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New insights into the structure and degradation of alizarin lake pigments: input of the surface study approach

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Abstract

Alizarin (Az) lake is a typical example of lake pigments that is widely used in cultural heritages. Previous work focused on the complex formation in solution, theoretical models and crystallized stoichiometric complexes. The real nature of lake pigments, as well as the reason for its degradation, are still controversial. In this work, we prepared the Az lakes from the ancient recipe with non-stoichiometric complexes, i.e., $R = 5-200$ (R refers to molar ratio of Al^{3+}/Az). The influences of the dye concentration, pH and basic reagents on the structures and chromatic properties of bulk Az lakes were studied by attenuated total reflection Fourier transform infrared spectrum (ATR-FTIR), nuclear magnetic resonance (NMR) and CIE 1976 color space. In order to investigate the ~~surface~~ interaction between Az and the alumina inorganic substrate, likely determining in the aging of pigments, Az was deposited on a thin Al_2O_x film prepared by physical vapor deposition method. Polarization modulation infrared reflection absorption spectroscopy (PM-IRRAS) was employed to characterize the chemical reactivity and degradation of Az- Al_2O_x film complex. Az derivatives interacted with Al_2O_x film like bulk lakes. Al^{3+} were coordinated with 1,2-dihydroxyl (catechol) group, instead of 1-hydroxy-9-keto group. Exposure under UV light resulted in degradation of Az- Al_2O_x film complex by oxidizing catechol groups to carbonyl. This study presented a new insight into the structure and degradation of Az lake.

Keywords: Lake pigments, Alizarin, Photostability, Al_2O_x film, UV light.

1 Introduction

The growing attention in the study of art objects for the purposes of conservation and restoration has raised much interest in investigating the structure and physical-chemical properties of the materials used in art works[1]. Particularly, organic dyes and pigments are key components in cultural heritage artifacts, such as easel paintings, polychrome arts, manuscript illuminations, textiles, etc. Since color is an important component in the perception of beauty and in an artist's original intent when creating a work, better conservation of the cultural heritage requires detailed knowledge of dyes and pigments, including the complex evolution they have endured over time[2].

Both dyes and pigments have distinct features: dyes are soluble in the medium in which they are present, while pigments are solid state[3]. But organic dyes can be fixed on inorganic substrates to build organic-inorganic hybrid pigments, which sometimes exhibit remarkable stability. Lake pigments[2, 4-6] and Maya blue[7-9] are the typical hybrid pigments prepared from natural dyes and inorganic substrates. Particularly, a famous red lake, known as madder lake, was used for painting by precipitating the madder dyestuffs (extracted from the roots of *Rubia tinctorum* L) with a metal salt to form insoluble metal complexes since antiquity[10, 11]. According to historical recipes, the most frequently used metal cation was Al^{3+} , added to the dyestuff as a potassium double sulphate ($\text{KAl}(\text{SO}_4)_2 \cdot 12\text{H}_2\text{O}$, called alum)[12].

Alizarin (Az, see Fig. 1), the main component of madder dyestuffs, has been classified among the organic dyes most resistant to light-induced fading, although degradation still happens[13]. Az lake is often considered as a typical example of lake pigments to investigate the formation [14], spectral properties[1, 2, 5, 11, 12, 15-20] and lightfastness[21-24] by using modern and complementary technologies, such as gas chromatography, mass spectrometry, UV-visible spectrophotometry, thin layer chromatography, high performance liquid chromatography.

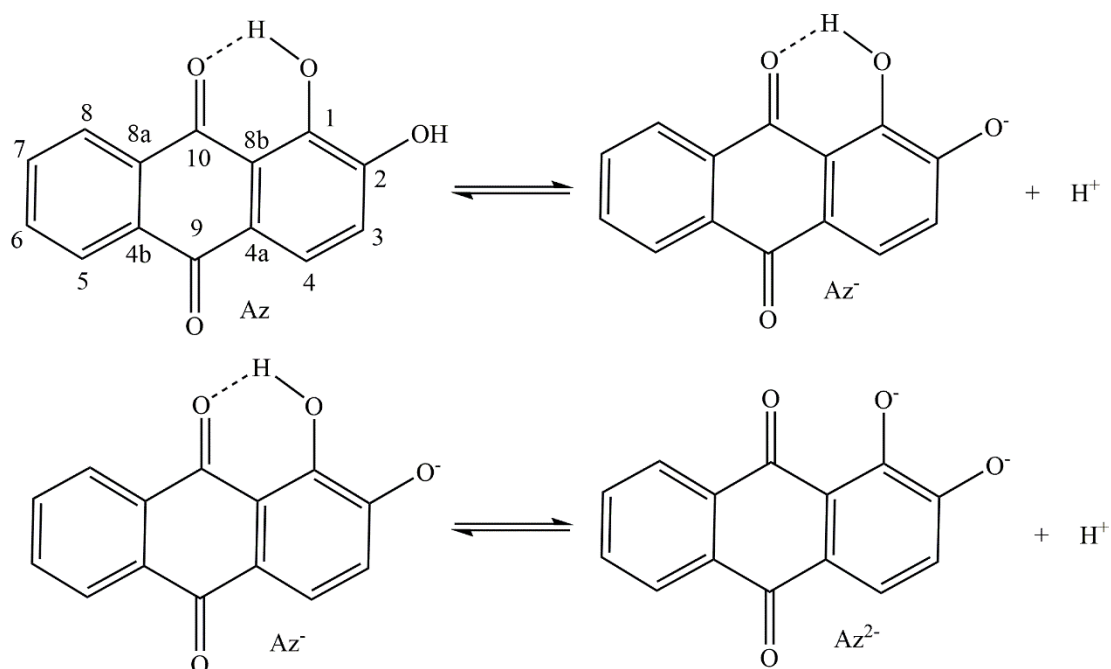


Fig. 1 The structure and ionization process of Az in aqueous solution

In 1920s, Weiser and Porter [25-27] pointed out that the lake formation is due to the adsorption of the Az anion by the hydrous oxides in varying amounts depending on the composition of the dye bath. Later, Kiel and Heertjes stated that the formula of the calcium-aluminum lake of Az (Turkey Red), i.e., $Ca_2Al_3AZ_4$ (where Az represents an alizarin molecule minus two hydroxylic hydrogen atoms), was wrong. Therefore,

they carried out a further study of this complex, and they illustrated the formula of the water-containing calcium-aluminum lake of Az is $\text{CaAl}(\text{OH})(\text{Az})_2 \cdot x\text{H}_2\text{O}$ [4]. The extremely slow reaction of alizarin with the chelate-forming metal aluminum alone due to the strong hydrogen bond between the 1-OH group and the 9-C=O group. This hydrogen bond has to be broken or at least weakened when a complex was formed. This phenomenon occurred when salt formation ($2\text{-OH} \rightarrow 2\text{-O}^-$) takes place before chelate formation [28]. The other similar Az lakes were also investigated by Kiel and Heertjes [29, 30].

