Measurement Report: Effects of transition metal ions on the optical properties of humic-like substances (HULIS) revealing structural preference-A case study of PM$_{2.5}$ in Beijing, China

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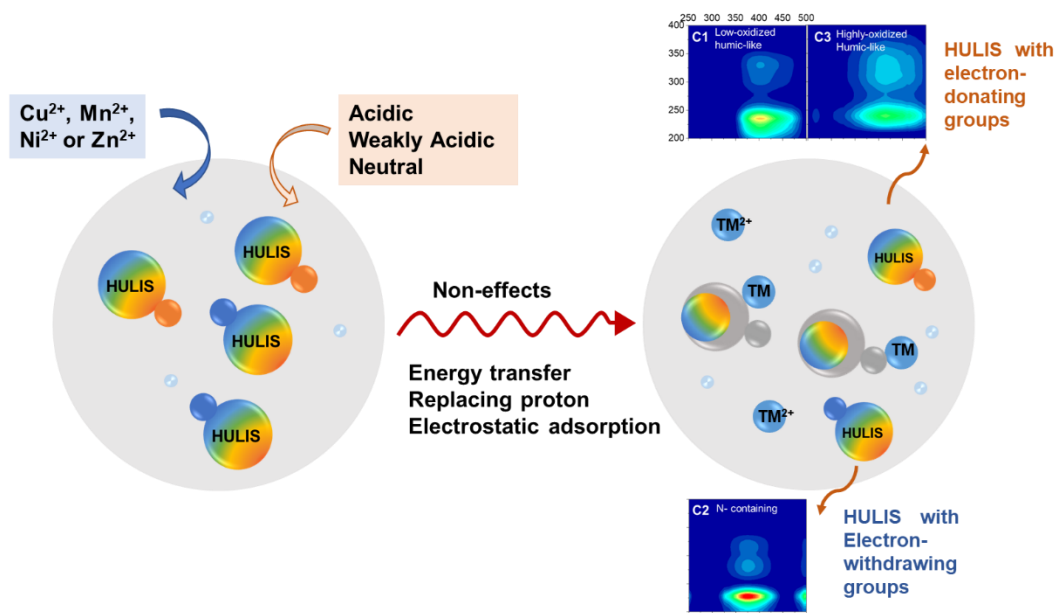
Abstract. Humic-like substances (HULIS) are complex macromolecules in water-soluble organic compounds containing multiple functional groups, and transition metal ions (TMs) are ubiquitous in atmospheric particles. In this study, potential physical and chemical interactions between HULIS and four TM species including Cu^{2+}, Mn^{2+}, Ni^{2+}, and Zn^{2+} were analyzed by optical method under acidic, weakly acidic and neutral conditions. The results showed that Cu^{2+}, Mn^{2+}, and Zn^{2+} only slightly enhanced mass absorption efficiency (MAE_{365}) of HULIS in winter, and had indiscernible effects on absorption Ångström exponent (AAE) of HULIS in both seasons under all acidity conditions. All four TMs had fluorescence quenching effects on winter HULIS, and only Cu^{2+} had similar effects on summer HULIS, with the highest quenching coefficients found under weakly acidic condition in both seasons. The [1]H-nuclear magnetic resonance (^{1}H-NMR) and Fourier-transform infrared (FT-IR) spectra revealed that Cu^{2+} mainly bound with aromatic species and tightened the molecule structures of HULIS. The parallel factor analysis (PARAFAC) results extracted four components of HULIS, including low-oxidized humic-like substances (C1), N-containing compounds (C2), highly-oxidized humic-like substances (C3), and the mixing residuals (C4), from the fluorescence spectra in both winter and summer. The spectral characteristic of HULIS with Cu^{2+} additions under three acidity conditions indicated that electron-donating groups of HULIS mainly corresponded to C1 and C3, with Cu^{2+} binding with HULIS by replacing proton, while electron-withdrawing groups of HULIS could correspond to C2, with its connection with Cu^{2+} through electrostatic adsorption or colliding induced energy transfer.

1 Introduction

Atmospheric particles, consisting of both organic and inorganic substances, have significant impacts on the natural environment, climate, and human health. Humic-like substances (HULIS) and transition metal ions (TMs) are among the major organic and inorganic fractions, and can substantially affect the physical and chemical properties of atmospheric particles (Hawkins et al., 2016; Zanca et al., 2017; Frka et al., 2018; Ma et al., 2019). HULIS and many TMs, like Fe, Cu, Mn, V, Cr, and Co, have negative impacts on human health since they can promote the generation of reactive oxygen species (ROS), which cause inflammatory response of human respiratory system (Verma et al., 2012; Gali et al., 2015; Lin and Yu 2019; Zhang et al., 2022a). HULIS are known as a mixture of macromolecular organic compounds, containing aromatics and aliphatic species with multiple oxygenated functional groups like carbonyls, hydroxyl, nitrate, and nitroxy-organosulfate (Win et al., 2018; Huang et al., 2022; Zhang et al., 2022b). TMs can transfer electrons and participate in chemical reactions or serve as catalyst, especially in atmospheric photochemistry (Mao et al., 2013; Guasco et al., 2014).

Considering that HULIS and TMs are both active chemical components of atmospheric particles, the significance of their co-effects is apparently conceivable. Early studies have demonstrated that HULIS can enhance the solubility of TMs by organic complexation (Paris and Desboeufs 2013; Scheinhardt et al., 2013), and TMs might mediate organic transformation processes and induce the formation of light absorbing secondary water-soluble organics and water insoluble abiotic polymers (Slikboer et al., 2015). More recent studies focused on the oxidation potentials, and ROS formation or other metal induced photochemical reactions of TMs (Fe^{2+}, Cu^{2+}, and Mn^{2+}) in combination with HULIS (Gonzalez et al., 2017; Lin and Yu 2019; 2020; Ye et al., 2021). The mixture of TMs and HULIS could exhibit synergistic or antagonistic effects on ROS formation associated with the

45 chemical composition of HULIS. Moreover, TMs can also bind with HULIS and directly affect the light absorption properties of HULIS (Fan et al., 2021; Wang et al., 2021).

To reveal the linkage between HULIS and TMs in atmospheric chemistry, this study characterized the varying optical and chemical properties of HULIS with addition of different TMs species under three different levels of acidity. The effects of TMs on light absorption of HULIS and the TMs-sensitive fraction of HULIS were determined by optical spectra, and parallel

50 factor analysis (PARAFAC) were performed to unveil the susceptive fluorescent components. The interaction between Cu^{2+} and HULIS were further analyzed for its prominent effects, and the potential reaction mechanisms were deduced based on the structural and optical variation scenarios. Results from the present study advanced our understanding on the physical and chemical characteristics and transformations of HULIS, especially the physical and chemical connections between HULIS and TMs.

55 **2 Methods**

2.1 Sample extraction and titration

In the present research, two concentrated HULIS solutions were extracted from daily $PM_{2.5}$ samples collected in ~~a~~summer and winter ~~and summer month~~of 2016 (30 samples in each season) in Beijing, and the detailed sample information can be found elsewhere (Qin et al., 2022). The HULIS isolation was mainly based on the method established by Lin et al., (2010) with some

60 adjusted procedures for practical reasons. Firstly, WSOC was ultrasonically extracted from 30 or 60 daily $PM_{2.5}$ filter samples (D = 2 cm, 30 pieces for winter and 60 pieces for summer), using 200 mL Milli-Q water three times, and filtered using 0.22 μm membranes. The liquids were then separated by SPE cartridges (Oasis HLB, 6cc, 500 mg/cartridge, Waters, USA), rinsed with 10 mL water, and pumped to dry. The cartridges were eluted with 10 mL methanol afterwards. Finally, the eluants were evaporated to dryness by N_2 at room temperature and redissolved into 200 mL ultrapure water to obtain a HULIS stock solution.

65 Before metal ion titration, the HULIS stock solutions were diluted by 10 folds, and the total organic carbon (TOC) concentration was determined by a TOC analyzer (Multi N/C 3100, Germany), with the carbon concentrations of 1.9 mg/L for winter HULIS and 1.6 mg/L for summer HULIS. The original acidity of HULIS was determined by a pH meter (Mettler Toledo, Swiss) before and after dilution. In both winter and summer, triplicate 600 mL of diluted HULIS solutions were separately added to 1 L sealed vials to obtain weakly acidic HULIS (pH = 5.65 for winter and pH = 5.92 for summer), acidic HULIS (pH

70 ≈ 3.0, by 0.05M H_2SO_4), and neutral (pH ≈ 7.5, by 0.1M NaOH), respectively. Considering the instability of natural pH gradient, excessive base solution was added to ensure the solution pH to be around neutral before measurement.

Four transition metals ions that are relatively rich in Beijing $PM_{2.5}$ and can complex with HULIS were considered including Cu^{2+}, Zn^{2+}, Mn^{2+}, and Ni^{2+} ~~were chosen as representative metal ions that might be reactive to HULIS, considering their richness in atmospheric environment or significant health risks (Fan et al., 2021;~~ (Wang et al., 2021). Titration experiments were

75 conducted by adding 0~~--~~ 526.32 μl corresponding metal sulfate solutions (0.01 M) and 9 ml diluted HULIS solution to a 10 ml amber volumetric flask, and ultrapure water was then added to obtain samples containing TMs concentrations of 0, 20, 50,

100, 200, 300 and 500 μM, respectively. The titrated solutions were shaken for 12 h at room temperature to ensure complexation equilibrium, and pH of all titrated samples were detected again before spectral analysis.

2.2 Spectral analysis

The optical properties of HULIS with addition of TMs (0–500 μM) were analyzed by UV-Vis and fluorescence spectrophotometers under acidic, neutral, and neutral pH. The chemical properties of HULIS before and after adding TMs were analyzed by [1]H-nuclear magnetic resonance (^{1}H-NMR) and Fourier-transform infrared (FT-IR).

