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Resonant Plasmonic-Biomolecular Chiral Interactions in the Far-Ultraviolet: Enantiomeric Discrimination of sub-10 nm Amino Acid Films

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ABSTRACT: Resonant plasmonic-molecular chiral interactions are a promising route to enhanced biosensing. However, biomolecular optical activity primarily exists in the far-ultraviolet regime, posing significant challenges for spectral overlap with current nano-optical platforms. We demonstrate experimentally and computationally the enhanced chiral sensing of a resonant plasmonic-biomolecular system operating in the far-UV. We develop a full-wave model of biomolecular films on Al gammadion arrays using experimentally derived chirality parameters. Our calculations show that detectable enhancements in the chiroptical signals from small amounts of biomolecules are possible only when tight spectral overlap exists between the plasmonic and

biomolecular chiral responses. We support this conclusion experimentally by using Al gammadion arrays to enantiomerically discriminate ultrathin (< 10 nm thick) films of Tyrosine. Notably, the chiroptical signals of the bare films were within instrumental noise. Our results demonstrate the importance of using far-UV active metasurfaces for enhancing natural optical activity.

Since the seminal works on optical chirality and superchiral light by Tang and Cohen,^{1,2} researchers have employed a myriad of nano-optics approaches aimed at amplifying the intrinsically small effects^{1,3} of natural optical activity (OA) – circular dichroism (CD) and birefringence (CB).⁴ Nanostructures can create electromagnetic near-fields with optical chirality density higher than circularly-polarized light (CPL);⁵ thus increasing the CD from chiral molecules in these locations.¹ Initially, most nano-optics studies consisted of plasmonic nanoparticles and nanoarrays.⁶ More recently, there has been growing interest in developing⁷ chiroptical dielectric platforms,^{8–11} owing in part to their ability to sustain both electric and magnetic resonances, which, when properly coupled, can lead to high optical chirality densities.¹² Unfortunately, these resonances mainly arise in the visible to the infrared (IR), leading to great spectral mismatches with natural OA.⁷ Most chiral analytes of interest (such as biomolecules) have absorption bands in the ultraviolet (UV).¹³ In this context, low loss high index dielectric materials are still limited for the UV region,^{14–16} especially in the far-UV regime ($\lambda < 250$ nm) where CD signals are used to assess biomolecular structure.¹³

As for plasmonics, achiral nanostructures have been the most thoroughly investigated.¹⁷ Assembling achiral metallic nanoparticles into chiral superstructures^{18–23} can lead to strong CD signals from dipole-dipole interactions among neighboring particles.²⁴ Furthermore, the presence of chiral molecules close to achiral metallic nanostructures can induce chiroptical signals at the plasmon frequency.^{25–29} The use of achiral nanostructures is motivated by a desire to avoid the

strong CD signals from their chiral counterparts which could overshadow the weak molecular chiroptical response^{7,17,30–33} or require data post-processing to assess the chiral analyte signal.^{10,17,32} Conversely, chiral plasmonic nanostructures can generate high optical chirality in their near fields.^{5,7,33,34} Previous works which utilized chiral plasmonic metasurfaces relied on the differences in the spectral shift induced by a chiral analyte when interacting with a pair of enantiomeric arrays^{35,36} or on the sum of the plasmonic chiroptical signal in the presence and absence of the chiral analyte.³⁷ The drawback of these approaches is that nanofabrication heterogeneities³² can perturb the perfectly mirror-symmetric spectrum of the enantiomorph arrays. Furthermore, regardless of the platform, the energy mismatch between natural optical activity and metamaterial resonances remains the most significant obstacle for chiral sensing.³³

The benefits of overlapping molecular OA and achiral nano-optical resonances were previously studied.^{11,38–40} Notably, detuning of the molecular and plasmonic resonances was detrimental to the observed CD signals.^{38,40} Experimentally,^{11,40} the evaluation of these resonant interactions was permitted by conjugating DNA to chromophores, where an induced OA in the visible was enabled by the chiral macromolecule.^{41,42} Therefore, the natural OA of the DNA molecule was not directly assessed. In this work, we study the resonant chiral interactions between biomolecules and plasmonic near-fields in the far-UV by means of electromagnetic calculations and experiments. To the best of our knowledge, this is the first time a platform enabling spectral overlap between biomolecular and plasmonic chiral resonances is used to assess natural OA directly. We determine the handedness of the analyte by analyzing the fingerprint-like chiral signature of the plasmonic-biomolecular system. Our biosensing scheme enables the enantiomeric discrimination of ultrathin (< 10 nm thick) amino acid films. Notably, the CB signal of the bare biomolecular films could not be reliably resolved by our instrument. Our work provides a critical step forward in understanding

resonant plasmonic-biomolecular interactions and developing metasurfaces for enhanced chiral biosensing.