In the beginning of the 21st century, the electronic structure of Az with eleven different transition metal complexes and five rare-earth complexes were studied using density functional theory (DFT) by Komih et al. [31]. The M–O (M refers to the metal cation) bonds involve *p* or *d* metal orbitals depending on whether the *d* shell is full or empty. The complexation effect leads to a red shift and hence to a color change of the solutions of the complexes. Theoretical calculation also demonstrated that complexation always occurs at the *peri*- or *ortho*-hydroxycarbonyl group and involves the tautomeric anthraquinoid forms [32].

Fain et al. (1996) [33] illustrated Az lake as two types of structure based on the ^{27}Al NMR spectra, named “closed structure” (with strongly chelated H_2O molecules and dibenzene sandwich Na^+) and “open structure”. Doskocz et al. (2009) [34] pointed out that Az acted as a bidentate ligand on forming complexes with metal ions based on the ^1H and ^{13}C NMR spectra results. The complex structure with the metal attached to two hydroxyl oxygen atoms was of lower energy. Wunderlich and

Bergerhoff [35] proposed a different structure in which the Al^{3+} cations coordinated with two Az molecules through the ortho-hydroxyl groups (α -diol position), forming a five-membered chelate cycle, and calcium ions complete the structure, coordinating two dye molecules in the β -ketol position. The recent investigation on the complexation reaction of Az, which had been carried out by Sanyova in 2001 [36] and 2008 [37], provided experimental evidence in favor of the β -ketol position of the dye as complexation site. In addition, his research also put forward the hypothesis of possible further aggregation of the complexes through Al–O–Al bonds and cross-linking.

Although much progress about the complex structure had been made in the past decades, the structure of Az lake and the nature of established chemical interactions are still a matter of controversy. Previous works only focused on the complex formation in solution, theoretical models and crystallized stoichiometric complexes. For example, the commercial lake pigments contained very low Az while the stoichiometric investigation always considered the complex structure with the Az/ Al^{3+} ratio of 1:1 and 2:1 [28, 32]. The structure and properties of a real Az lake pigment, with a very low content of Az dye, have never been studied.

What is more important, understanding the degradation pathway of Az lake is a crucial issue for the conservation and restoration of artworks which contain lake pigments. But it is difficult to evaluate the photostability and chemical changes of Az lake powders during light exposure. Dworak et al. (2009) [38] tried to investigate the photoinduced dynamics of Az in Al_2O_3 and TiO_2 mesoporous films that strongly

coupled with Az. Although the thickness of these films exceeded 1 μm , their research suggested that the interaction between Az and aluminum oxides might be an efficient way to study the structure and interaction of Az lake under light exposure. Aluminum oxides (Al_2O_x) exhibit the similar coordination with Al^{3+} complex in basic solutions. The use of Al_2O_x thin film to study the interaction between Az and Al^{3+} presents two essential advantages. First, Al_2O_x may be under the form of ultrathin layers and permit to investigate the interaction at the molecular scale. Previous quartz crystal microbalance measurements have demonstrated highly uniform layer-by-layer growth of the alumina on Ag nanoparticles with a growth rate of ~ 0.1 nm per deposition per second [39]. The sub-1-nm thickness is extremely advantageous in preserving integrity of the solid and increasing contact area and reactivity. In addition, the ultrathin film of alumina deposited on a reflective surface should enhance the sensitivity and accuracy of spectrometry [40], because only the surface atoms can react with Az molecules while the internal (bulk) atoms do not. In addition, polarization modulation infrared reflection absorption spectroscopy (PM-IRRAS) can be employed, with a very high sensitivity, to investigate the interfacial interaction between Az dye and Al_2O_x thin film.

Aiming to investigate the structures and interactions of Az lake pigments, a series of bulk Az lakes were prepared. The influence of dye concentration, pH and basic reagents on the structures and chromatic properties has been investigated. In addition, Az molecules were deposited onto Al_2O_x thin film prepared by physical vapor deposition (PVD) method. The interaction between Az and Al_2O_x thin film and

the influence of ultraviolet light on the degradation of Az-Al₂O_x film complex have also been highlighted.

2 Materials and methods

2.1 Materials

Az (97%), KAl(SO₄)₂·12H₂O (98%), K₂CO₃ (98%) and KOH (99%) were purchased from Sigma-Aldrich, Inc, France. These materials were used without further purification.

2.2 Preparation of bulk lake pigments

The lake pigments were prepared as the following procedure: KAl(SO₄)₂ (alum) was dissolved into water with the concentration of 0.5 mol/L; Az was dissolved into water with the concentration of 0.005 mol/L (pH=10); Az solution was added into alum solution with the molar ratio of Al/Az (R value) from 5 to 200; a basic reagent (KOH or K₂CO₃, 0.5 mol/L) was added into the previous solution dropwise after 10 min stirring; The addition was stopped when pH value reached at 5.5, 6.5, 7.5 and 8.5 corresponding to different speciation states of Az; The solution was stirred for 30 min and then stood for 12 h; finally, the precipitate was filtered out, washed 3 times and dried at 50°C for 24 h.

2.3 Preparation of aluminum oxide films

Preparation of aluminum oxide thin films was performed by physical vapor deposition (PVD) and a scheme of the apparatus is shown in Fig. 2(A) (details can be seen in reference [41]). An Al foil, source of Al atoms, was placed on a tungsten

filament connected to a DC power supply. Au thin film of $1 \times 1 \text{ cm}^2$ and few hundreds of nm thickness on a glass substrate were exposed to flame for three times before use. The Au substrate was placed in the vacuum chamber equipped with a quartz crystal microbalance placed laterally. Once the sample was inserted into the chamber, the vacuum was maintained for 2 h to reach a pressure of 10^{-3} Pa . Then the current was slowly increased to a few tens of A to allow Al evaporation and condensation on the Au film. The Al thin film was oxidized in air and a Al_2O_x thin film was formed.

