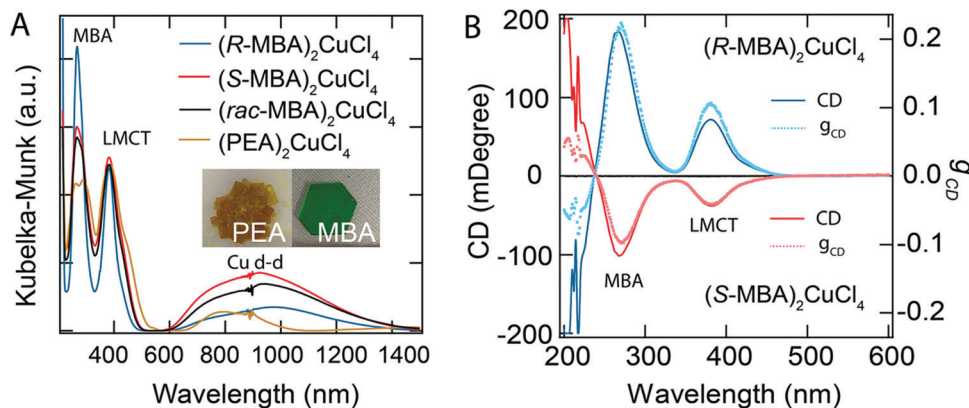


Fig. 12 (A) UV-vis-NIR spectra, and (B) CD spectra with the corresponding anisotropy factors of $(R-/S-/rac\text{-MBA})_2\text{CuCl}_4$ and $(\text{PEA})_2\text{CuCl}_4$. Optical images of $(\text{PEA})_2\text{CuCl}_4$ and $(R\text{-MBA})_2\text{CuCl}_4$ crystals are shown in the inset. Reprinted with permission.¹⁰⁴ Copyright 2021, American Chemical Society.

under an inert atmosphere (Fig. 14a). The detection system was calibrated to generate identical excitation L/R-CPL intensities. The CPL detection experiment for the heterojunctions of $R\text{-MBA}_2\text{CuCl}_4/\text{SWCNT}$ exhibited a clear difference in photocurrent responses for right and left-handed CPL light. The photocurrent response (I_{ph}) as a function of time (s) under applied bias for R-CPL was found to be larger than that for L-CPL (Fig. 14b and c). When $S\text{-MBA}_2\text{CuCl}_4/\text{SWCNT}$ was tested, the photocurrent anisotropy was reversed.

In the recent past, molecular spintronics has seen immense activity. A spintronic device utilizes both the spin and charge of the electron. Molecular chirality can be used in spintronic-based devices to control the spin-selective charge transport studies; however, the mechanism by which spin and chirality regulate the charge transport is still not fully understood.^{105,106}

Findings from the Faraday rotation experiments suggested that spin-polarized excitons could be generated if circularized polarized light is used to photoexcite the quantum dots (QDs).¹⁰⁷ Waldeck and co-workers investigated the profound effect that chiral QDs can exhibit spin-selective charge transport phenomena using molecular-scale spintronic devices.¹⁰⁸ Chiral CdSe QDs were employed as electron acceptors templated on silica surfaces (SiO_2 microbead, Fig. 15). Upon illumination with circularly polarized light, the QDs initiated the electron transfer. Corresponding electron-transfer rates for left and right CP light photoexcitation were measured and then used to calculate the electron-transfer rate asymmetry (P_{et}) and a 15-fold difference was observed. As can be seen in the scheme depicted above, the acceptor QD is ligand exchanged with cysteine making it chiral, whereas the donor QD is left achiral.