Aluminum is a promising metal for UV plasmonics,^{43–45} displaying excellent optical properties in the UV, especially when low base pressures and fast deposition rates are employed.⁴⁶ Additionally, its passivating oxide layer promotes the stability of Al nanostructures to air and organic solvents.⁴⁷

To better understand resonant plasmonic-biomolecular chiral-chiral interactions, we start by performing full-wave electromagnetic (EM) simulations of Al gammadion arrays coated with 20 nm thick achiral and chiral biomolecular films (Figure 1a). The latter were implemented in the finite-element solver JCMsuite (JCMwave GmbH, Germany) with bi-isotropic constitutive relations,^{48,49} and initially modeled as a homogeneous material with index of refraction $n = 2.045 + 0.230i$, and chirality parameter $\kappa = 0.004$ or $0.004i$. For purely real ($\text{Im}\{\kappa\} = 0$) and imaginary ($\text{Re}\{\kappa\} = 0$) chirality parameters, the film only displays CB and CD, respectively^{50–52} (Figure 1b). To verify the effects of the chiral medium on the plasmonic-biomolecular system, we use as reference the chiroptical signals of the nanostructures coated with a racemic film,³² which exhibit no net chirality, i.e., $\kappa = 0$. The nanostructures had a total thickness of 105 nm (5 nm Ti and 100 nm Al) and were on top of a fused silica substrate (see Figure S1 for dimensions).

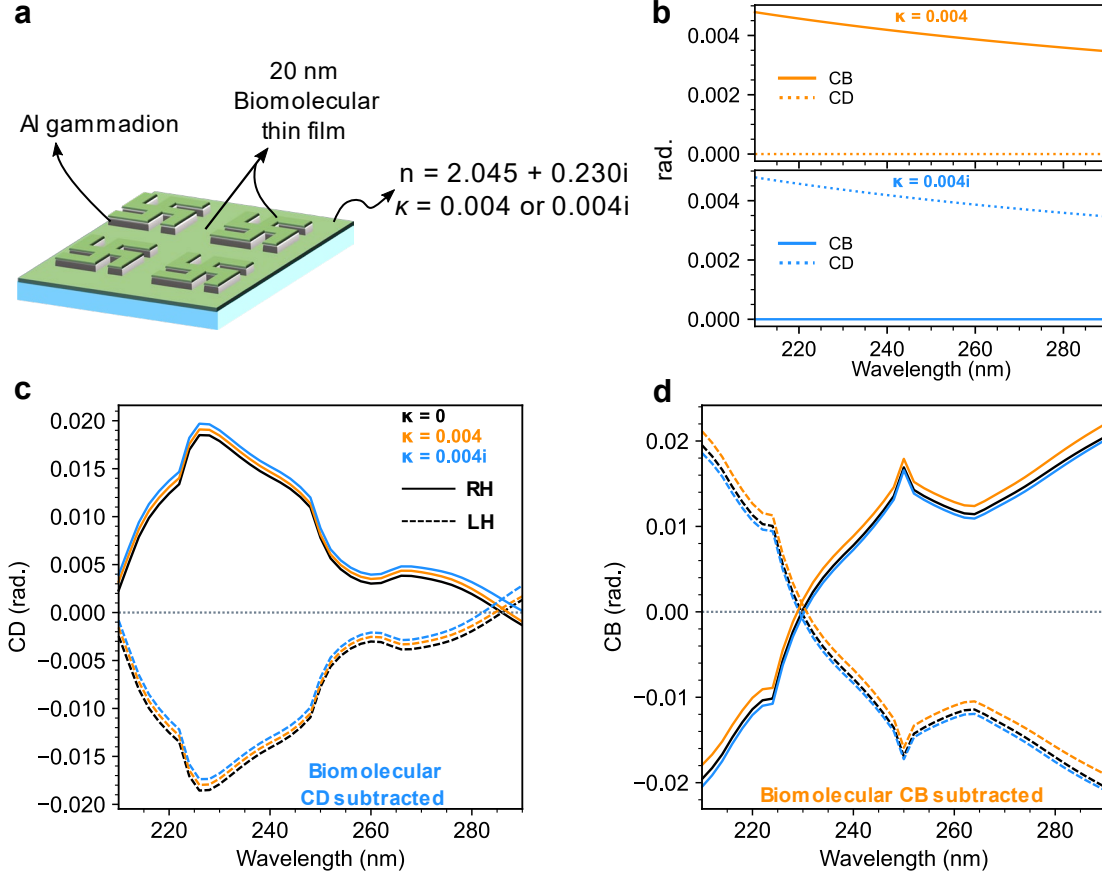


Figure 1. Simulated far-field optical response of the UV plasmonic-biomolecular chiral system.

(a) Schematic of a LH Al gammadion array coated with 20 nm of biomolecular layers, which we initially model with constant optical properties. (b) For chirality parameters $\kappa = 0.004$ and $0.004i$, the molecular layers display only CB and CD, respectively. (c,d) Effects of biomolecular chirality on the plasmonic chiroptical response. When the films are racemic, i.e., achiral, the (c) CD and (d) CB signals of the RH (solid black) and LH (dashed black) gammadions display the expected mirror symmetry. Conversely, when the films are chiral, this mirror symmetry is perturbed. In (c), even though the biomolecular κ is purely real, it asymmetrically affects the CD of the system when it is composed of RH (solid orange) and LH (dashed orange) gammadions. The lack of mirror symmetry is more pronounced when $\text{Im}\{\kappa\} \neq 0$, i.e., when both biomolecules and plasmonic structure have a CD response (solid and dashed blue lines). Note here, that we subtracted the CD

signal arising from the bare biomolecular film (the observed differences are not simply an additive effect). (d) The same remarks are also valid for the CB signal of the plasmonic-biomolecular system.