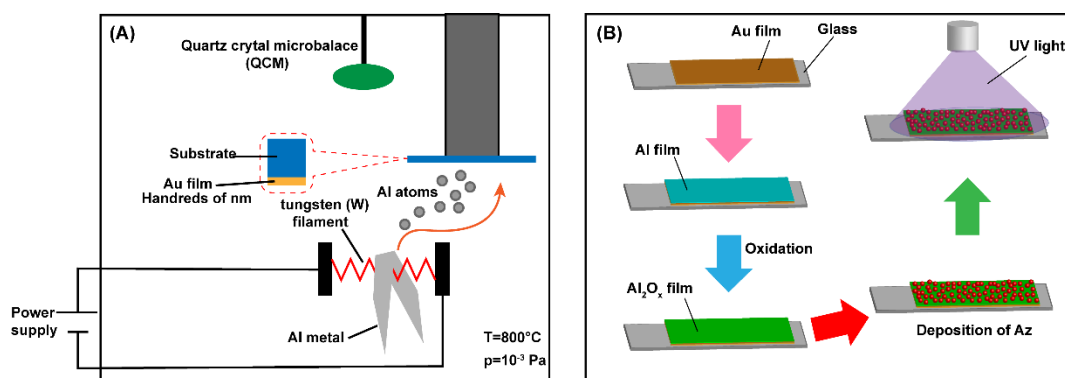


Fig. 2 (A) PVD apparatus for preparation of Al thin film on gold (Au) surface and (B) preparation process of alumina-Az complex samples.

2.4 Deposition of dyes onto the thin film

In order to study the properties of the dye on an aluminum oxide film surface, 20 μL of Az aqueous solution (0.5 mol/L) was dropped on the surface of the thin film prepared by PVD (see Fig. 2(B)). After 15 min, the excess of solvent was evaporated under Ar flux. To study the properties of the Dye- Al^{3+} complex on gold surface, a complex of Az- Al^{3+} was prepared in water by mixing Az and $\text{KAl}(\text{SO}_4)_2$ with Az/ Al^{3+} ratios of 1:1 and 2:1. The pH was adjusted to 7.5 by addition of KOH solution. 20 μL

of Az-Al³⁺ complex solution (0.5 mol/L) was dropped on the surface of the thin film previously exposed to flame for three times. The excess of solvent was evaporated under Ar flux after 15-min deposition.

2.5 Photostability

The thin film was exposed to UV light. The distance from the light source and the film was 10 cm. The UV light was the Lightningcure LC-L1V3 LED source from Hamamatsu Photonics with irradiance of 14000 mW/cm² at 365 ± 5 nm. The sample was characterized by PM-IRRAS after 0.5 h, 1 h, 2 h, 4 h, 7 h, 24 h, 33 h, 48 h, 76 h, 168 h and 264 h.

2.4 Characterization

Attenuated total reflection Fourier transform infrared spectrum (ATR-FTIR) spectra were collected from a Cary 630 FTIR spectrometer in the range of 4000-650 cm⁻¹.

CIE (Commission Internationale de L'Eclairage) parameters were obtained from an Ocean Optics spectrophotometer USB4000. The light source of Ocean Optics HL2000 was employed for CIE results. CIE 1976 color space system was applied to evaluate the color of the pigments. Lightness (L*) represents the brightness (+) or darkness (-) of the color, i.e., more positive L* values refer to whiter while more negative L* values represent darker. The values of a* and b* indicate the color details: +a* is the red direction, -a* the green direction, +b* the yellow direction, and -b* the blue direction. Before the reflectance and CIE tests, the samples were hand-pressed

into pellets using 3 mm die set and a metal collar for Quick Press from Perkin Elmer (ref. 01862295) with the pressure about 10 MPa.

^{13}C and ^{27}Al magic angle spinning (MAS) nuclear magnetic resonance (NMR) spectra were obtained on a Bruker Advance III spectrometer equipped with a 2.5 mm and 4 mm H-X MAS probe, operating at frequency of 125.76 MHz (^{13}C) and 130.30 MHz (^{27}Al). Chemical shifts were calibrated using the carboxyl signal of adamantane (38.52 ppm) for ^{13}C and $\text{Al}(\text{NO}_3)_3$ (0 ppm) for ^{27}Al as external standard. The ^{13}C cross-polarization (CP) spectra were acquired with a MAS rate of 14 kHz, a ramp-CP contact time of 2 ms and a 2 s of recycle delay and with a ^1H decoupling spin. Over an acquisition time of 40 ms, the number of scans to obtain the spectra depends on the signal to noise ratio obtained for each sample. Spectra were processed with a zero-filling factor of 2 and with an exponential decay corresponding to a 10 Hz line broadening in the transformed spectra. Only spectra with the same line broadening are directly compared. The ^{27}Al experiment using 30° pulse was recorded in the same condition of spinning rate (14 kHz), with a number of scans of 2048, 263 kHz spectral width and with 500 ms of recycle delay.

PM-IRRAS spectra were recorded using a Nicolet 5700 spectrometer equipped with a nitrogen-cooled MCT wide-band detector. The incident beam was modulated between p and s polarization using a ZnSe grid polarizer and ZnSe photoelastic modulator placed before the sample. The spectrometer was interfaced to the UHV chamber via ZnSe windows. The reflected light was focused onto the detector at an optimal incident angle of 85° . All spectra were obtained after 512 scans with a

resolution of 8 cm^{-1} .

3 Results and discussion

3.1 Influence of the Az concentration in lake pigments

3.1.1 Chromatic analysis

The CIE parameters (Table 1) of lakes pigments witness their colors. In the case of $R=*$, the precipitation shows L^* , a^* and b^* values of 99.7, 0.2 and -1.6, respectively, indicating a white color for the substrate Al^{3+} -X complex (Fig. 3). With the addition of Az, the CIE parameters of lake pigments changed. The lightness (L^*) of lake pigments decreased while the red ($+a^*$) and blue ($+b^*$) color components increased. Interestingly, the a^* values of lake pigments are much larger than b^* values, indicating that the lake pigments display red color (Fig. 3). The color of Az aqueous solution is readily affected by the pH value. In solid state, Az is yellow powder. However, the color of Az aqueous solution changes from yellow to red, and further to violet and blue with the increase of pH value. While, in the solid state, Az is a yellow powder, the color of Az aqueous solutions changes from yellow to red, and further to violet and blue, when increasing the pH value. In a basic solution, the chemical equilibrium of Az ionization was shifted to the right, generating more Az^- and Az^{2-} anions, resulting in a red color. Hence, the Al^{3+} -Az complex also displayed a red color. However, the real lake pigments were prepared with a very low concentration of Az, suggesting that only a part of Al^{3+} cations were coordinated with Az and others were still under the $AlO(OH)$ complex form. With the increase of Az concentration, the L^*

value of lake pigments increased while the a^* and b^* values were almost stable. Hence, the concentration of Az influenced the color nature of these lake pigments by changing their lightness. Less Az resulted in a brighter color.