The UV-vis absorption spectra of WSOC and HULIS were measured by UV-VIS photometer in Agilent Technologies covering wavelength range of 200 nm–500 nm with an interval of 1nm, using a scanning speed of 200 nm/min. The three-dimensional fluorescence spectra were measured by an Agilent Technologies photometer under the following conditions: excitation wavelength (EX) 200 nm–400 nm, step length 5 nm, slit width 10 nm; emission wavelength (EM) 250 nm–500 nm, step length 5 nm, slit width 5 nm; scanning speed: 2400 nm/min; and response time: automatic. The sample pool for UV-vis and three-dimensional fluorescence uses a quartz colorimeter with a light path of 1cm. The limit of detection for UV-Vis and fluorescence analysis were estimated as 2 times of their respective standard deviation of blanks, which were 0.005 and 0.01 absorption unit, respectively.

The infrared spectra of HULIS for wavenumber between 4000–400 cm^{-1} were measured by an FTIR spectrometer (Thermal Fisher, Nicolet IS10) at room temperature. The essential extraction of HULIS for FT-IR measurements was the same as those described in section 2.1 above, except that around 1 mg of the HULIS (dried by gentle stream of N_2 at room temperature 25 ℃) was directly mixed with 60 mg KBr and dried in a drying oven, rather than redissolved by water. The mixtures were then ground and pressed into firm tablets for analysis. A 600 MHz NMR spectrometer (Bruker, AVANCE III 600 M) was applied for ^{1}H-NMR spectra measurements. About 5 mg HULIS samples were redissolved into 1 mL D_2O for detection.

2.3 Data Analysis

2.3.1 Mass absorption efficiency and absorption Ångström exponent

The mass absorption efficiency (MAE, $10^3 \cdot AU \cdot cm^2/mg$) of HULIS and the absorption Ångström exponent (AAE) were obtained as follows (Kirillova et al., 2014; Saleh et al., 2014):

$$MAE = (A_\lambda - A_{700}) \times \frac{\ln{(10)}}{C_{spcies} \times L} \tag{1}$$

$$A_\lambda = K\lambda^{-AAE} \tag{2}$$

Where A_λ is light absorbance at wavelength λ, $C_{species}$ is the chemical concentration of organic compounds (WSOC and HULIS in the present research), L is the light path length (1 cm), K is a scaling constant, and the fitting wavelength of AAE is 300–400 nm.

2.3.2 Fluorescence indices

Fluorescence indices based on intensity ratios may provide clues about the condensation state of WSOC. The average fluorescence intensity is the mean intensity of an excitation-emission matrix (EEM) spectra; the specific fluorescence intensity (SFI) is the fluorescence intensity per unit TOC (mg/L) (Kalbitz et al., 2000). The biological index (BIX) reflects the freshness of microbially produced DOM (Parlanti et al., 2000), with a high BIX value (> 1) corresponding to a predominantly microbial derived organic matter, and a value of <0.6 indicating fewer microbial originated DOM (Qin et al., 2018). BIX is calculated as follows:

$$\text{BIX} = \frac{EEM_{Ex_{200-310},Em_{250-380}}}{EEM_{(max\ (Ex_{200-310},Em_{420-435}))}} \tag{3}$$

The fluorescence lifetime for HULIS was estimated by a formula derived from the Strickler–Berg equation (Strickler and Berg 1962). The τ_0/ρ ratio comprehensively reflects the fluorescence intrinsic lifetime (related to the fluorophore type) and fluorophore density (associated with the molecular structure), a large τ_0/ρ can be resulted from a long fluorescence lifetime or a small fluorophore (Xiao et al., 2018), it was a representation of fluorescence lifetime in this research.

$$\tau_0/\rho = c2.88 \times 10^{-12}n^2 \frac{\int_{Em} FI(v)dv}{\int_{Em} v^{-3}FI(v)dv} \int_{Ex} UVA(v)dv^{-1} \tag{4}$$

Where τ_0 is the intrinsic lifetime (ns), ρ is the density of fluorophore in the organics (mmol-fluorophore per gC), c is the total organic concentration (mgC/L), n is the refractive index of the solution ($n^2 \approx 1.8$), v is the excitation or emission wavenumber (cm^{-1}), UV is the UV-Vis absorbance (A.U.), and FI is the fluorescence intensity (R.U.).

Stokes shift (SS) can reflect the fluorescence energy of organics (Xiao et al., 2020; Yu et al., 2020), and is defined as Ex^{-1}-Em^{-1} nm^{-1}, demonstrating the energy loss of the excited fluorophore due to internal conversion during relaxation (Lakowicz 2006). AvgSS were defined as the intensity-weighted average Stokes shifts of HULIS (Xiao et al., 2019).

2.3.3 Parallel factor analysis (PARAFAC)

PARAFAC model can decompose complex EEM spectra into several main components by statistical method. The excitation spectrum, emission spectrum, and scores of each component are as follows:

$$x_{ijk} = \sum_{f=1}^{F} a_{if}b_{jf}c_{kf} + \varepsilon_{ijk} \qquad i=1,...,I\ j=1,...,J\ k=1,...,K \tag{5}$$

Where x represents the fluorescence intensity, f is the number of components resolved by PARAFAC, a is proportional to the concentration of the f-th component, and b and c are the scaled estimating of the emission and excitation spectra. The subscript i is the sample number, and j and k represent emission and excitation wavelength, respectively. Before performing PARAFAC, all EEM data were normalized to unit norm to reduce concentration-related collinearity and avoid extremely different leverages (Wang et al., 2020). Tucker congruence coefficient (TCC) was determined for each excitation spectrum and emission spectrum, and a threshold of 0.95 was applied to confirm the spectral congruence. The model was determined by half-split validation.

3 Results and ~~discussion~~Discussion

The light absorption and fluorescence efficiencies of winter and summer HULIS under acidic, weakly acidic, and neutral environments are depicted in Figure 1, and the spectra derived indices are listed in

Table 1. Generally, the overall light absorption efficiencies (represented by MAE) of winter HULIS were higher than those in summer, and the AAE showed a reverse trend, indicating higher light absorption abilities of winter than summer samples (Mo et al., 2021). The MAE and AAE of HULIS under three acidity levels in both seasons were consistent with those reported in our previous research, with the average MAE_{365} at 0.011±0.00 in winter and 0.005±0.001 in summer, and the corresponding average $AAE_{300-400nm}$ at 6.46±0.86 and 6.97± 0.83, respectively (Qin et al., 2022), which were comparable with those of Gosan Korea (Kirillova et al., 2014), Hongkong China (Ma et al., 2019), and Tibet China (Wu et al., 2019), but lower than those of biomass burning sources (Park et al., 2016).

The fluorescence efficiencies of winter HULIS (represented by SFI and AFI/TOC) were higher than those of summer HULIS as well (Figure 1), and the fluorescence indices reveled some structural differences. The SFI spectra of HULIS mainly exhibited two fluorescence peaks at wavelength of Ex = 225–230 nm, Em = 380–400 nm, and Ex = 325–330 nm, Em = 395–400 nm in both seasons, with the peaks being roughly characterized as fulvic-acid like group and protein-like groups, respectively. The Stokes shift index of AvgSS could be associated with the hydrophobicity/hydrophilicity of organics, with a low AvgSS implying a relatively high hydrophilicity and a high one implying a high hydrophilicity (aromaticity or humification degree) (Xiao et al., 2019). Thus, the higher AvgSS of winter HULIS indicated a higher hydrophilicity compared to that in summer. The different optical properties of HULIS between winter and summer were mainly resulted from divergent chemical structures. Our previous research had revealed that winter HULIS contained more unsaturated structures and were relatively hydrophobic, leading to the higher light absorption and fluorescence efficiencies, while summer HULIS were rich with oxidized species and functional groups and were more hydrophilic (Qin et al., 2022). Baduel et al., (2011) evaluated the varying optical properties of HULIS during O_3 oxidation, and found the C=O and COOH groups of HULIS increased with oxidation. However, Yang et al., (2021) found that O_3 oxidation of humic acid could consume some functional groups like aromatic C=C, OH, C=O, COOH and COO^-, causing a decrease in MAE and an increase in AAE, and these divergences were ascribed to the different structures of HULIS. The different structures of HULIS between winter and summer could not only lead to the variated optical properties but also the distinctive behaviors of HULIS with other coexist species in atmospheric environment. Therefore, it is necessary to analyze the relations between TMs and HULIS for winter and summer HULIS separately.