As expected, when the right- (RH, solid line) and left-handed (LH, dashed line) gammadion arrays are coated by racemic layers (black curves), their CD (Figure 1c) and CB (Figure 1d) spectra are mirror images of each other. The effects of the biomolecular chirality are displayed in the orange and blue curves of Figure 1c and d. Notice that we subtracted the CD and CB signals arising from the isolated 20 nm biomolecular films from the relevant spectra presented in Figure 1c and d, respectively. These plots show that the mirror symmetry among the chiroptical signals of the coated RH and LH gammadions is broken when the molecular film is chiral. These changes are most noticeable near the crossing point of the two enantiomers, i.e., at 285 nm for Figure 1c and at 230 nm for Figure 1d. In Figure 1c, even though the chiral medium has no CD of its own, it asymmetrically perturbs the electromagnetic response of the RH (solid orange) and LH (dashed orange) nanostructures to CPL. $\text{Re}\{\kappa\}$ can modify the CD response of plasmonic-biomolecular systems.⁵³ When the film has a purely imaginary κ , the lack of mirror symmetry is more substantial (Figure 1c, blue curves), even after considering the background biomolecular chiroptical signal. The same observations are valid for the CB signals (Figure 1d). For real biomolecules, with dispersive n and κ , we expect to see the largest deviations from the reference spectra, when the CD (or CB) response of the plasmonic nanostructure spectrally overlaps with the respective chiral signal of the biomolecule. Recently, Both *et al.*⁵³ theoretically described the multiple mechanisms contributing to the CD signal of chiral plasmonic nanostructures interacting with a chiral medium.

We expect the same effects to be present in our work because both nanostructure and biomolecular films are chiral. An exhaustive mechanistic evaluation is, however, beyond the scope of this work.

To better understand our resonant plasmonic-biomolecular system we modified our simulation to account for dispersion in the chiral parameter of real materials, using Tyrosine (Tyr) as our model biomolecule. We extracted $\text{Im}\{\kappa\}$ from the CD and $\text{Re}\{\kappa\}$ from the CB signal of a 90 nm thin film of L-Tyr on top of a fused silica substrate. Figure 2a,b displays the experimental and numerical values for CD and CB, respectively (See Methods in Supporting Information for details). To simulate D-Tyr, we flipped the sign of κ determined for the L- enantiomer. We note that at resonance (Figure 2c), the highest absolute value for $\text{Re}\{\kappa\}$ and $\text{Im}\{\kappa\}$ of L-Tyr are approximately 1×10^{-3} and 7×10^{-4} , respectively (See Supporting Information for tabulated data). At 310 nm, which is still close to the resonant frequencies, $\text{Re}\{\kappa\}$ quickly decays to $\sim 4 \times 10^{-5}$, and additionally, the film is essentially transparent, i.e., $\text{Im}\{n\} = 0$ (Figure S3). We observe the same effects for the amino acid Tryptophan (Trp); see Figure S4. Aromatic amino acid (such as Tyr and Trp) side chains contribute⁵⁴ and influence^{54,55} the CD spectrum of proteins. Therefore, if one aims to sense the chirality of small amounts of biomolecules, it is paramount that the metasurface is active in the far-UV spectrum region.³³

In Figure 2 d,e we present the CD and CB spectra of the RH (solid) and LH (dashed) gammadions coated with racemic (purple), L- (blue) and D-Tyr (red). The dotted curves represent the chiroptical signal of the bare amino acid film. Like in the previous simulation (Figure 1c,d), the reference consists of the chiroptical response from the Al gammadions surrounded by racemic layers. When the nanostructures are coated with L- or D-Tyr, the mirror symmetry among the CD and CB signals of the gammadion enantiomorphs are perturbed. For a fixed-handedness of gammadion (e.g., RH), the concavities of the CD spectra in the region $210 \leq \lambda \leq 220$ nm are altered

with respect to the reference. Furthermore, the peak at around $\lambda \approx 228$ nm is increased or decreased depending on the combination of chiral Tyr and gammadion enantiomers. Because of symmetry, the CD and CB signals for the pair L-Tyr-RH-gammadion and D-Tyr-LH-gammadion are mirror symmetric to one another (holding also for L-Tyr-LH-gammadion and D-Tyr-RH-gammadion). Similar findings are also present in the CB signals (Figure 2e). This can be a mechanism used for the enantiomeric discrimination of the analyte.