Table 1 CIE parameters of lake pigments prepared with Az.

R	L^*	a^*	b^*
*	99.7	0.2	-1.6
5	33.9	43.6	8.5
25	39.5	43.5	10.9
50	45.6	45.6	10.5
100	53.9	41.5	8.7
200	61.6	40.6	7.8

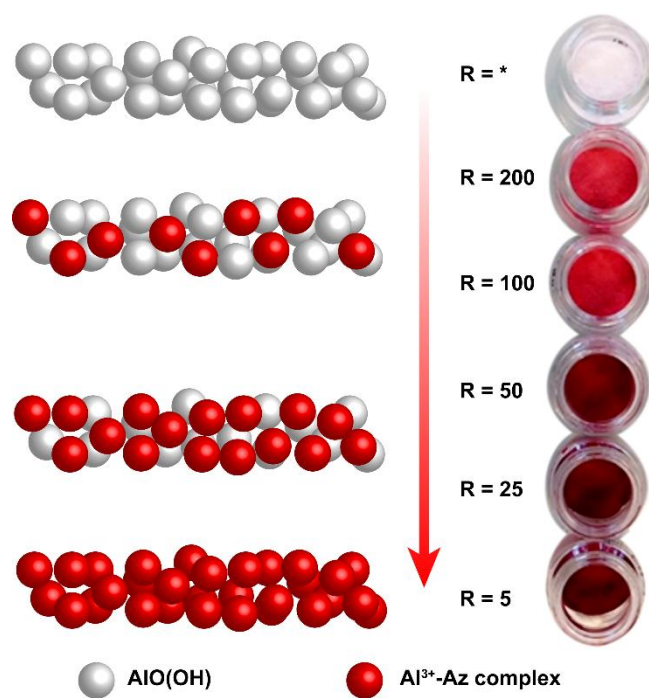


Fig. 3 Digital pictures of lake pigments and corresponding interpretive diagrams.

3.1.2 ATR-FTIR

The FTIR spectrum of Az is presented in Fig. 4. In agreement with previous reports [4, 42, 43], the assignments are summarized in Table 2. The absorption band at 3359 cm^{-1} belongs to the vibration of phenolic hydroxyl groups. The characteristic bands are $\nu(\text{C}=\text{O})$ at 1661 cm^{-1} (C10) and $\nu(\text{C}=\text{O})$ at 1631 cm^{-1} (C9); $\nu(\text{C}=\text{C})$ of benzene ring at 1584 and 1449 cm^{-1} , in addition to out of plan deformation bands at 1194 , 1179 and 1158 cm^{-1} , and group bands at 1047 , 1031 , 1012 cm^{-1} ; $\delta(\text{O}-\text{H})$ at 1349 cm^{-1} (C2) and 1329 cm^{-1} (C1); $\nu(\text{C}-\text{O})$ at 1280 cm^{-1} (C2) and 1263 cm^{-1} (C1).

In the case of R = * (without the addition of Az), four bands are observed in the FTIR spectrum. A wide band ($3000\text{-}3750\text{ cm}^{-1}$) centered at 3414 cm^{-1} is attributed to the stretching vibration of O-H from coordinated hydroxyl group (Al-OH) and water, corresponding to the bending vibration at 1104 cm^{-1} ($\delta(\text{Al}-\text{OH})$) and 1638 cm^{-1} ($\delta(\text{H}_2\text{O})$), respectively[44-46]. This result demonstrated that both -OH and H_2O coordinated with Al^{3+} . Furthermore, the signals of $\nu(\text{Al}-\text{O}-\text{Al})$ at 978 cm^{-1} and $\nu(\text{Al}-\text{O})$ at 852 cm^{-1} are observed, indicating the formation of a polymeric Al^{3+} complex. The lake pigment substrate is loosely described as hydrated alumina, but in practice is difficult to characterize, partly because it is amorphous and highly variable state [47]. It does not correspond to any of the various crystalline aluminum (oxy)hydroxides or anhydrous oxides, many of which are known mineral species such as gibbsite $\text{Al}(\text{OH})_3$ or boehmite ($\gamma\text{-AlOOH}$)[48-52]. The bands at 1508 and 1417 cm^{-1} are assigned to the stretching vibration of CO_3^{2-} group, implying that carbonates were included in the substrate.

When Az was added (R in the range 5 to 200), the FTIR spectra of samples present some shifts in comparison with that of Az. Firstly, the absorption bands referring to Az show lower intensity with the increase of R values. This phenomenon not only demonstrates the immobilization of Az molecules in the precipitation, but also testifies that smaller R value results in more Az included in the lakes. Secondly, compared with the FTIR spectrum of Az, the bands referring to C=C, C=O, and C—O and O—H groups shifted to higher or lower wavenumbers in the lake spectra. The two bands referring to the two C=O groups merged at 1638 cm^{-1} . The $\nu(\text{C}=\text{C})$ band from benzene ring splits into four bands at 1590, 1519, 1469 and 1435 cm^{-1} . The bands referring to $\delta(\text{O}—\text{H})$ shifted to 1355 and 1321 cm^{-1} from 1349 and 1329 cm^{-1} . The C2 $\nu(\text{C}—\text{O})$ signal shifted to 1287 from 1280 cm^{-1} . This fact indicates the interaction between Az and Al^{3+} . The bands representing the vibrations of $\delta(\text{Al}—\text{OH})$ at 1104 cm^{-1} , $\nu(\text{Al}—\text{O}—\text{Al})$ at 978 cm^{-1} and $\nu(\text{Al}—\text{O})$ 852 cm^{-1} remained in the ATR-FTIR spectra of Az lakes, demonstrating no changes happened to the substrate. But these bands exhibited lower intensity with the increase of R value, because of the low concentration of Al^{3+} (high concentration of Az). Particularly, the lake pigments with R=50 and 200 showed the remarkable signal of Al—OH group. This fact demonstrates that less Az molecules results in more Al—OH groups. The Az competed with —OH for coordinating with Al^{3+} .