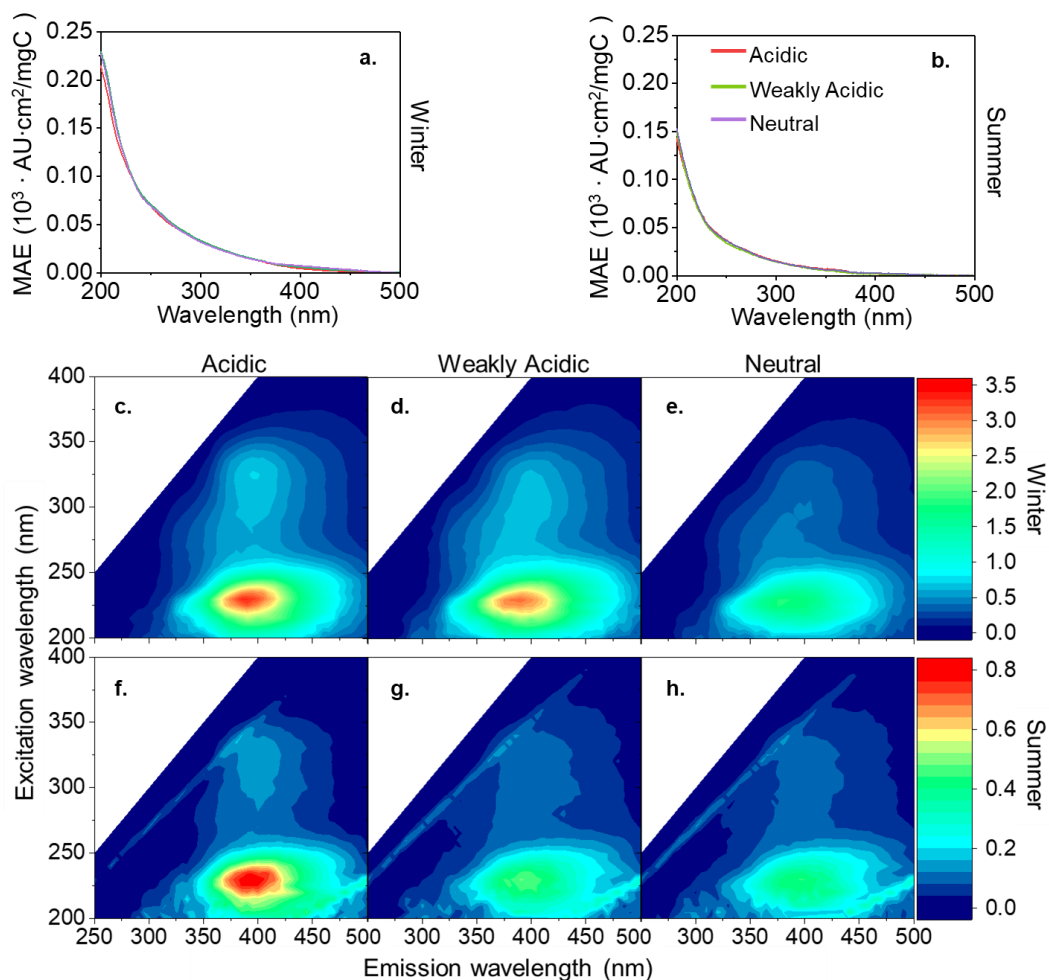



Figure 1 The MAE (a-b) and SFI (c-h) spectra of winter and summer HULIS under three acidity conditions.

Table 1 The light absorption and fluorescence indices of HULIS under three acidity conditions.

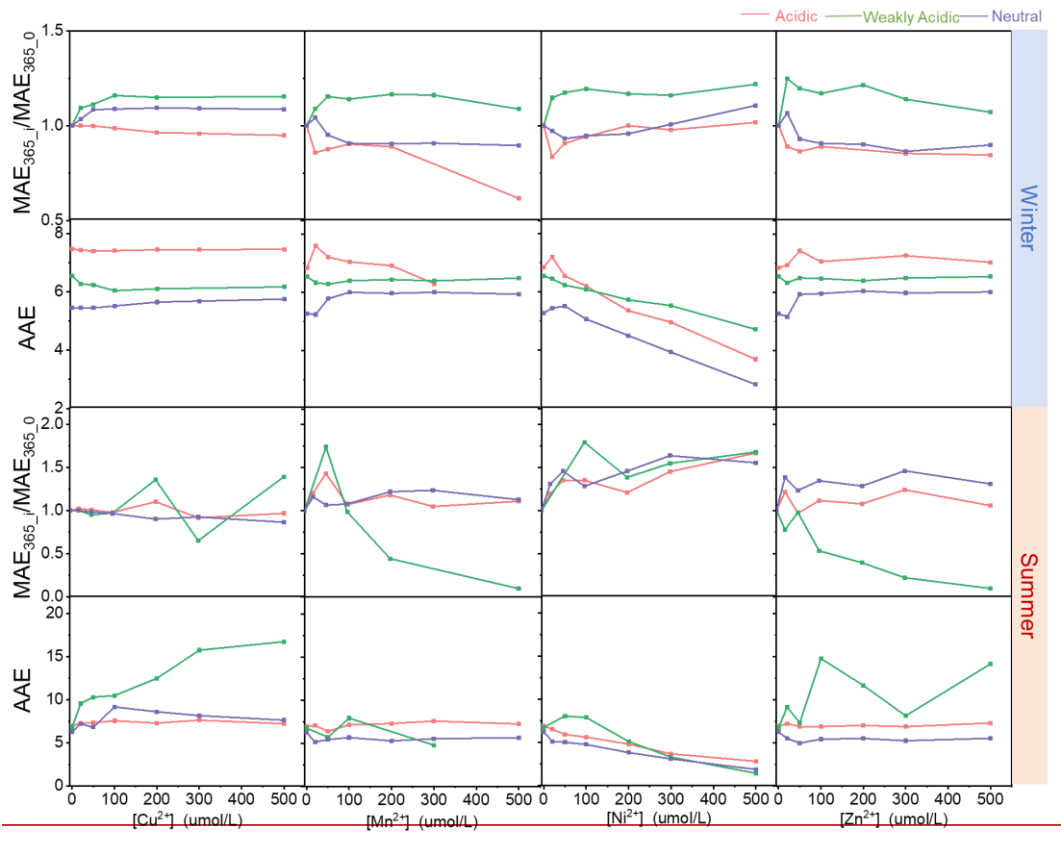
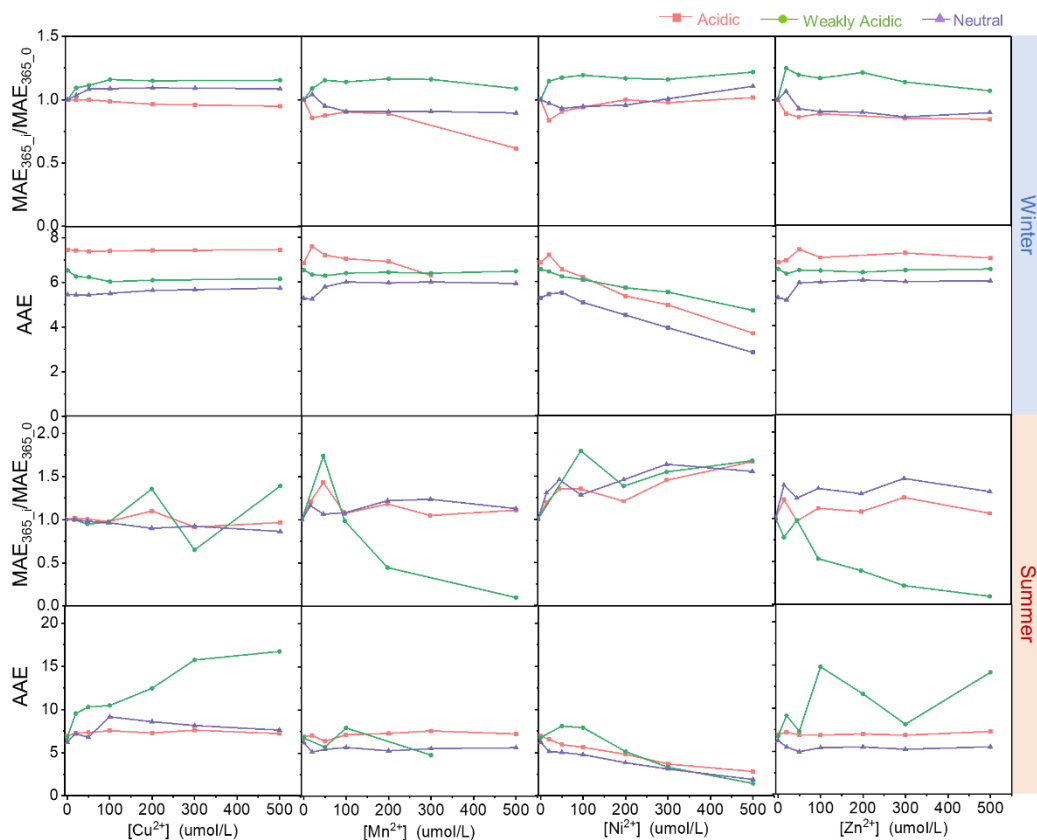
	Winter				Summer			
	Acidic	Weakly acidic	Neutral	Mean ± S.D.	Acidic	Weakly acidic	Neutral	Mean ± S.D.
MAE$_{365}$[a]	0.011	0.011	0.011	0.011 ± 0.00	0.006	0.004	0.006	0.005 ± 0.001
AAE$_{300-400nm}$	7.05	6.86	5.47	6.46 ± 0.86	6.97	7.80	6.15	6.97± 0.83
AFI/TOC[b]	0.79	0.62	0.58	0.66 ± 0.11	0.10	0.10	0.09	0.10 ± 0.01
BIX	1.07	0.96	0.99	1.00 ± 0.05	0.98	0.90	0.90	0.93 ± 0.05
AvgSS	1.54	1.54	1.54	1.54 ± 0.00	1.01	0.90	0.91	0.94 ± 0.06