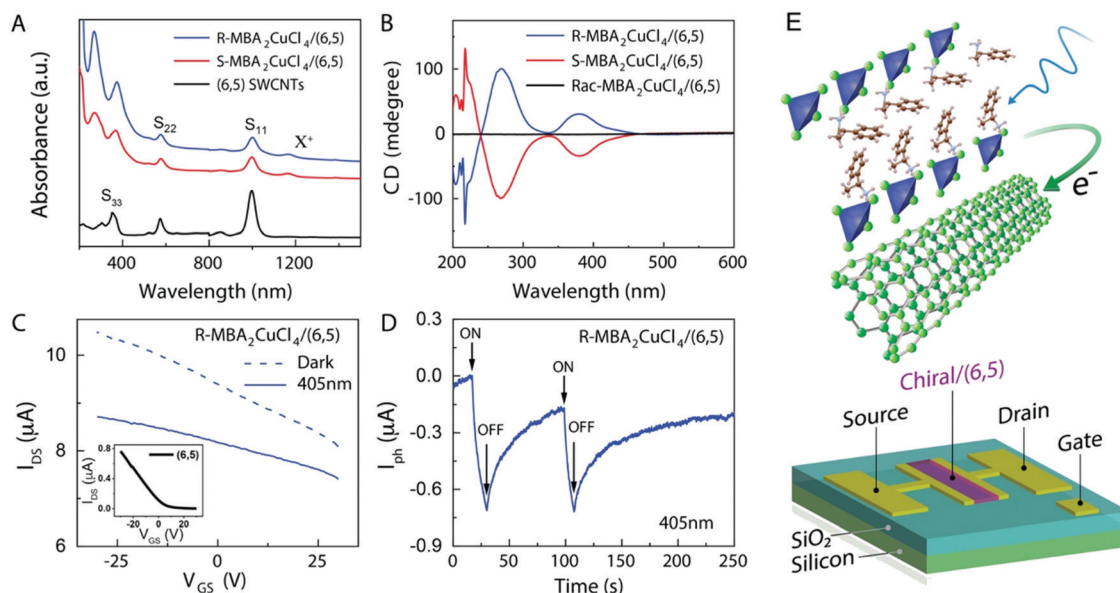


Fig. 13 (A) UV-Vis-NIR absorption spectra of SWNT, R - and S - $\text{MBA}_2\text{CuCl}_4/\text{SWCNT}$. (B) CD signals of R - and S - $\text{MBA}_2\text{CuCl}_4/(6,5)$ SWCNT and $(rac\text{-MBA})_2\text{CuCl}_4/(6,5)$ SWCNT. (C) A comparison of the effects of dark and photoresponse conditions on the FET transfer curve. (D) The photocurrent response of $(R\text{-MBA})_2\text{CuCl}_4/\text{SWCNT}$ over time. (E) A schematic description of a three-terminal FET device. Reprinted with permission.¹⁰⁴ Copyright 2021, American Chemical Society.