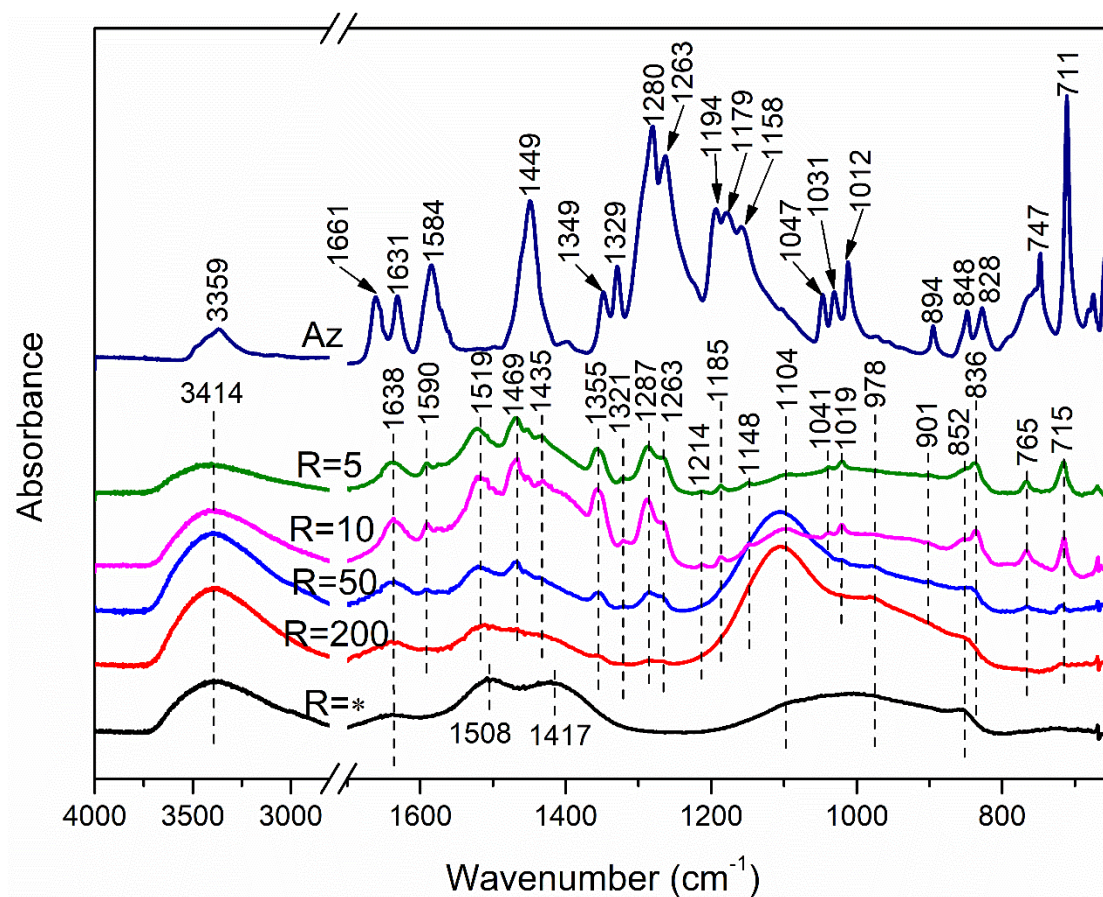


Fig. 4 ATR-FTIR spectra of Az and lakes with different R values.

Table 2 The assignments of FTIR bands of Az and the corresponding shifts in lakes.

Absorption band (cm^{-1})		Assignment
Az	Lake pigment	
3359		$\nu(\text{O-H})$
1661		$\nu(\text{C=O})$ C10
	1638	$\nu(\text{C=O})$ C9/ $\delta(\text{O-H})$ water
1631		$\nu(\text{C=O})$ C9
1584	1590	$\nu(\text{C=C})$
	1519	$\nu(\text{C=C})$

	1469	$\nu(\text{C}=\text{C})$
1449	1435	$\nu(\text{C}=\text{C})$
1349	1355	$\delta(\text{O}-\text{H})$ phenol C2
1329	1321	$\delta(\text{O}-\text{H})$ phenol C1
1280	1287	$\nu(\text{C}-\text{O})$ C2
1263	1263	$\nu(\text{C}-\text{O})$ C1
1194/1179/1158	1185/1148	$\nu(\text{C}=\text{C})/\delta(\text{CCC})/\delta(\text{C}-\text{H})$
	1104	$\delta(\text{Al}-\text{OH})$
1047/1031/1012	1019	$\nu(\text{C}=\text{C})/\delta(\text{CCC})/\delta(\text{C}-\text{H})$
894/848/827		$\gamma(\text{C}-\text{H})/\delta(\text{C}=\text{O})/\gamma(\text{C}=\text{C})/\delta(\text{CCC})/\gamma(\text{C}-\text{H})$
	978	$\nu(\text{Al}-\text{O}-\text{Al})$
	852	$\nu(\text{Al}-\text{O})$
748	766	$\delta(\text{CCC})$
712	715	$\gamma(\text{C}-\text{O})/\gamma(\text{C}=\text{O})$

Note: ν —stretching vibration, δ —bending in-plane, γ —bending out-of-plane.

3.1.3 NMR

The ^{27}Al MAS NMR spectra of Az lake pigments (Fig. 5 (A)) reveal the coordination nature of Al^{3+} in Az lakes. Three signals are observed in the spectrum of precipitation without Az ($R = *$), i.e., Al(VI) (hexa-coordinated Al^{3+}) at 7.1 ppm, Al(V) (penta-coordinated Al^{3+}) at 36.1 ppm and Al(IV) (tetra-coordinated Al^{3+}) at 66.7 ppm [53]. Comparing the area of NMR signals by integration, 89% of Al(VI), 7% of Al(V)

and 4% of Al(IV) were included in the substrate. Previous work also proved the presence of Al(VI) in the lakes. However, the authors ignored the minor presence of Al(V) and Al(IV) in spite of the ^{27}Al NMR spectra indicating weak signals. The most Al(VI) demonstrated major octahedral $[\text{AlO}_6]$ structure in the framework of inorganic substrate, indicating a polymeric complex. On the other hand, the presence of Al(V) and Al(IV) suggested some defects in the complex structure. The lake pigments (R=25 and 50) displayed similar ^{27}Al MAS NMR spectra with the precipitation without Az, demonstrating the similar coordination environment of Al^{3+} in the lake pigments. Most of Al^{3+} cations are coordinated with 6 O atoms, but some are coordinated with 5 or 4 O atoms. Thus, the introduction of Az did not influence the coordination structure of Al^{3+} complex.