| τ_0/ρ | 0.15 | 0.24 | 0.13 | 0.17 ± 0.06 | 0.56 | 0.92 | 0.60 | 0.69 ± 0.19 |

a: $10^3 \cdot AU \cdot cm^2/mgC$ b. $RU \cdot L/mgC$

3.2 Effects of TMs on the light absorption properties of HULIS

The gradients of MAE_{365_i}/MAE_{365_0} and $AAE_{300-400nm}$ for HULIS with TM (Cu^{2+}, Mn^{2+}, Ni^{2+}, and Zn^{2+}) concentrations ranging from 0 to 500 μM under three acidity conditions are depicted in Figure 2 and the whole MAE spectra are exhibited in SI Figures S1 and S2. For winter HULIS, under acidic environment, the addition of Mn^{2+}, Ni^{2+}, and Zn^{2+} at low concentrations (< 50 μM) could induce a decrease of 14%–16% in MAE_{365}, and further increasing TM concentrations had little effects on MAE_{365}, whereas Cu^{2+} showed minimal effects on MAE_{365}, which only decreased MAE_{365} by 5% with increasing TM concentrations.; these results were consisted with former study who found that the light absorption capacity of the atmospheric HULIS solution in the $Cu^{2+}/Al^{3+}/Zn^{2+}$-coupled system can be enhanced to $5-35\%$ (mean: 19%) (Li et al., 2022). Under weakly acidic environment, a small amount of any TMs (0–50 μM) could induce evident increase of MAE_{365} at a range of 16% to 20%, and then MAE_{365} became steady or slightly decreased with further increasing TM concentrations. As for the neutral environment, the addition of Cu^{2+} could lead to up to 9% increase in MAE_{365}, but Cu^{2+} showed inhibiting effects on MAE_{365} in cases with additions of Mn^{2+}, Ni^{2+}, or Zn^{2+}, with the maximum decrease at 11%, 7%, and 14%, respectively. Nevertheless, TMs could only change the $AAE_{300-400nm}$ of HULIS slightly, and the solution acidity sieved the initial $AAE_{300-400nm}$ of HULIS; besides, the TM concentrations had inconspicuous effects on the regularity. An exception was observed for Ni^{2+} because of its self-absorption between 350–400 nm, leading to the continuously reducing of $AAE_{300-400nm}$ with increasing $[Ni^{2+}]$.

For summer HULIS, Cu^{2+}, Mn^{2+}, and Zn^{2+} could cause fluctuations in MAE_{365} with increasing metal ion concentrations under acidic and neutral environments, and they inhibited the light absorption efficiencies of HULIS under weakly acidic environment. Likewise, Ni^{2+} could enhance the MAE_{365} of HULIS under all acidity conditions because of the increasing absorption generated by increasing Ni^{2+} concentrations. The $AAE_{300-400nm}$ of HULIS had subtle variation when adding Cu^{2+}, Mn^{2+}, and Zn^{2+} under acidic and neutral environments, indicating that the TMs-HULIS mixtures might not compose complexations or their complexes exhibited no structural difference. Yet when the solution was weakly acidic, the $AAE_{300-400nm}$ exhibited sharp fluctuations with increasing Mn^{2+} and Zn^{2+}, and even increased with increasing Cu^{2+} concentrations.

190 **Figure 2 The light absorption indices of HULIS with additions of TMs under different acidity conditions.**

3.3 Effects of TMs on the fluorescence properties of HULIS

The effects of TMs on the fluorescence indices of HULIS under different acidity environments are shown in Figure 3 and Figure 4, and the fluorescence spectral variations are provided in SI Figures S3 and S4. Obviously, all four TMs showed quenching effects on the AFI of HULIS with fiddling seasonal or acidity dependent differences, with the effectiveness ranking in the order of $Cu^{2+} > Mn^{2+} \sim Ni^{2+} > Zn^{2+}$. At the maximum TMs concentration of 500 μM, AFI of winter HULIS reduced by 5%–21%, 5%–11%, and 7%–28%, and of summer HULIS reduced by 8%–22%, 12%–39%, and 5%–26% under acidic, weakly acidic, and neutral environments, respectively.

The effects of TMs on fluorescent HULIS could be evaluated by the Stern-Volmer equation. Supposing metal ions only singly bonded with HULIS molecules, the quenching constants (k) of TMs species to the total fluorescence of HULIS were obtained and are listed in Table 2. Briefly, for winter HULIS, all TMs species had significant liner correlations with AFI/AFI_0 ($R^2 >$ 0.6). Cu^{2+} and Mn^{2+} had largest k constants under acidic and neutral environments and relatively low k under weakly acidic environment, indicating that Cu^{2+} and Mn^{2+} might mainly bond with organic groups that dissociated under acidic or neutral environments. Ni^{2+} had the same k under weakly acidic and acidic environments and larger k under neutral environment,

indicating that Ni^{2+} might prefer to bond with neutral HULIS. Similarly, Zn^{2+} had larger k under acidic environment, indicating a preference of acidic environment for the complexation of Zn^{2+} and HULIS. For summer HULIS, Cu^{2+} still had significant and large k under acidic and neutral environments and relatively low k under neutral environment. Mn^{2+} only had a relatively large k under acidic environment and a low k under neutral environment, but no statistically significant k under weakly acidic environment. Besides, AFI/AFI$_0$ of HULIS with Ni^{2+} and Zn^{2+} also had non-significant k, indicating negligible interactions between fluorescent HULIS and these two ions.

Although TMs showed quenching effects on HULIS, the locations of fluorescence peaks rarely moved with increasing TM concentrations. Thus, the fluorescence spectra-based indices only show minimal variations under different acidity conditions with increasing metal ions concentrations. In Figure 4, the Stokes shift spectra of HULIS separated into groups under three acidity conditions for both seasons. Only addition of Cu^{2+} could hierarchically distribute the spectra slightly. The other three TMs showed inconspicuous effects, indicating that they could not cause substantial structural variations on fluorescent HULIS, as can be more visually presented by the ratio of AvgSS$_0$/AvgSS$_i$ (Figure 5). Additionally, the addition of Mn^{2+}, Ni^{2+}, and Zn^{2+} didn't change the τ_0/ρ of HULIS either, which might confirm that they did not bound with HULIS molecules since fluorescent lifetime is a sensitive indicator vulnerable to the micro-environment surrounding detected objects (Coble, 2014).

Table 2 The fluorescence quenching coefficients of TMs and HULIS obtained from Stern-Volmer equation

Stern-Volmer	Acidity	Cu^{2+}		Mn^{2+}		Ni^{2+}		Zn^{2+}	
		$k \cdot 10^3$	R^2	$k \cdot 10^3$	R^2	$k \cdot 10^3$	R^2	$k \cdot 10^3$	R^2
	Acidic	0.60	0.99	0.10	0.98	0.10	0.95	0.20	0.98
Winter	Weakly Acidic	0.10	0.86	0.08	0.91	0.10	0.95	0.07	0.83
	Neutral	0.60	1.00	0.10	0.68	0.20	0.90	0.07	0.58
	Acidic	0. 60	0.99	0.20	0.38	-0.02	0.79	0.01	0.02
Summer	Weakly Acidic	1.00	0.82	-0.10	0.02	-0.04	0.05	-0.20	0.13
	Neutral	0. 40	0.88	0.05	0.63	0.05	0.08	-0.20	0.16

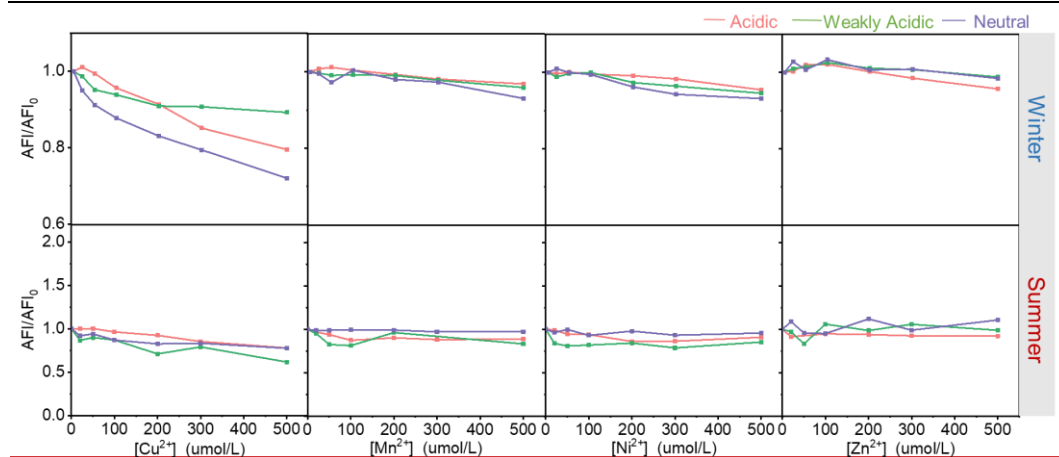
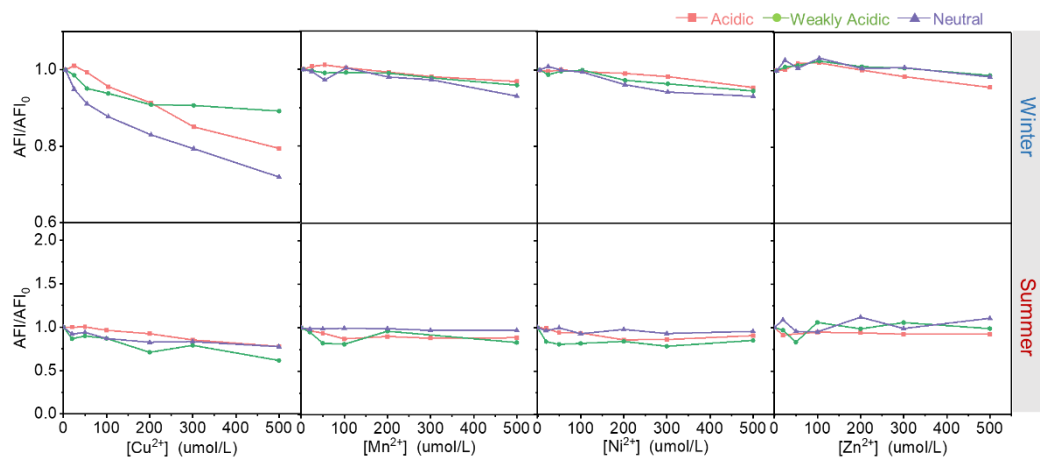



Figure 3 The average fluorescence intensities of HULIS with adding of Cu^{2+}, Zn^{2+}, Mn^{2+}, and Ni^{2+} under different acidity conditions in winter and summer.