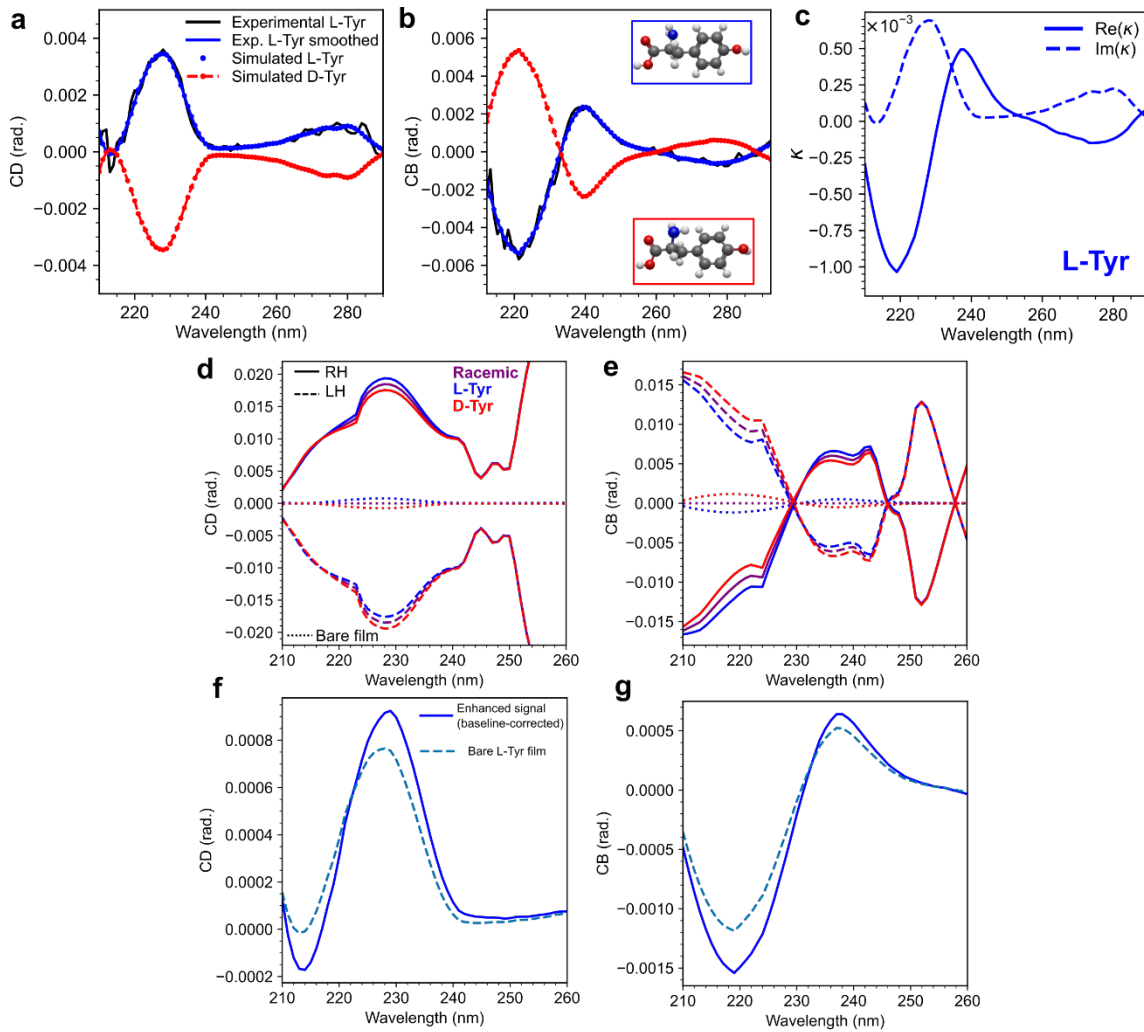


Figure 2. Far-field simulations of a resonant plasmonic-biomolecular system with real-world chirality parameter (κ). We employed Tyrosine as our model biomolecule. Experimental and

numerical (a) CD and (b) CB of a 90 nm-thick L-Tyr film on top of a fused silica substrate. The experimental CB and CD signals of the film were used to retrieve (c) $\text{Re}\{\kappa\}$ and $\text{Im}\{\kappa\}$ for L-Tyr, respectively. To simulate D-Tyr, we simply flipped the sign of κ determined for the L-enantiomer. (d) CD spectra of bare films (dotted lines) and of RH (solid lines) and LH (dashed lines) gammadion arrays coated with Racemic Tyr (purple curves) L-Tyr (blue curves) and D-Tyr (red curves). We use as reference the CD spectrum of the arrays coated with racemic layers, which form a perfect mirror image. Dissimilarly, when the plasmonic-molecular system is composed of chiral films, the CD spectra of the gammadion enantiomorphs lose this symmetry. Compared to when it is coated with a racemic film, a given gammadion enantiomer will interact differently with L- and D-Tyr (f,g) For the RH gammadion coated with L-Tyr, we perform a baseline-correction procedure to remove the bare signal of the plasmonic nanostructure. The baseline-corrected (f) CD and (g) CB are larger than the signal of the bare film, which clearly indicates enhancement.

These observed effects are more pronounced in the regions where the amino acids have stronger OA. At longer wavelengths, where biomolecular OA is vanishing, the mirror symmetry between the chiroptical signals of RH and LH gammadion arrays essentially reemerges (Figure 2d,e for $\lambda > 250$ nm and Figure S5). Previous works,^{38,56} have studied the resonant interaction of achiral nanoparticles with chiral molecules in the visible range. They verified that driving a plasmonic-molecular system out of resonance had detrimental effects to the observed CD enhancements. Our computational results suggest that when chiral nanostructures are involved, a resonant system with good spectral overlap is likewise important for superior chiral biosensing.