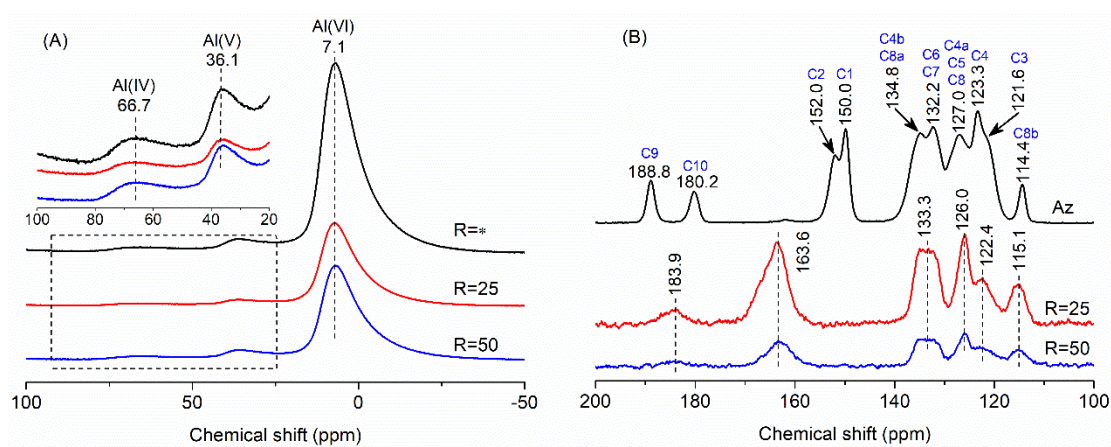


Fig. 5 (A) ^{27}Al and (B) ^{13}C MAS NMR spectra of Az lake pigments.

The ^{13}C MAS NMR spectra of Az and lake pigments are presented in Fig. 5 (B). The chemical shifts of C1 is smaller than that of C2, due to the hydrogen bond (Fig. 1). Accordingly, C9 shows higher chemical shift than C10, because the intramolecular hydrogen bonds decrease the electron density of C9 by delocalization. Compared with

Az, the NMR spectra of lake pigments exhibit some shifts. The chemical shifts of Az remained in the lake pigments, except for the signals of C1, C2, C9 and C10. Both chemical shifts of C1 and C2 increased to 163.6 ppm from 150.0 and 152.0 ppm, implying that the electron density of C1 and C2 decreased when Az interacted with Al^{3+} . C9 and C10 signals shifted to 183.9 ppm in lake pigments. This suggested the conjugative effect, which averaged the electron distribution in Az molecules coordinated with Al^{3+} .

Az has two phenolic hydroxyl groups, which can release protons (Fig. 1). Two acid-basic equilibria happen when Az is in aqueous solution. Kuban and Havel (1973) [39] reported the $pK_{a1} = 5.25$ and $pK_{a2} = 11.50$ of Az. Later, Miliani et al. (2000) [9] reported the pK_{a1} and pK_{a2} values as 6.57 and 12.36, respectively. Hence, pH value directly influences the proportion of Az, Az^- , Az^{2-} and H^+ . Besides, two C=O groups are also included in an Az molecule. Thus, Az can be considered as a bidentate ligand [4, 32-35, 54, 55]. For metal ion fixation, Az presents two potential sites in competition: the 1,2-dihydroxyl (catechol) and the 1-hydroxy-9-keto groups [34]. If a monodentate fixation mode could be considered, bidentate complexes are usually the most stable ones, for such ligands. Two contradictory hypothesis are proposed in the literature, an Al^{3+} fixation with the catechol group [55, 56] and a metal chelation with the 1-hydroxy-9-keto group [4, 28, 31]. Both chemical shifts of C1 and C2 increased from 150.0 and 152.0 ppm to 163.6 ppm while the chemical shifts of C9 (188.8 ppm) and C10 (180.2 ppm) equalized to 183.9 ppm. This fact demonstrates that Al^{3+} cations are fixed with the catechol groups in Az molecules, instead of 1-hydroxy-9-keto

groups. This result is in accordance with a recent report[34]

There are 6 possible coordinated structures of Al^{3+} - Az^{2-} complexes (Fig. 6). Al^{3+} cations coordinated with O atoms from H_2O or OH^- , i.e., $[\text{AlX}_6]$, $[\text{AlX}_5]$ and $[\text{AlX}_4]$ (X refers to H_2O or OH^-). The structures of lake pigments can be described as the replacement of Al-X complexes by some Az^{2-} anions. $\text{C}-\text{O}^-$ (C1 and C2) groups donated the electrons to the unoccupied orbitals of Al^{3+} , resulting in the chemical shift of C1 and C2 to higher values. In Al^{3+} - Az^{2-} complexes, the distribution of electrons became delocalized due to the coordination with Al^{3+} , resulting in the identical chemical shift of C1 and C2, as well as C9 and C10.