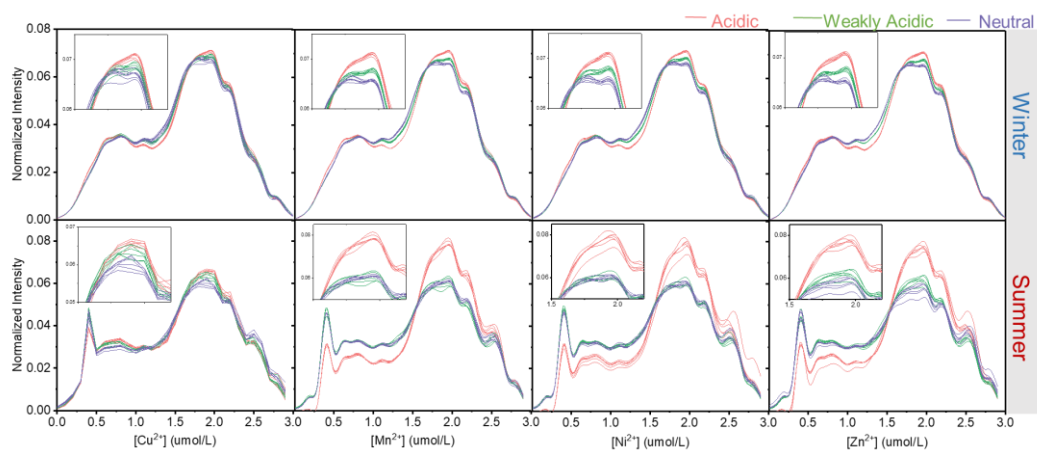



Figure 4 The Stokes shift spectra of HULIS with adding of Cu^{2+}, Zn^{2+}, Mn^{2+}, and Ni^{2+} under different acidity conditions.

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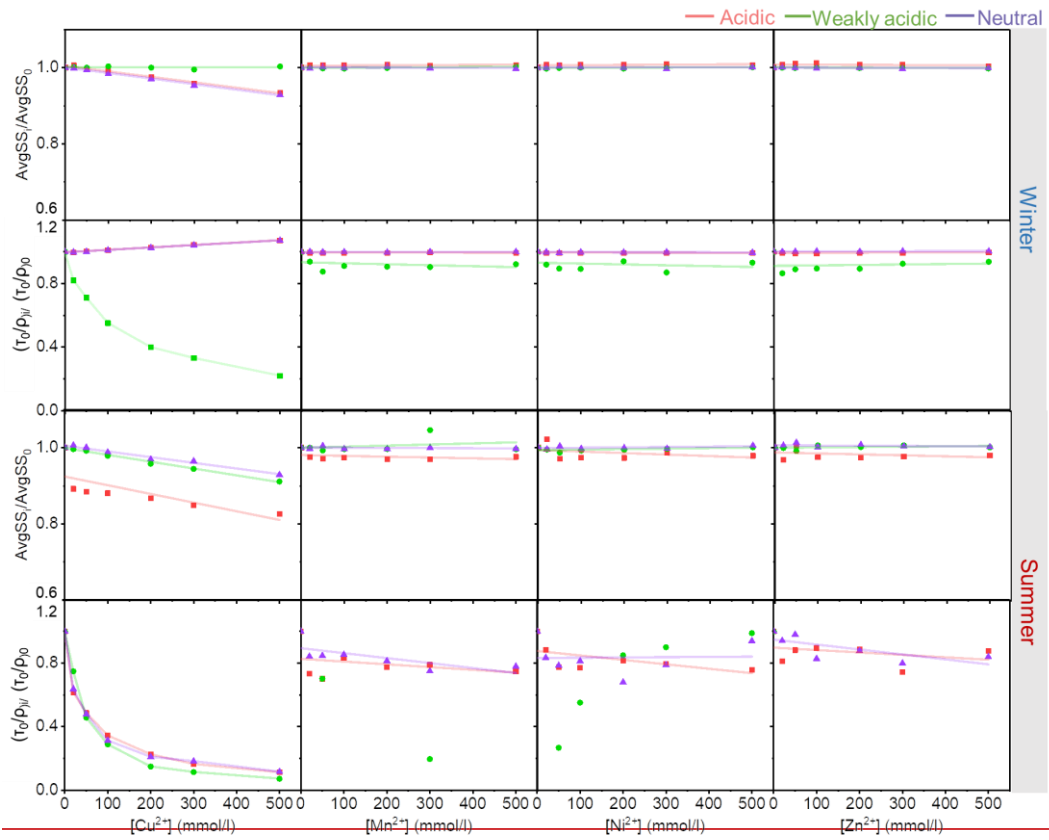
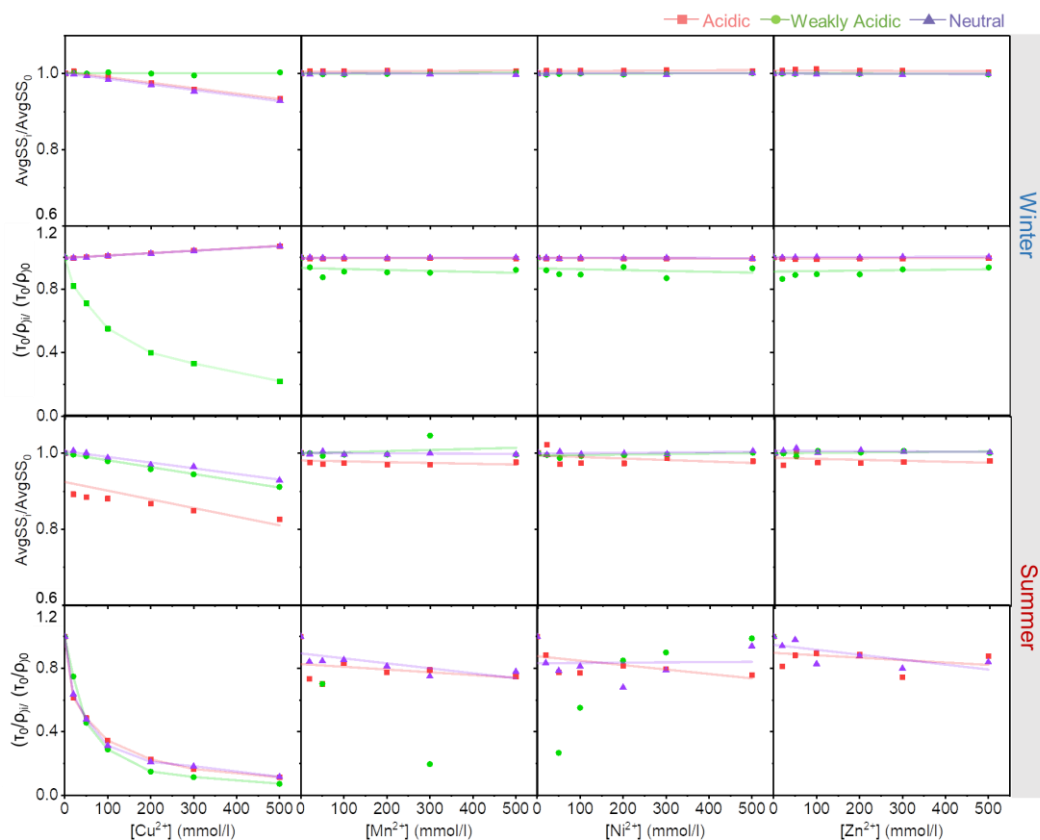


Figure 5 The fluorescence indices of HULIS with increasing Cu^{2+}, Zn^{2+}, Mn^{2+}, and Ni^{2+} concentrations under different acidity conditions.

3.4 Distinguishing the varying fractions of HULIS with additions of TMs

230   To further identify the response of specific HULIS compounds to TMs, PARAFAC methods were applied to distinguish the varying groups of HULIS with addition of different TM species. In general, four compounds (C1-C4) were sorted out from EEM spectra of winter and summer HULIS, and they were characterized as low-oxidized humic-like substances (C1), N-containing compounds (C2), highly-oxidized humic-like substances (C3), and the mixing residuals (C4), respectively (Chen et al., 2016). The constitution of C1-C4 in winter and summer HULIS under different acidity levels are listed in

235   Table 3.

    Form the correlations between F/F_0 ratio and TM concentrations of fluorophores C1-C4 shown in Figure 6 and Figure 7, it was obvious that the intensities of four fluorophores had increasing or decreasing tendencies in response to the acidity and TM species because of the divergent acid/base groups of HULIS. In the case of winter HULIS under acidic environment, Cu^{2+} could induce a substantial fluorescence quenching effects on C1 and C4 and a concurrent modest fluorescence increasing

240   effects on C3, Mn^{2+} and Ni^{2+} exhibited weak fluorescence quenching effects on C1 and C3, and Zn^{2+} exhibited weak fluorescence quenching effects on C1 and C4. Under weakly acidic environment, all of the four TMs exhibited discernible

quenching effects on C2 and C3, with Cu^{2+} manifest as the most potent quencher. Under neutral environment, Cu^{2+} significantly quenched the fluorescence intensities of C1, C2, and C4, Ni^{2+} displayed modest quenching effects only on C3, and other metal ions showed inconspicuous fluorescence quenching/enhancement effects.