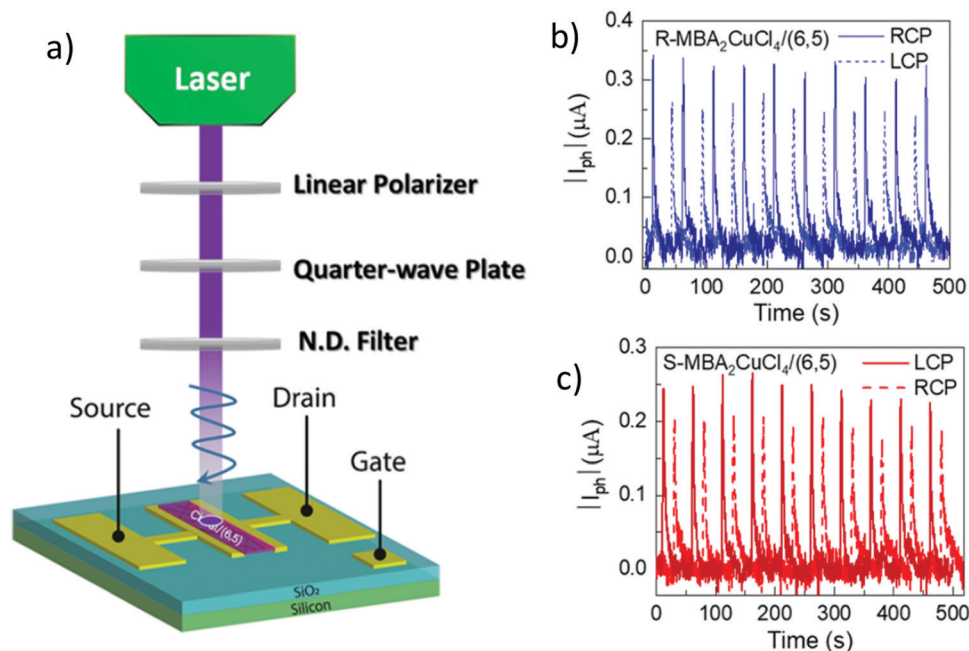


Fig. 14 (a) Schematic description of an experimental set-up used for creating left and right CPL. (b) Photocurrent (μA) variation as a function of time (s) using $R\text{-MBA}_2\text{CuCl}_4/\text{SWCNT}$, and (c) $S\text{-MBA}_2\text{CuCl}_4/\text{SWCNT}$ when illuminated with a 405 nm pulsed laser light under $V_{\text{DS}} = 2$ V. Reprinted with permission.¹⁰⁴ Copyright 2021, American Chemical Society.

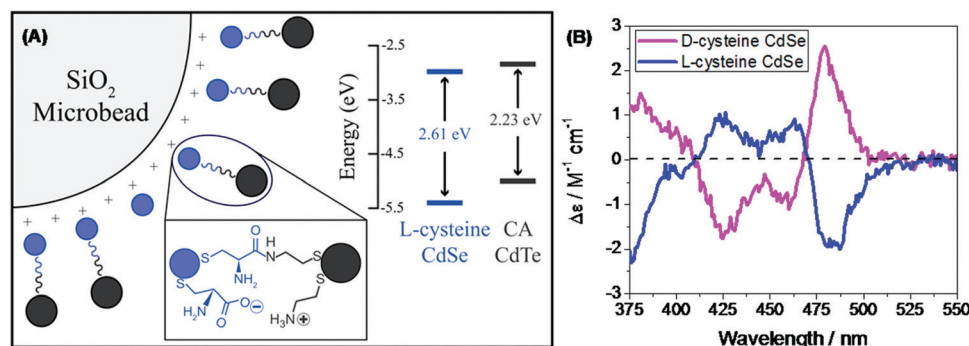


Fig. 15 (A) Schematic showing achiral donor CdTe QDs and the surface-functionalized chiral acceptor CdSe QD dyad assembled over a silica bead. The inset shows the energy levels of donor and acceptor QDs. (B) Mirror symmetry CD signals of the L-/D-Cysteine functionalized acceptor QD displaying opposite trends. Reprinted with permission.¹⁰⁸ Copyright © 2017, American Chemical Society.

The CD spectra, which show a very distinctive band around 475 nm, indicated that ligand chirality is imprinted on QD (Fig. 15B). The fluorescence lifetime for QD dyad assemblies containing achiral donor QD and acceptor QD (ligand exchanged with chiral capping ligand *L*-cysteine/*D*-cysteine or achiral ligand mercaptopropionic acid) were studied using excitation light as linear, clockwise (CW), and counterclockwise (CCW) circularly polarized forms (Fig. 16). It was found that when polarized light was used, the lifetime decay profile remained almost unchanged for the achiral acceptor and changed significantly for the chiral acceptor dyad assembly. For the *D*-cysteine functionalized acceptor QD, the decay was fastest when excitation light was CW-CPL and slowest when CCW-CPL was used. The *L*-cysteine capped acceptor QD dyad

assembly showed the opposite behavior. The authors further demonstrated that asymmetry in electron transfer rates (P_{et}) is not only influenced by the chirality of the capping ligand but also depends on the QD exciton's CD intensity. The results confirmed that in quantum dyads comprised of chiral acceptor QD and achiral donor QD, electron transfer varies with the direction of circularly polarised excitation light and acceptor chirality. This work unequivocally demonstrates that not only bio-molecules but also chiral QDs can show spin-selective electron transfer processes supporting the chirality-induced spin selectivity (CISS) phenomena.^{33,109–111}

Hybrid perovskites, where a cationic methylammonium is incorporated within the metal-halide structure, have attracted great attention for their applications in solar cells and other