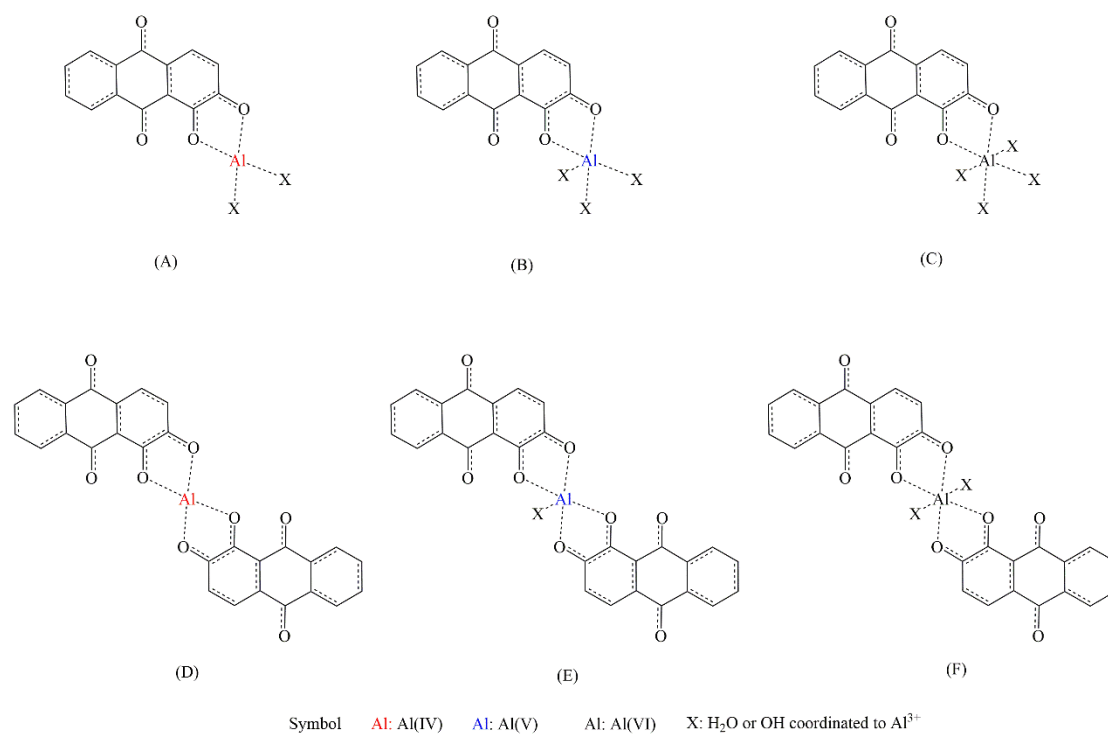


Fig. 6 The possible coordinated structures of $\text{Az}-\text{Al}^{3+}$ complexes: (A) $\text{Al}[\text{AzX}_2]$, (B) $\text{Al}[\text{AzX}_3]$, (C) $\text{Al}[\text{AzX}_4]$, (D) $\text{Al}[(\text{Az})_2]$, (E) $\text{Al}[(\text{Az})_2\text{X}]$ and (F) $\text{Al}[(\text{Az})_2\text{X}_2]$.

3. 2 Influence of basic reagents

3.2.1 Chromatic analysis

The CIE parameters of lake pigments prepared with K_2CO_3 and KOH are listed in Table 3. At the same pH value, the lake pigments prepared with K_2CO_3 showed lower L^* and b^* values, but larger a^* value than those of lake pigments prepared with KOH. Hence, K_2CO_3 resulted in redder but less bright color than KOH. The pH value also affected the color of lake pigments. With the increase of pH value, the L^* decreased while the a^* and b^* value increased, suggesting that the increase of pH value leads to a lower brightness and blue color. The bases and pH value influenced the color of lake pigments, with minor differences between K_2CO_3 -lakes and KOH-lakes (Fig. 7).

Table 3 Influence of basic reagents on the CIE parameters of Az pigments prepared at different pH values.

pH	L^*		a^*		b^*	
	K_2CO_3	KOH	K_2CO_3	KOH	K_2CO_3	KOH
5.5	51.2	55.7	41.8	41.8	7.0	10.9
6.5	44.7	53.1	47.5	45.1	9.4	15.8
7.5	44.9	56.4	47.0	39.7	8.5	10.7
8.5	45.4	47.5	48.4	43.3	9.3	12.9

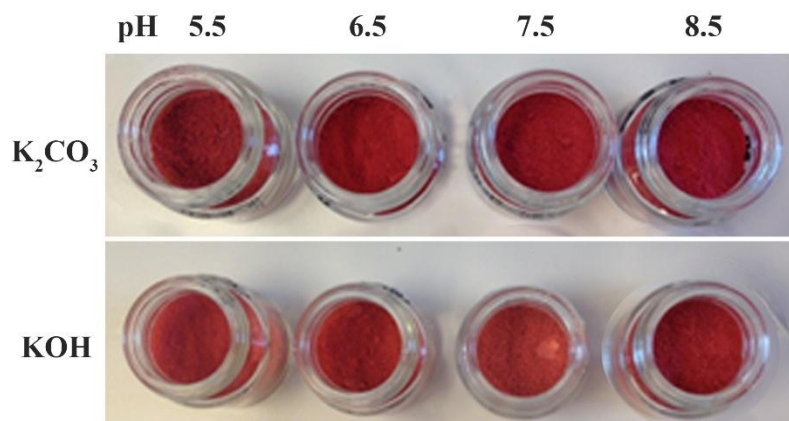


Fig. 7 Digital pictures of lake pigments.

3.2.2 ATR-FTIR

The precipitation of Al³⁺ is not only resulting from carbonate (K₂CO₃) but also from alkali (e.g. KOH). The ATR-FTIR spectra of lake pigments prepared with K₂CO₃ and KOH are given in Fig. 8. In Fig. 8(A), the band at 3396 cm⁻¹ refers to the stretching vibration of coordinated H₂O, whose bending vibration appeared at 1639 cm⁻¹. In addition, the ν(Al—OH) signal was overlapped by the wide band at 3396 cm⁻¹, corresponding to the δ(Al—OH) at 1103 cm⁻¹. The band at 1103 cm⁻¹ is assigned to the bending vibration (in-plane) of Al—OH group, the band at 974 cm⁻¹ is attributed to Al—O—Al group, and the band at 854 cm⁻¹ belongs to Al—O groups. In addition, the bands at 1499 and 1430 cm⁻¹ are assigned to CO₃²⁻ groups coordinated to Al³⁺. Carbonates are absent in the precipitation prepared at pH=5.5 while they present at pH value above 6.5. This demonstrated that carbonates were included in the precipitations when using K₂CO₃ as the base reagent. However, carbonates can only be present at pH > 6.5, because they are instable in acidic condition. The precipitation at pH = 5.5 showed a great signal at 1103 cm⁻¹ (Al—OH) and 974 cm⁻¹ (Al—O—Al),

while the band at 850 cm^{-1} (Al—O) was very weak. With the increase of pH value, the intensities of Al—OH and Al—O—Al groups decreased dramatically while the signal of Al—O group increased. This phenomenon implied that polymeric $\text{Al}^{3+}\text{-X}$ complexes formed at a low pH value, e.g., $\text{pH} = 5.5$, while individual chelates tended to be synthesized at pH above 6.5.