245 As for summer HULIS, only Cu^{2+} exhibited notable quenching effects on fluorophores C1-C4, while Mn^{2+}, Ni^{2+}, and Zn^{2+} showed rather minimal or erratic effects under all acidity conditions. Under acidic and weakly acidic environments, Cu^{2+} could reduce the fluorescence intensity of C1-C3, and contributions of C4 to total fluorescence intensities gradually increased. For neutral environment, Cu^{2+} exhibited pronounced fluorescence quenching effects on C2, and mild reduction tendencies on C1 and C3. Because of the minimal quenching effects of Mn^{2+}, Ni^{2+}, and Zn^{2+} on C1-C4, most of the correlation coefficients (R^2)

250 between F/F_0 and metal ions concentrations were small, indicating the insignificant effects of Mn^{2+}, Ni^{2+}, and Zn^{2+} on HULIS.

Table 3 The PARAFAC decomposed fractions of HULIS for winter and summer under three acidity conditions.

		C1	C2	C3	C4	Total fluorescence intensity
	Acidic	42%	19%	24%	15%	805.52
Winter	Weakly acidic	29%	24%	27%	19%	869.69
	Neutral	22%	27%	29%	22%	587.49
	Acidic	50%	28%	20%	2%	192.09
Summer	Weakly acidic	23%	26%	18%	32%	179.17
	Neutral	16%	32%	16%	36%	161.00

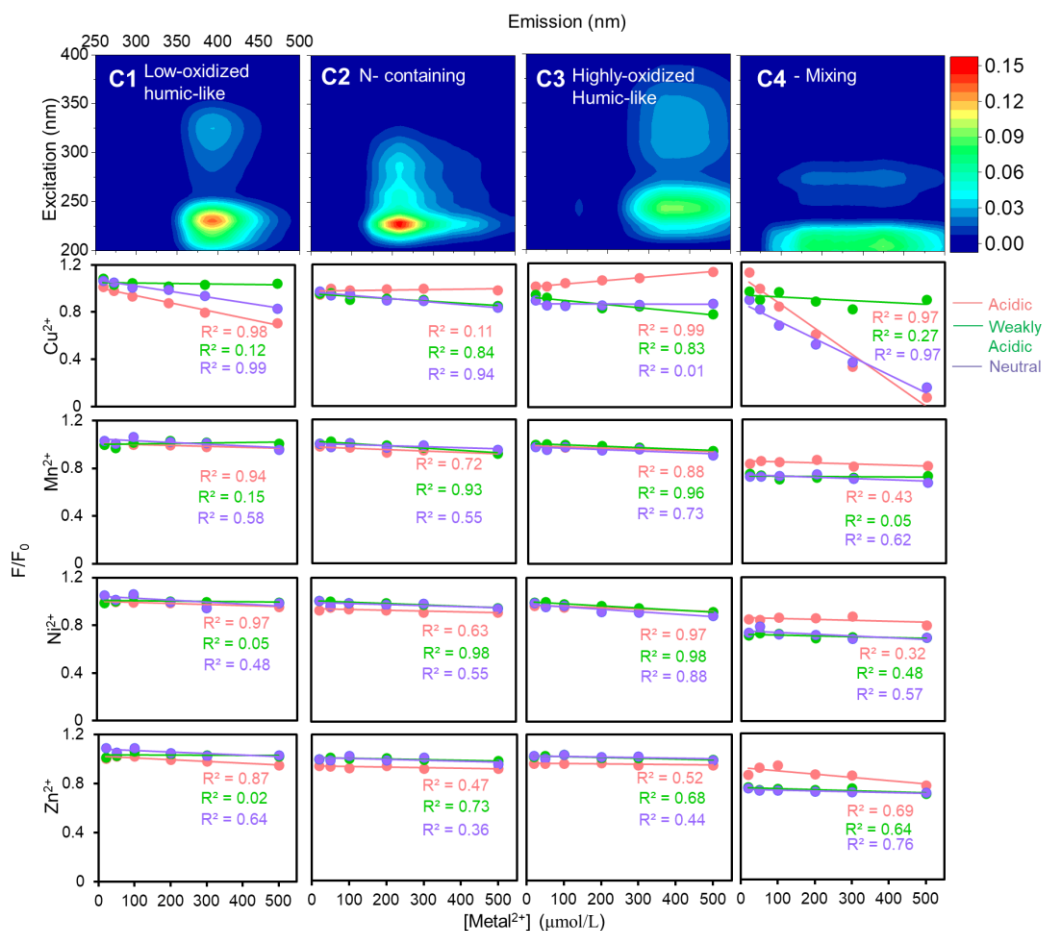



Figure 6 The changing patterns of four fluorophores with TMs under different acidity conditions for winter HULIS.

In light of the above analysis, it was obvious that Cu^{2+} demonstrated much more significant effects on optical properties of HULIS than the other three metal ions, which might be because of their disparate complexing abilities (Fan et al., 2021). Li et al., (2022) found that with increasing Fe^{3+}, Al^{3+}, and Cu^{2+} concentrations, the fluorescence intensities of highly oxidized HULIS decreased, and those of low oxidized HULIS gradually increased, and they ascribed these phenomena to the fluorophore blue-shifts of highly oxidized HULIS induced by complexation. Likewise, some other studies observed strong bondage between Cu^{2+} and atmospheric organic matter (Wang et al., 2021; Ma et al., 2022). However, the detailed mechanisms causing the differed complexation results of TMs remained unclear.

It is noticed that studies on the effects of TMs on optical properties of fulvic acid and humic acid had been conducted in earlier years in surface water than atmospheric environment, which provided an inspiration to the present study. For example, it was reported in 1980s that Cu^{2+} posed greater fluorescence quenching effects on fulvic-acid than Co^{2+} and Mn^{2+}, and it was thus proposed that Cu^{2+} bound internally to fulvic-acid, while Co^{2+} and Mn^{2+} might only externally attached to organics via electrostatic attraction (Ryan et al., 1983). Other studies found that Cu^{2+} had stronger complexation capabilities compared to

the other divalent transition metal ions, such as Pb^{2+}, Cd^{2+}, Ni^{2+}, and Zn^{2+} (Yamashita and Jaffe 2008; Chen et al., 2015; Huang et al., 2018). With application of two-dimensional infrared correlation spectroscopy and synchronous/asynchronous fluorescence spectroscopy analyze, carboxyl and polysaccharide groups were found to bind fast with Cu^{2+}; and bondage of Cu^{2+} with phenolic and aromatic carboxyl groups altered the molecule vibration of organics, inducing fluorescence quenching.

Besides, Cu^{2+} could bind with several amide and aliphatic groups (protein-like), leading to fluorescence quenching. More recently, Zhu et al. (2021) suggested that the small atomic radius and paramagnetic properties might result in the unique response of Cu^{2+} to carboxyl groups, thus a strong complexation ability with DOM. Considering the smaller molecules and aromatic structures of HULIS compared to humic substances in water, Cu^{2+} continued its specialty of significant fluorescence quenching effects on HULIS, whereas bondages between Zn^{2+} (or Mn^{2+} and Ni^{2+}) and responsive HULIS might be too weak to be observed.