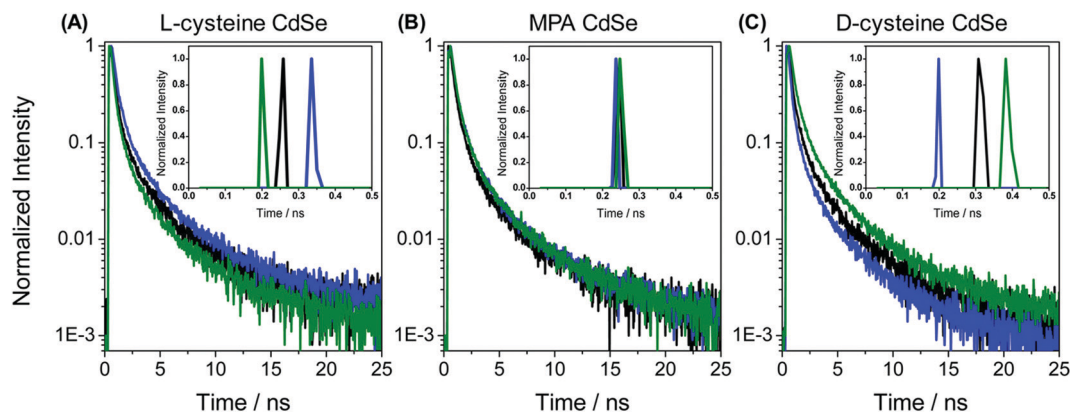


Fig. 16 (A–C) Lifetime decay profiles for acceptor CdSe QD functionalized with chiral L-cysteine, achiral mercaptopropionic acid, and chiral D-cysteine, respectively. Data represented in black, blue, and green correspond to the linear, clockwise, and counter-clockwise circularly polarized excitation light, respectively. Insets show the short parts of the lifetime distributions. Reprinted with permission.¹⁰⁸ Copyright 2017, American Chemical Society.

organic electronic devices.¹¹² In this context, a lack of inversion symmetry and an additional feature of optical activity in chiral systems, particularly in hybrid perovskites obtained by using chiral organic cations, make them an apt system for spin-optoelectronic device fabrications. A recent perspective article by Wei and Ning discusses the design and application of chiral perovskites towards spin-optoelectronic and spintronic devices.¹¹³ Although chiral perovskites have been applied in spin-optoelectronic systems and there has been an increase in such devices, the theoretical aspects behind the work principle and factors affecting its efficiency have not been explored much.^{114–117} In this regard, Ning and co-workers performed theoretical calculations on a Pb–I-based structure and studied the impact on the spin filtering property of the system by changing halogen substitution on the chiral part, which is methylbenzylammonium (MBA), of the hybrid perovskite.¹¹⁸ In their study, they focused on the structure of chiral molecules responsible for inducing chirality in the hybrid perovskite. Following experimental results from $(R\text{-MBA})_2\text{PbBr}_4$ that indicated an increase in CD intensity upon halogen substitution at the *para*-position, the Ning group first studied the hybrid perovskite systems obtained by substituting the *para*-position H-atom of *S*-MBA with F, Cl, and Br, and then substituting the H-atom that is near to the chiral carbon of *S*-MBA, respectively. Their investigation revealed the enhancement of the spin splitting energy from 13 meV to 73, 90, and 105 meV when the H-atom was substituted with F, Cl, and Br, respectively. This indicates the possibility of significant modulation of the output in spin-optoelectronic devices based on the chiral perovskites.

So far, we have discussed the induction of chirality mostly through the surface functionalization of the nanoparticles. We know that it is also possible to induce plasmonic chirality in the nanoparticles from the chiral molecules adsorbed on the surface through a dipole–dipole interaction so, in some cases, chiral induction could be achieved even in the absence of any chiral adsorbate.¹⁴ An interesting review by Liz-Marzán and co-workers discussed the recent progress in the templated assembly of achiral plasmonic nanoparticles exhibiting chiroptical activities.¹¹⁹ In an impressive example of utilizing supramolecular