Given that the chiral nanostructure exhibits CD and CB signals of its own, we performed a baseline correction method⁵⁷ to take into consideration the "background" chiroptical response of

the gammadion. In Figure 2f, we present the results of this procedure for the RH gammadion coated with L-Tyr, where $CD_{\text{Baseline-corrected}} = CD_{\text{RH-gammadion-L-Tyr}} - CD_{\text{RH-gammadion-Racemic-Tyr}}$; in addition to the CD signal of the bare L-Tyr film (20 nm on top of a fused-silica substrate). Clearly, the overall CD is boosted in comparison to the isolated amino acid film. In Figure 2g, the same procedure also reveals an overall enhancement for the CB signal. We posit that the enhancement is due to resonant chiral plasmonic-molecular near-field interactions.

To test this hypothesis, we performed simulations of the RH Gammadion with 20 nm of L-Tyr film on top. We varied the dimensions (e.g., arm width and pitch) of the RH gammadion and calculated the impact on the $CD_{\text{Baseline-corrected}}$ signal (see Figure S6). Our simulations confirm that spectral overlap between plasmonic and biomolecular resonances is of great importance. When we increased the number of gammadions (145 nm arm width) in a fixed space by reducing its pitch from 500 to 475 nm, the signal enhancement was not increased given that the system was driven off-resonance. Negligible enhancements were observed for the array with 550 nm pitch regardless of the nanostructure dimensions due to lack of plasmonic OA at ~ 228 nm, i.e., where the L-Tyr films exhibit OA. Overall, the L-Tyr CD signal was only enhanced when the system displayed a clear spectral overlap between biomolecular and plasmonic OA resonances. The net OA enhancement can arise due to multiple mechanisms, among them are resonance shifts and modifications in the excitation and emission of the resonator modes.⁵³ These individual calculations are, however, beyond the scope of this work.

We then investigated the near fields of the gammadions. Chiral plasmonic nanostructures generate electromagnetic fields with enhanced optical chirality ($C_{\text{enh}} = C/|C_{\text{CPL}}|$),⁵ i.e., "superchiral fields"^{1,2} where the optical chirality density is higher than that for CPL ($|C_{\text{enh}}| > 1$). We illuminate the nanostructure with circular polarization of negative (e^-) and positive (e^+) ellipticities at $\lambda = 228$

nm; C_{enh} is computed only in regions accessible by the biomolecules. Figure 3a displays the side view of a LH gammadion array coated with 20 nm of racemic Tyr; two reference points are also provided. In Figure 3b, we present C_{enh} plots for a xz-cut at the middle of the unit cell. Figure 3c,d show xy-plots of a 5 nm-cut above the Al gammadion. The top amino acid layers display large and homogeneous values for C_{enh} , with good positional overlap with the enhancements in the electric field (Figure S7a,b). Conversely, the bottom Tyr layer display small (Figure 3b) and inhomogeneous (Figure S8) values of C_{enh} , with electric field enhancements confined only to the gammadion perimeter (Figure S7c,d). We expect the plasmonic-molecular interactions from the top layer to primarily contribute to the observed results. Furthermore, if a nanostructure is to be utilized in enantiomeric discrimination, a desired characteristic is that its near-field C_{enh} flips in sign when alternating between the two circular polarizations.⁵ From Figure 3c,d the sign is clearly reversed, leading to a homogeneous difference in optical chirality enhancement,⁵ i.e., $\Delta C_{\text{enh}} = C_{\text{enh}+} - C_{\text{enh}-}$ for the top amino acid layer (Figure S9).

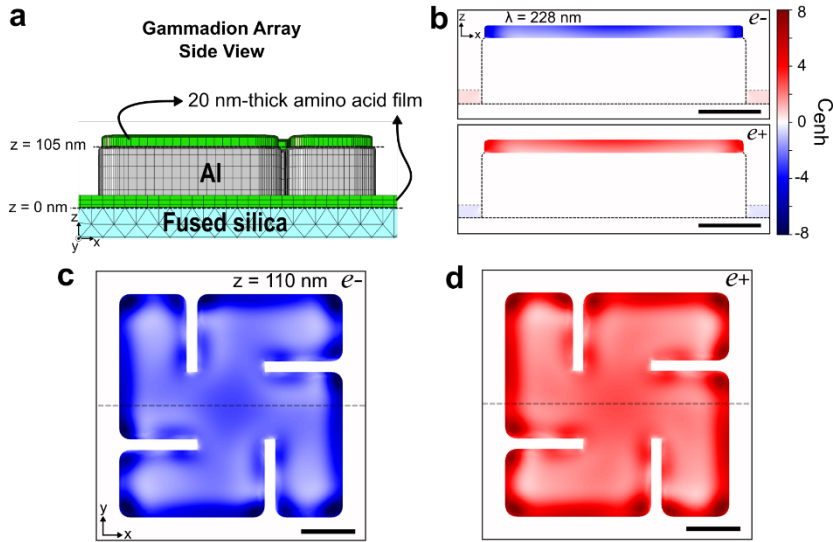


Figure 3. Near-field optical chirality calculations. (a) Gammadion array side view with reference point (b-d) Near-field optical chirality enhancement of a LH gammadion coated with 20 nm of racemic Tyr illuminated at resonant frequency with CPL. The top amino acid layer experiences high and homogeneous values of C_{enh} . Furthermore, as we switch the CPL state (handedness) from negative (e^-) to positive ellipticity (e^+), C_{enh} flips in sign. Scale bar: 100 nm.