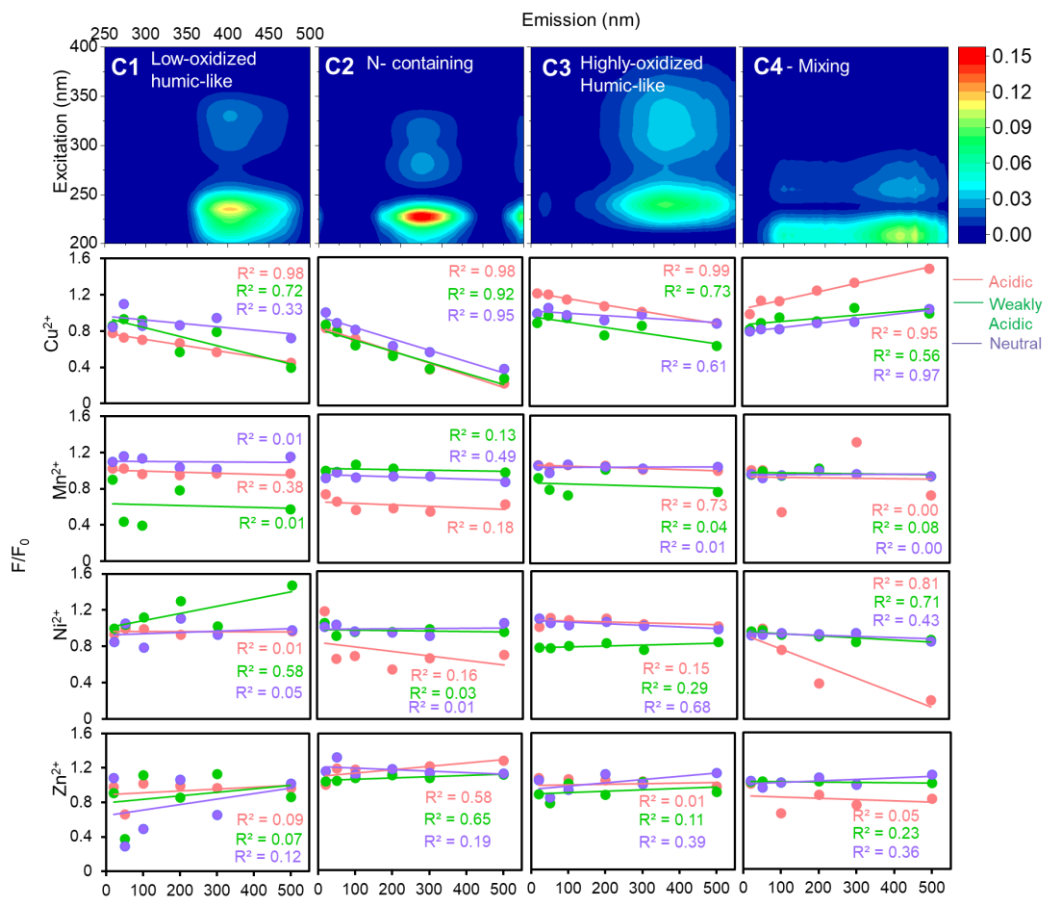
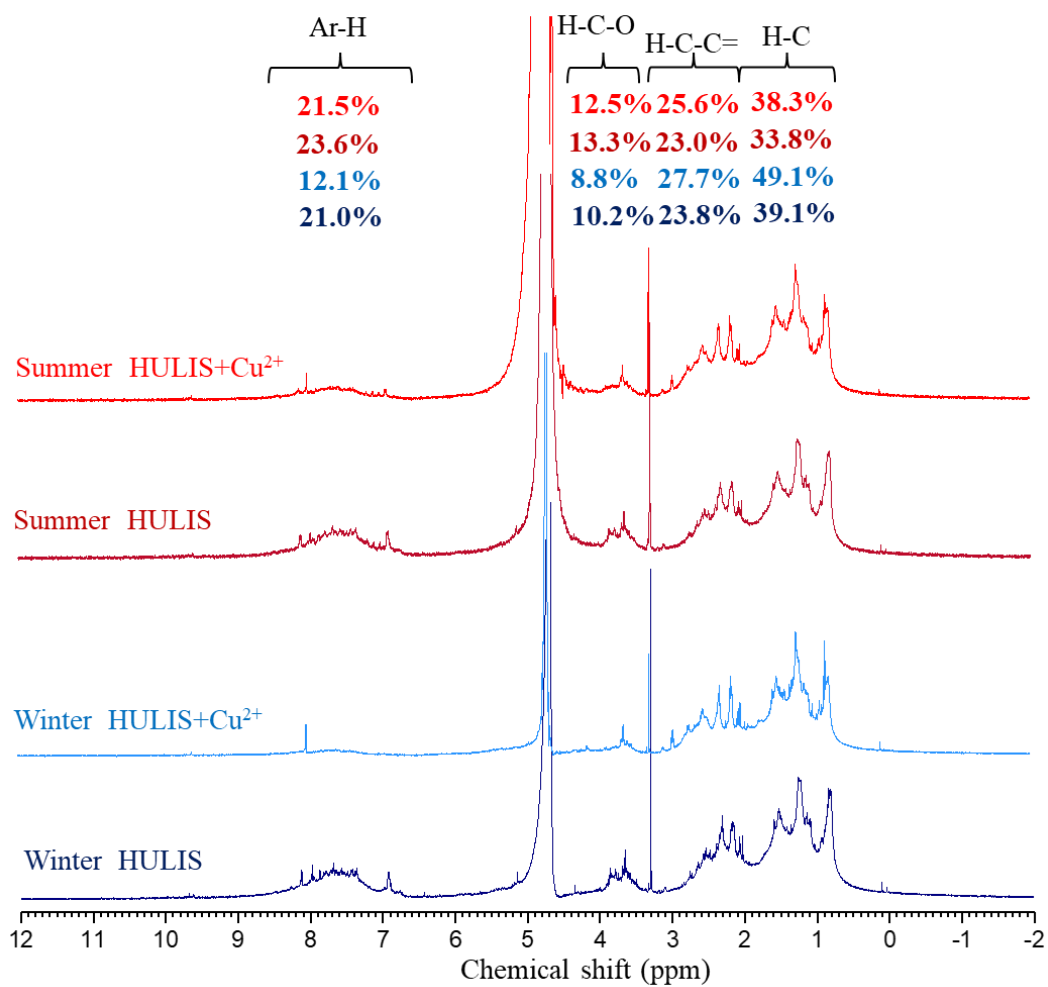



Figure 7 The changing patterns of four fluorophores with TMs under different acidity conditions for summer HULIS.

3.5 The hidden mechanism of HULIS with addition of TMs - taking Cu2+ as an example

To further understand the complexation mechanisms of Cu^{2+} and HULIS, the structural variations of HULIS before and after adding 100 μM Cu^{2+} were evaluated by ^{1}H-NMR and FT-IR methods, respectively (Figure 8 and Figure 9). The ^{1}H-NMR results showed that HULIS were rich with protons of H-C, H-C=C-, Ar-H, and H-C-O and functional groups of -OH or -NH, Ar-C=O, Ar-C-O-C, and C-O in both seasons, which were similar to the results of former researches (Fan et al., 2013; Hawkins et al., 2016; Kumar et al., 2017). The addition of Cu^{2+} mainly affected the protons bound with aromatics and oxygen in HULIS, with the signals located between δ7-δ8 and δ3.4-δ4 ppm trailing off, for which the Ar-H signal decreased by 8.9% and 2.1%, and H-C-O signal decreased by 1.4% and 0.8% in winter and summer, respectively. According to the basic principles of NMR spectroscopy, coordination or chemical exchange reactions between HULIS molecules and Cu^{2+} might cause the chemical shift signals vanished or observed as new signals (Günther 2013). Meanwhile, the functional groups of HULIS, revealed by FT-IR spectra, mainly included -OH/-NH, -CH, Ph-C=O, C=O, Ph-C-O-C, and -C-O in both seasons, and the summer HULIS contained more oxygen-containing functional groups than winter HULIS because of the high atmospheric oxidization capacity (Chen, et al. 2016; Haynes et al., 2019). With the addition of Cu^{2+}, the whole FTIR spectra of winter and summer HULIS blue shifted to higher energy, especially for O-containing aromatic groups, but the peak shapes remained unchanged, indicating that Cu^{2+} might not directly lead to the transformation of functional groups, but could result in an enhancement of chemical bonds in HULIS molecules.

295 **Figure 8 The ¹H-NMR spectra of HULIS and HULIS with addition of 100 μM Cu²⁺ in winter and summer.**

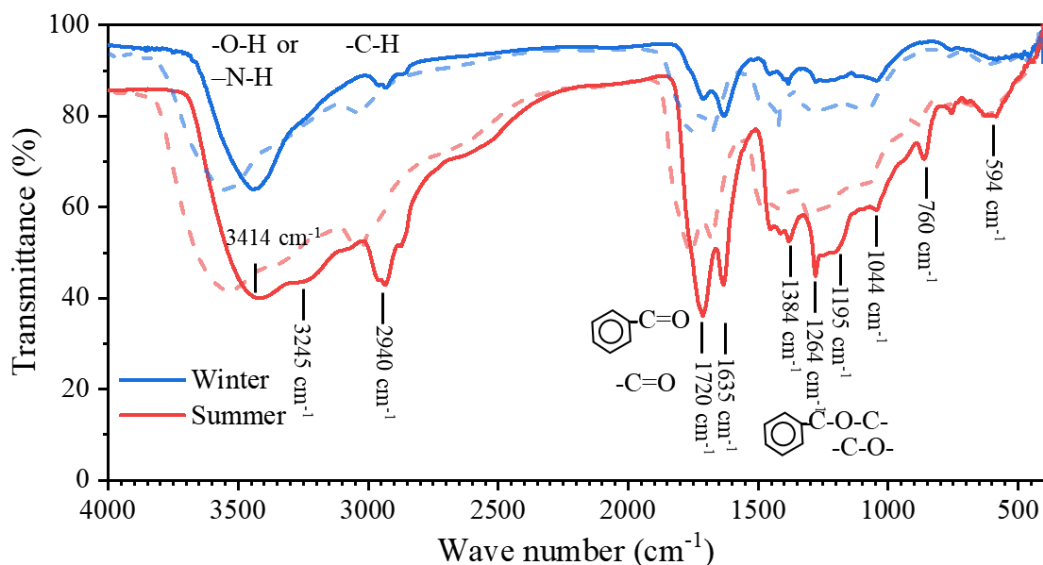
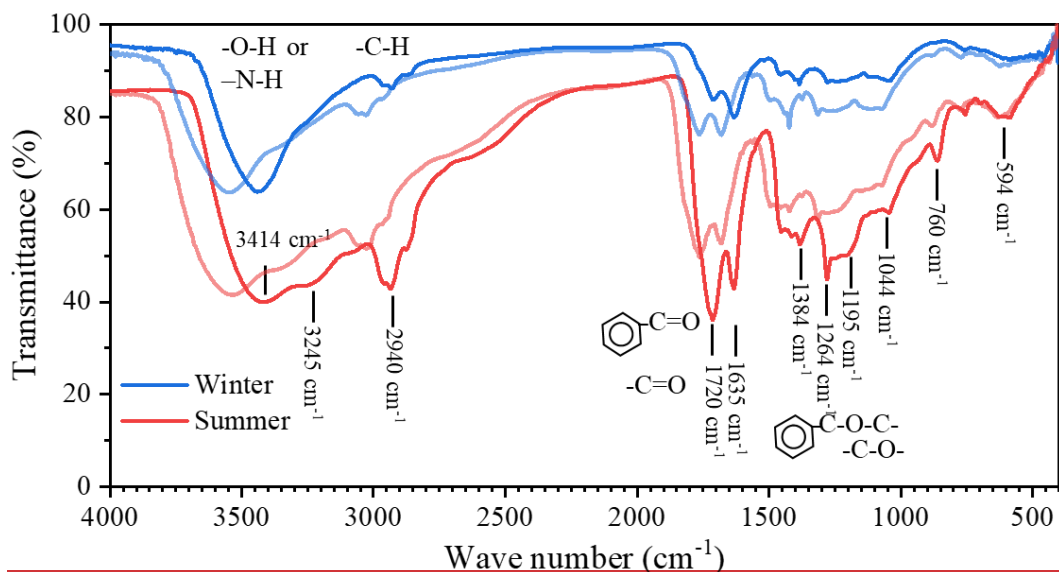



Figure 9 The FT-IR spectra of HULIS and HULIS with addition of 100 μM Cu^{2+} (dash lines) in winter and summer.