nanostructures to build a variety of nanoparticle assemblies in the cylindrical confinements, Xu and co-workers successfully prepared nanoparticle stacked-rings and helical nano-patterns.¹²⁰ These assemblies displayed a chiral plasmonic response that was superior to the original material. A supramolecular assembly consisting of polystyrene-*block*-poly(4-vinylpyridine) with 3-*n*-pentadecylphenol was mixed with 9 vol% of dodecanethiol-capped gold nanoparticles of 4 nm. The mixture was drop-casted on the 35 nm Al_2O_3 membrane and annealed at 110 °C. The supramolecular part formed cylindrical hexagonal structures that were used to form one-dimensional nanowires of Au-NPs (Fig. 17). As can be seen in the in-plane TEM image (Fig. 17b), Au-NP forms a single helix having a pitch between 20 to 50 nm. By modifying the NP loading, the assembly pattern could be varied to yield stacked rings or single/double-helical structures. Nanoparticle helices generally display chiroptical activity in the 500–600 nm range. Using dark field-particle CD spectroscopy, the optical activity of a helical NP assembly was measured. A designated dip-peak was observed at around 800–1000 nm with a dissymmetry factor of 0.31 at $\lambda = 895$ nm, and 0.17 at $\lambda = 960$ nm. Near-infrared (NIR) chiral activity is highly promising towards the bimolecular application of nanoparticle-based devices.

Lin and co-workers investigated the formation of chiral assemblies of nanocomposites comprised of the achiral block copolymer and nanoparticles.¹²¹ They used theoretical calculations to get the predicted chirality and CD responses of the nanoparticle assembly in block copolymer nanochannels and then

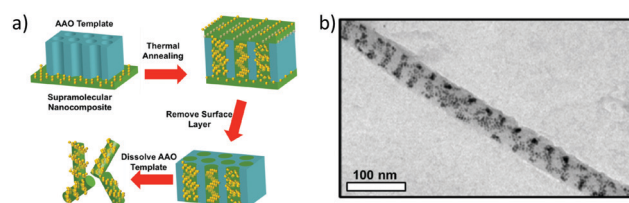
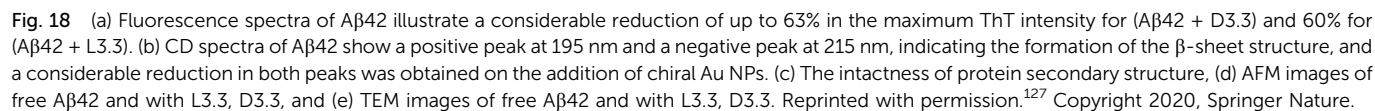


Fig. 17 (a) Schematic diagram showing the steps leading to the formation of Au-nanowires; (b) TEM image of a single-helical nanowire. Reprinted with permission.¹²⁰ Copyright 2017, American Chemical Society.



The effective approach to uplifting the treatment of Alzheimer's disease (AD) is by precluding the aggression of amyloid-beta (A β). In this context, Tang and co-workers synthesized the achiral Au NPs, followed by the induction of chirality by functionalization with the chiral 3.3 nm L- and D-glutathione (GSH) (abbreviated as L3.3 and D3.3 respectively).¹²⁷ Chiral Au NPs have been observed to enantioselectively curb the A β aggression and permeate the blood-brain-barrier (BBB) through several GSH channels that reside in the brain, resulting in the treatment of AD. The thioflavin T (ThT) fluorescence assay elucidated the free L- or D-GSH, indicating the sparse hindrance on A β 42 fibrillization.

Due to the exciting optical features and biocompatibility, chiral inorganic NPs have emerged as a tremendously efficient and