To corroborate the results of the far-field EM simulations, we performed analogous experiments. We started by sublimating^{32,58,59} 30 nm of Tyrosine over lithographically defined Al gammadions. We characterize the samples with Mueller matrix (MM) polarimetry in transmission and retrieve the chiroptical signals with an analytic inversion of the MM⁶⁰ (See Supporting Information for detailed sample fabrication and optical characterization). This avoids potential artifacts to the chiroptical signals arising from linear effects⁶¹ from either the gammadion arrays or biomolecular films. The nanostructures without any biomolecular coating display optical activity from the far-UV ($\lambda < 250$ nm) to the visible part of the spectrum (Figure S11b,c). The fabricated gammadions have arms with width of ~ 150 nm, ~ 15 nm of gap between adjacent arms and periodicity of 600 nm (see Figure S11d,e for SEM micrographs). To allow a reliable comparison between each amino acid film (L-, D- or racemic Tyr), we used the same gammadion arrays throughout the experiments. We removed the tyrosine coatings by immersing the sample in dimethylformamide (DMF)⁶² for 15 minutes with gentle agitation, followed by a DMF and IPA rinse and N_2 drying. We kept track of the chiroptical signals of the bare arrays before each biomolecular deposition step. We did not observe changes in the CD or CB signals of the bare samples after cleaning them with DMF (Figure S12).

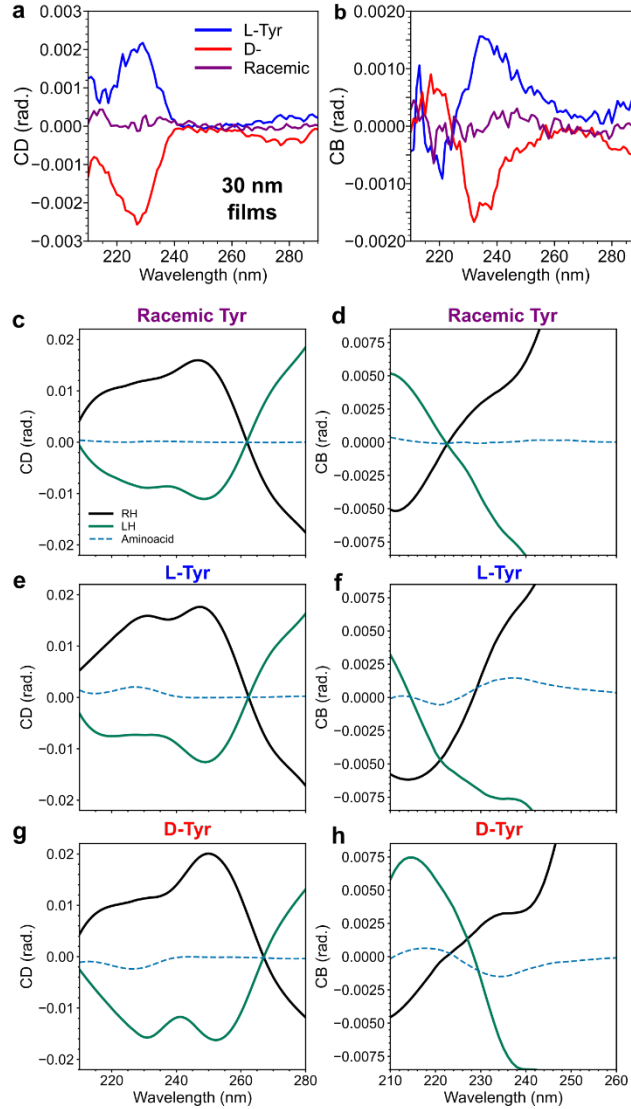


Figure 4. Experimental results of the plasmonic-biomolecular system. (a) CD and (b) CB of 30 nm-thick films of L-, D- and racemic Tyrosine on top of a fused silica substrate. The CD and CB peaks previously obtained for the 90 nm L-Tyr film are still observable, although the CB signal is relatively noisy, especially for the peak at ~ 218 nm. CD and CB spectra of the bare amino acid films and the Al gammadion arrays coated with (c, d) Racemic Tyr (e, f) L-Tyr and (g, h) D-Tyr. (c, d) When the nanostructures are surrounded by racemic layers, the CD (c) and CB (d) spectra of the coated gammadion enantiomorphs display good mirror symmetry. Perfect mirror symmetry is limited by nanofabrication heterogeneities. (e-h) Conversely, when the films are chiral, the

mirror symmetry is lost, as also observed on the simulations. These effects go beyond an additive effect. The perturbations arising from the chirality of the biomolecular film can be used to determine its handedness.