Our previous research had characterized the pK_a and corresponding acidic/basic functional groups of winter and summer HULIS, and proposed a hypothetical HULIS structure with unsaturated aromatic main body holding pH responsive functional groups (Qin et al., 2022). Briefly, the pK_a of 3.7 – 3.8 and 8.0 – 8.1 were assigned as electron-donating groups (represented by H$_1$A$_1$ and H$_3$A$_3$, respectively), and the pK_a of 5.0 – 6.3 were assigned as electron-withdrawing groups (represented by H$_2$A$_2$) for winter HULIS. Likewise, the pK_a of 6.4 was assigned as electron-donating groups (represented by H$_4$A$_4$) and pK_a of 8.4 – 8.5 was assigned as electron-withdrawing groups (represented by H$_5$A$_5$) for summer HULIS. By synthetically considering the versatile optical properties of HULIS with addition of Cu^{2+} under three acidity conditions, and its inherent acid/base groups,

the Cu^{2+} and HULIS complexation or interaction mechanisms were explored and the schematic diagram is depicted in Figure 10.

(1) Firstly, for winter HULIS under acidic environment, the acidity-sensitive HULIS molecules were hard to deprotonate when H^+ was crowding around. Most of HULIS existed as forms of H_1A_1, H_2A_2, and H_3A_3. Cu^{2+} could easily be absorbed by the electron-donating groups of HULIS to form $Cu-A_1H_1$ complex, or replace the H atom to form $Cu-A_1$, causing static fluorescence quenching of C1 and C4 which slightly increased the lifetime of HULIS. Simultaneously, Cu could alter the charge densities of the fluorescent aromatic molecules, leading to an enlargement of conjugated structures and enhancing the fluorescence of C3.

(2) Secondly, for winter HULIS under weakly acidic environment, some of the acidity-sensitive HULIS deprotonated and the solution contained A_1^-, H_2A_2, and H_3A_3. When added with Cu^{2+}, A_1^- complexed with Cu^{2+} to form $Cu-A_1$, whereas the sensitive structures of H_2A_2 were electron-donating groups which could hardly provide electrons to Cu^{2+}. However, Cu^{2+} could still be attracted by protons in the unsaturated aromatic structures of HULIS, causing dynamic fluorescence quenching of C2 and C3. With regard to the fast-decreasing lifetime and steady AvgSS of HULIS, chemical reactions might not be involved in these processes, and thus flexible structures of $Cu-A_2H_2$ by electrostatic adsorption or colliding induced energy transfer could be the possible explanation.

(3) Thirdly, for winter HULIS under neutral environment, the acidity-sensitive HULIS existed as A_1^-, A_2^-, and H_3A_3. The deprotonated HULIS had become non-fluorescent, and collision of A_1^- and A_2^- with added Cu^{2+} might not be observed. The fluorescent H_3A_3 could provide electrons to Cu^{2+}, leading to partially fluorescence quenching effects on C1 and C2, and the potential complexation mechanism might be similar to H_1A_1.

(4) Lastly, for summer HULIS, the quenching mechanism of Cu^{2+} and acidity-sensitive HULIS were similar to those of winter HULIS. Under acidic and weakly acidic environment, H_4A_4 and H_5A_5 of summer HULIS were not deprotonated, the quenching effects could be resulted from the replacement of H atom in H_4A_4 (electron-donating) with Cu^{2+}, causing decrease of C1-C3. Under neutral environment, Cu^{2+} complexed with deprotonated A_4^- and aromatic structure of H_5A_5 (electron-withdrawing), mainly causing decrease of C2.

Thus, it was speculated that HULIS with electron-donating groups mainly exhibit fluorescence signals in C1 and C3, which were mainly low and highly oxidized humic-like substances, respectively. Whereas for HULIS with electron-withdraw groups, the fluorescence signals were mainly exhibited in C2, which was N-containing species or protein-like species. Besides, although very few studies can explicit the interaction mechanisms between HULIS and heavy metals, the hypotheses of present study are aligned with a recent soil study, exploring the retention of heavy metals by humic acid using atomic force microscopy who found that some metals (Pb^{2+} and Cd^{2+}) formed strong adhesion stems from the synergistic metal-humic acid complexation and cation-π interaction at pH 5.8, leading to significant retention, while other metals (As^{5+} and Cr^{6+}) only bound weakly through hydrogen bonds (Wang et al., 2024).

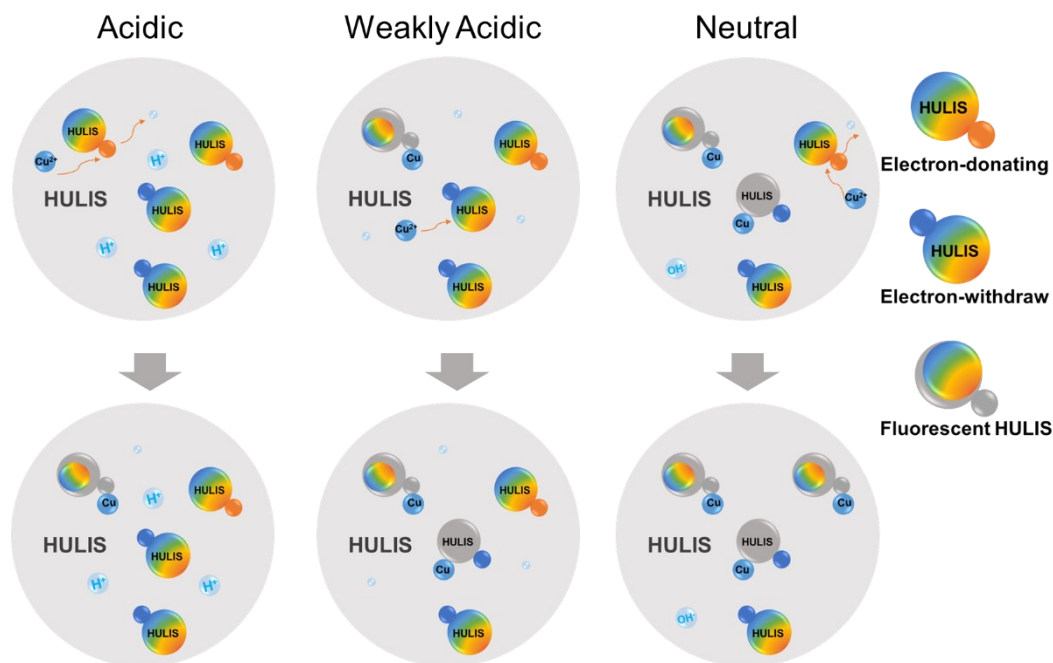



Figure 10 The schematic diagram of HULIS complexation with Cu^{2+} under different acidity conditions

4 Conclusions and Implications

In this study, the interrelations between four TMs and HULIS were analyzed by optical methods under acidic, weakly acidic, and near neutral environments. The results elucidated that the light absorption and fluorescence properties of HULIS were substantially influenced by the solution acidity. TMs either enhance or attenuate the signal strength, contingent on the metal species, but with negligible impact on the spectral indices. Cu^{2+} demonstrated the most conspicuous influence on the optical properties of HULIS, while Mn^{2+}, Ni^{2+}, and Zn^{2+} proved to be less responsive. Cu^{2+} could promote light absorption ability of HULIS at weakly acidic to neutral environment, but the effect was negligible under acidic environment. The ^{1}H-NMR and FT-IR spectra of HULIS and HULIS-Cu indicated that Cu^{2+} mainly bound with aromatic HULIS, resulting in more compacted molecule structures of HULIS. The distinct tendencies of PARAFAC-separated fluorophores with increasing Cu^{2+} under three acidity conditions indicated that electron-donating groups might correspond to low and highly oxidized humic-like substances, and Cu^{2+} coupled with HULIS by substituting H atom; while electron-withdraw groups were N-containing species or protein-like species, with their fluorescence quenching likely caused by electrostatic adsorption or colliding induced energy transfer. Results from this study not only revealed that the complexations of HULIS and TMs could be fragile, but also connected the fluorescence spectra of HULIS with structural characterizations through the micro-effects of TMs on HULIS. HULIS and TMs are both well-known ROS generative species, and the combined effects of HULIS and TMs on ROS generation can be synergistic or antagonistic depending on the TMs species. The ROS generation abilities of TMs and HULIS are essentially

determined by their physical and chemical properties and interactions, the complexation, electrostatic adsorption, and colliding induced energy transfer processes could enhance or prohibit ROS generation and resulted to synergistic or antagonistic effect. Our results ~~could~~ provide ~~some perception~~perceptions from chemical analysis perspective on interactions between HULIS and TMs, however, ~~the exact reaction mechanisms still require further~~due to the intricate nature of $PM_{2.5}$, many crucial transition metals, such as Fe^{2+}/Fe^{3+} and Cr^{6+}, which play pivotal roles in the chemical transformation of $PM_{2.5}$ or pose high risks to human health, are not addressed in the current study. Furthermore, comprehensive research on worldwide HULIS samples is also imperative for future investigations.

Code availability

Data availability

The data used in this study are available on the Zenodo data repository platform: https://doi.org/10.5281/zenodo.10460562

Author contribution

TJ and WX designed the experiments, QJ and QY carried them out, SZ and GY helped with the experimental issues, LJ and SS finished the data analysis. QJ prepared the manuscript with contributions from all co-authors, ZL, TJ and WX supervised the writing, and QT provided professional advices on the discussion section.

Competing interests

At least one of the (co-)authors is a member of the editorial board of Atmospheric Chemistry and Physics.

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