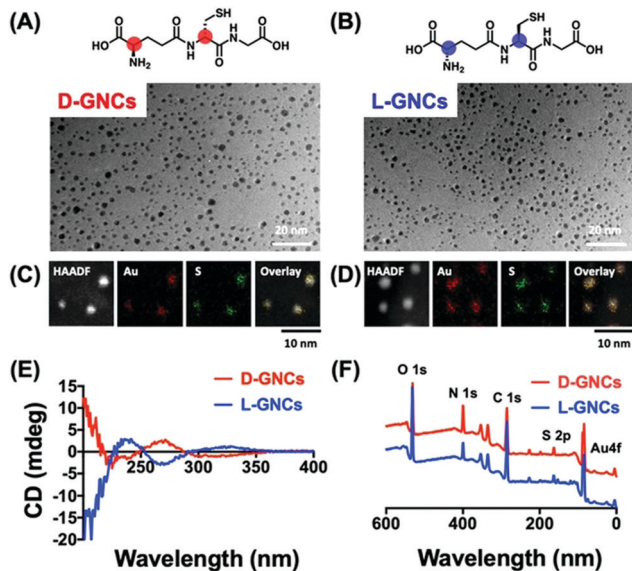


Fig. 20 (A and B) Chemical structure of chiral ligand D-/L-glutathione, (C and D) TEM images, and the corresponding elemental mapping analysis of the corresponding GNC's. (E) and (F) show the characterization of D-/L-GNCs by CD and X-ray photoelectron spectroscopy (XPS). Reprinted with permission.¹³⁴ Copyright 2021, Wiley-VCH.

long retention time increases the circulation time in the blood and hence leads to a broader organ distribution of GNCs. The study demonstrated that by utilizing surface functionalization, the nanomaterials' chirality could be reversed and their biosafety and *in vivo* performance could be drastically improved.

4. Conclusion and outlook

In summary, benefiting from controllable nanoscale size, large surface area, morphology, and quantum confinement, the metallic or semiconductor NPs have manifested excellent performance in various aspects of optics, physics, material science, and life science. The application of nanomaterials becomes more widespread when chirality is induced in achiral nanomaterials or intrinsic chiral nanomaterials are employed. Small chiral organic ligands are employed to induce chirality in achiral NPs, which act as symmetry breakers. Chiral nanomaterials possess intriguing optical features represented by circular dichroism, circular plasmonic luminescence, *etc.* The recent advances in this area have revealed that besides their optoelectronic device applications, chiral NPs are a boon towards the fields of disease diagnosis and therapies. The chiral NPs have been proved to be excellent candidates for the treatment of Alzheimer's Disease. In addition, the 3D chiral Au NRs-protein hybrid structures can act as a rich source for amyloid detection and expand the scope for the implication of template-driven plasmon chirality. Designing capable and influential antibacterial nano-scale agents that can cure bacterial infection is now possible due to the availability of such NPs. The future scope of this line of research is to study the applied mechanisms demonstrated by the chiral NPs in detail

to comprehend the micro-steps and variations. The premise of this research is to evolve the application of this phenomenon and boost its activity for better results. Though the research progress made with the chiral NPs is laudable, this domain is still in the nascent phase, as the systematic understanding of chiral imprinting is crucial. Therefore, more experimental and theoretical studies are needed. During the process of nanoimprinting by chiral inducers, the formation of the cluster may occur instead of distinct NPs development. To prohibit such cluster formation, the concentration of the corresponding metal salt solution can be diluted. This induced chirality study has been experimentally performed on selected metals and semiconductors with mostly L and D cysteine amino acids. Thus, more experimental and theoretical studies are welcome with various other metals such as Cu, Ni, Fe, using chiral inducers, to understand their growth mechanisms. To make them fruitful bio-applications, more such studies are also necessary using such chirality-induced NPs. There is relatively little known about the magnetic field effect on chiral NPs, which is crucial for the rational design and construction of nanoscale spintronic devices. It seems that this domain is quite challenging because all the ferromagnetic metals (Fe, Co, Ni) are prone to oxidation during the formation of NPs. A feasible solution could be the electrochemical reduction of such metal salts followed by capping with such chiral ligands. Furthermore, this concept can be deployed to build heterostructures where at least two different NPs can be assembled. Since semiconducting NPs can conduct electricity and display good charge carrier concentration, NPs can also be applied in making electrode materials for devices. However, scalability, reproducibility, and operational aspects can be of major concern and would need to be resolved.

Author contributions

PCM: conceptualization; funding acquisition; investigation; supervision; writing – original draft. DA: writing – original draft. RKP: visualization; writing – review and editing. SJ: visualization; writing – review and editing.

Conflicts of interest

There are no conflicts to declare.

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