Figure 4a,b shows the CD and CB signals of 30 nm-thick films of L-, D- and racemic Tyrosine (see Figure S13 for their extinction spectra). The spectral features previously obtained for the thick L-Tyr film (Figure 2a,b) are still resolved, i.e., CD peak at 228 nm and CB peaks at 218 and 237 nm. This was expected because of the low thickness (< 100 nm) of the amino acid films.⁵⁸ When the arrays are coated with racemic Tyr, the CD and CB of the RH and LH nanostructures display good mirror symmetry (Figure 4c,d). Conversely, the handed amino acid films (L- or D-Tyr) notably perturb the chiroptical signals of the gammadions enantiomorphs. As shown in Figure 4e-h, the CD and CB of RH and LH nanostructures coated with L- or D-Tyr lose their spectral mirror image. The most pronounced features are the intense peaks appearing at $\lambda \approx 228$ nm in the CD spectra of the RH gammadion coated with L-Tyr and of the LH gammadion coated with D-Tyr (Figure 4e,g). Simultaneously, the CD signals for the other gammadion-amino acid combinations are relatively flat in this region, due to the opposite signs of the plasmonic and amino acid CD. At longer wavelengths, where the chiral amino acid films have vanishing OA, the chiroptical signals of the RH and LH gammadions are essentially mirror symmetric, regardless of the amino acid handedness or lack thereof (Figure S14). These results are consistent with the numerical calculations. Here, we do not perform any spectral summations of the chiroptical signals from the gammadion enantiomers in order to avoid data post processing, previously criticized due to the heterogeneities of nanofabricated chiral samples. Instead, we envision a platform where enantiomeric discrimination is enabled by "fingerprint-like" perturbations to the chiroptical signal

of the system, which will depend on the overlap of plasmonic and biomolecular chiral resonances. Nevertheless, to show that the effects in Figure 4e-h go beyond an additive effect, we subtracted the corresponding CD or CB signals of the bare chiral Tyr film from the chiroptical signals of the chiral Tyr coated gammadion enantiomers (Figure S15).

One desirability of plasmonic biosensors is the amplification of weak signals and the detection of small quantities of analyte.⁶³ Therefore, we shifted to evaporate ultrathin (average of 9.4 nm in thickness) films of Tyr, for which the CD spectra were barely resolved, and CB signal could not be reliably determined in our polarimeter (Figure S16). We estimate the quantity of Tyr to be on the order of ~ 10 attomole/per unit cell. Here, we focused on the CB signals of the system, given that the gammadion arrays did not display strong CD peaks at the region of interest (Figure S17). Figure 5a-c presents the CB spectra of the RH and LH gammadion arrays coated with racemic, L- and D-Tyr films, respectively. For plots with each amino acid, we also included a reference CB spectrum of a thick film to aid in the visualization of the molecular OA signals (or lack thereof). Clearly, the chiral nanostructures interact differently with the handed amino acids compared to when the gammadions are coated with a racemic film. Again, following the preceding theoretical and experimental results, these variations are pronounced in the biomolecular chiral resonance region, specifically at $\lambda \approx 240$ nm where there is a good overlap between the plasmonic and molecular CB peaks.

To further understand the effects of plasmonic-biomolecular OA spectral overlap, we scaled the unit cell of the fabricated arrays by factors of $7/8$ and $9/8$ (Figure 5d-i), intentionally altering the spectra of the bare gammadions. Figure 5d-f presents the CB spectra of the smallest gammadion arrays (e.g., $7/8$ scaled) coated with racemic, L- and D-Tyr films, respectively. When compared to the CB signal of the RH nanostructures coated with a racemic film, the spectrum of the same array

coated with L-Tyr (Figure 5e) displays a distinct concavity for $\lambda < 230$ nm (blue markers), due to the relative amplitudes and opposing signs of the plasmonic and biomolecular OA spectra in this wavelength regime. Furthermore, a distinct peak appears at $\lambda \approx 237$ nm (blue arrow). Similar results were obtained for the LH gammadions and D-Tyr combination (Figure 5f), effectively permitting the arrays to be used for enantiomeric discrimination. For the largest gammadion array (Figure 5g-i), the noticeable differences in respect to the control experiment (Figure 5g) were in the concavities of the CB (blue and red markers) spectra of the gammadions coated with chiral films for $\lambda < 237$ nm (Figure 5h,i). We reproduced the results of Figure 5d-i in eight sets of measurements for each amino acid (Figure S18). Furthermore, as predicted in the numerical calculations, at longer wavelengths where biomolecular OA is vanishing, we see no differences in the CB spectra of a gammadion array (i.e., RH) coated with racemic, L- or D-Tyr (Figure S19).

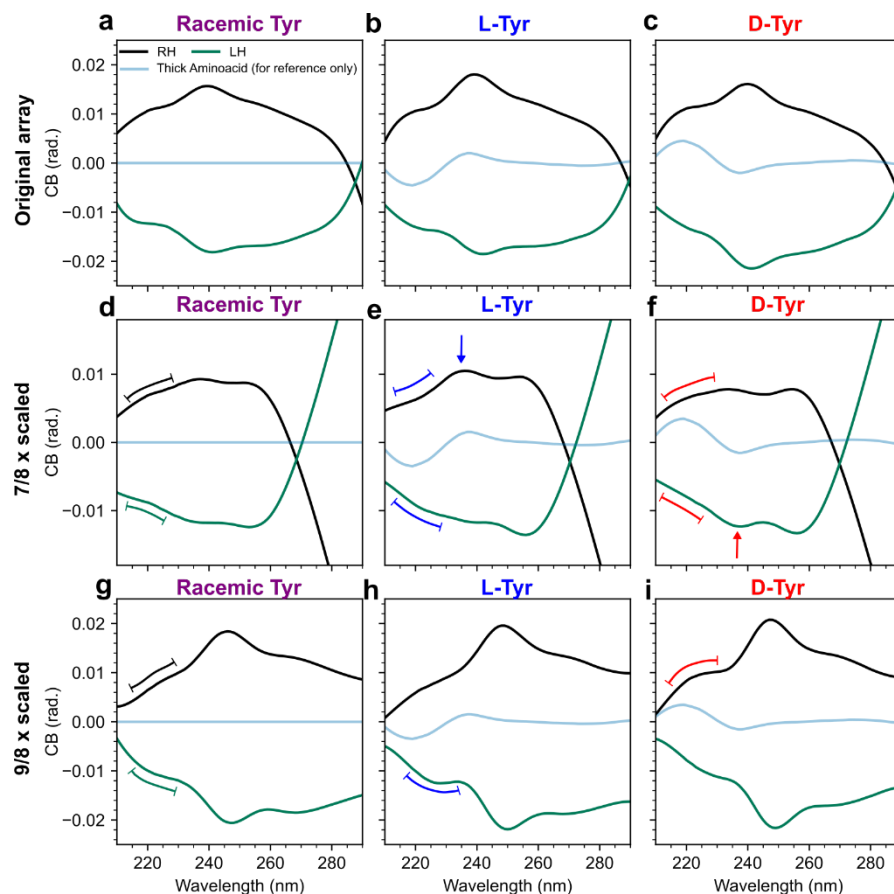


Figure 5. Experimental CB of the plasmonic-biomolecular system ultrathin (< 10 nm) films of (a) Racemic Tyr, (b) L-Tyr, and (c) D-Tyr films deposited on the gammadion array. Note the CB peaks for the bare films are not reliably determined in our instrument. A reference CB spectrum for a thick Tyr film is shown in light blue. When the nanostructures are surrounded by racemic Tyr the CB (a) spectra of the gammadion enantiomorphs display good mirror symmetry. Note, that perfect mirror symmetry is limited by nanofabrication heterogeneities in the gammadion. (b-c) Conversely, when (b) L-Tyr or (c) D-Tyr films are deposited on the gammadions the resulting left- and right-handed CB spectra show clear differences near the peaks of the (light blue) thick Tyr reference spectra. (d-f, g-i) CB results for 7/8 and 9/8 scaled gammadion arrays, with 9nm of (d,g) Racemic Tyr, (e,h) L-Tyr, and (f, i) D-Tyr films respectively, displaying similar trends.

In summary, we have shown that Al plasmonic nanostructures with far-ultraviolet chiroptical response can provide enantiomeric discrimination of ultrathin amino acid films. The unenhanced signals from the bare films (i.e., on top of the fused-silica substrate) were within the noise level of our instrument. This enhanced chiral biosensing is enabled by resonant interactions between chiral biomolecules and chiral plasmonic near-fields. Our simulated and experimental results confirm that the spectral overlap between plasmonic and natural OA is of great importance for enhanced biosensing. By evaporating ultrathin films (< 10 nm) of Tyrosine on the nanostructures, we show that minute quantities of biomolecules can perturb the plasmonic chiroptical signal and enable enantiomeric discrimination. The chiral signature of the plasmonic-biomolecular system strongly depends on how the resonances of these two components overlay on one another.

ASSOCIATED CONTENT

Supporting Information. Supporting Information Available: Experimental section – Electromagnetic simulations, chiroptical characterization with MM polarimetry, chirality parameter retrieval, sample fabrication, optical properties of modelled amino acid. Supplementary electromagnetic simulations figures including simulated chiroptical signals at longer wavelengths of the coated gammadions, electric-field and optical chirality enhancements. Supplementary experimental figures including chiroptical signals of bare Al gammadions, SEM micrographs of the structures, extinction spectra of the Tyr films, CD and CB spectra of coated gammadions at longer wavelengths, CD and CB spectra of ultrathin Tyr film, reproducibility study.

Refractive index and chirality parameter of L-Tyr.

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Author Contributions

K.M.M. and T.R.L. conceived the idea. T.R.L. fabricated the plasmonic arrays, measured their chiroptical response, and performed the electromagnetic simulations. L.Z. implemented the bi-isotropic model. T.R.L and O.K. deposited the amino acid thin films. T.R.L. wrote the manuscript with contributions from all co-authors. K.M.M. supervised and edited the manuscript. All authors have given approval to the final version of the manuscript.

Notes

The authors declare no competing financial interest.

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