Compared with the FTIR spectra of precipitates prepared with K_2CO_3 , those of precipitates prepared with KOH (Fig. 8(B)) did not display the signals of carbonates. The bands of coordinated water and —OH group emerged at 3370 and 1640 cm^{-1} . The bands at 1093 and 1054 cm^{-1} are assigned to the symmetric and asymmetric bending vibration of Al—OH group. The signal at 976 cm^{-1} belongs to the stretching vibration of Al—O—Al group. The stretching vibration of Al—O group occurred at 880 cm^{-1} . With the increase of pH value, the intensity of absorption at 3370 and 1640 cm^{-1} decreased, as well as the signal of Al—OH group, suggesting the loss of —OH groups coordinated with Al^{3+} and/or coordinated water. In addition, the precipitations prepared with KOH showed stronger signal of $\nu(\text{Al—O—Al})$ than those precipitations prepared with K_2CO_3 , suggesting more —OH groups coordinated with Al^{3+} in the presence of KOH.

The lake pigments prepared with K_2CO_3 showed complex FTIR spectra of Az and precipitates (Fig. 8(C)). Except for the bands of substrate, the signals of Az were observed. The stretching vibration of C=O was overlapped at 1633 cm^{-1} . The stretching vibrations of benzene ring appeared at 1590 , 1518 and 1469 cm^{-1} . The bending vibration of phenol —OH group appeared at 1354 cm^{-1} . The wide band

between 1261 and 1285 cm^{-1} are assigned to the stretching vibrations of C1 and C2 C=O groups. The broad band between 1400 and 1500 cm^{-1} is assigned to the stretching vibration of CO_3^{2-} groups. Compared with the FTIR spectrum of Az, the lake pigments prepared with K_2CO_3 exhibited some shifts: (i) The bands referring to two $\nu(\text{C}=\text{O})$ vibrations at 1661 cm^{-1} (10-C=O) and 1638 cm^{-1} (9-C=O) merged and shifted to 1633 cm^{-1} ; (ii) The benzene ring $\nu(\text{C}=\text{C})$ at 1584 and 1449 cm^{-1} split and shifted to 1590, 1518 and 1469 cm^{-1} ; (iii) The $\delta(\text{O}-\text{H})$ at 1349/1329 cm^{-1} shifted to 1354 cm^{-1} , and the $\nu(\text{C}=\text{O})$ at 1281 and 1263 cm^{-1} merged into a broad band. These changes demonstrated the conjugative effect which averaged the electron distribution of Al^{3+} -Az complex. Particularly, we also noticed the wide band between 1400 and 1500 cm^{-1} assigned to CO_3^{2-} groups. The FTIR spectra of lake pigments prepared with KOH (Fig. 8(D)) are similar to those of lakes prepared with K_2CO_3 , except for the absence of the wide bands of CO_3^{2-} . The key difference between the lake pigments prepared with K_2CO_3 and KOH lies in the coordinated $-\text{OH}$ groups (1000-1200 cm^{-1}). With the increase of pH value, Az lakes prepared with KOH showed higher absorption than those prepared with K_2CO_3 , demonstrating more $-\text{OH}$ groups included in the complexes.

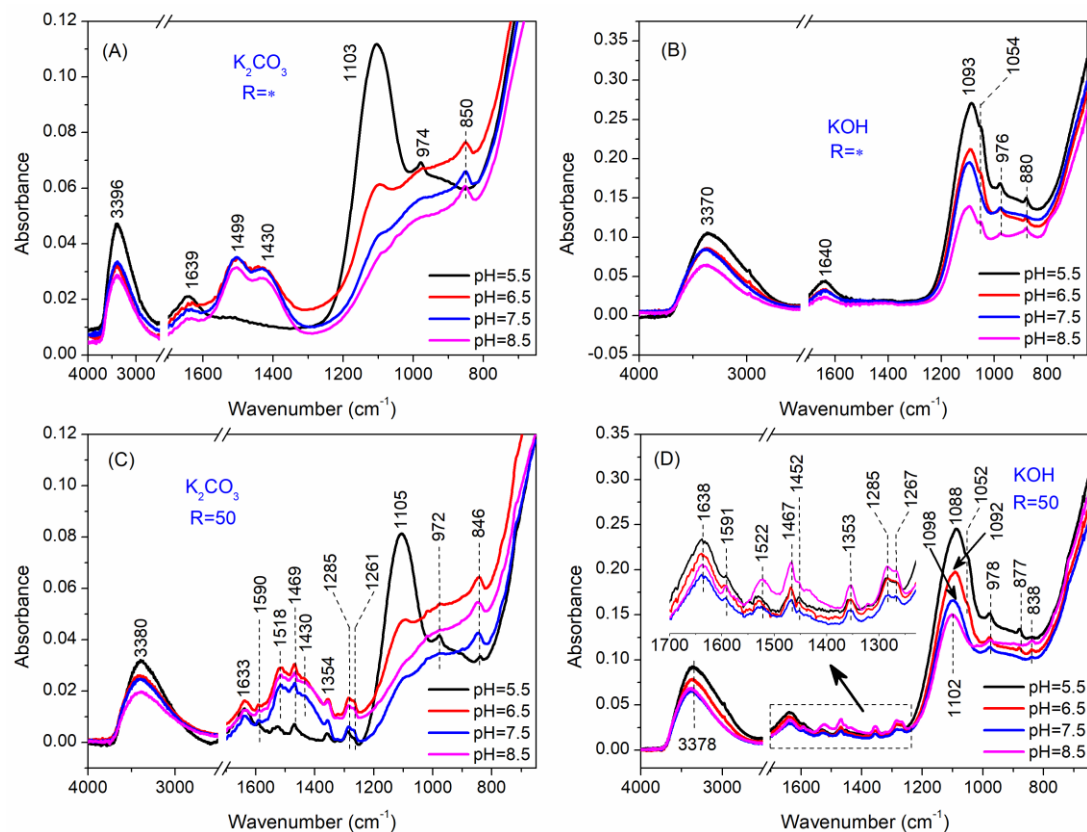


Fig. 8 ATR-FTIR spectra of lake pigments prepared with different bases.

3.3 Interaction between Al_2O_x film and dyes

The PM-IRRAS spectra of Al^{3+} -Az complexes on Au film (Fig. 9 (A)) indicate the interaction between Al^{3+} -Az complex and Au film. The Al^{3+} -Az complexes ($\text{Al}^{3+}/\text{Az} = 1:1$ and $1:2$) on Au films, i.e., $\text{Au-Al}^{3+}_1/\text{Az}_1$ and $\text{Au-Al}^{3+}_1/\text{Az}_2$, show characteristic bands of Al^{3+} -Az complex. The band at 1643 cm^{-1} can be attributed to the $\nu(\text{C}=\text{O})$ of Az and to the $\delta(\text{—OH})$ of coordinated water. The $\nu(\text{C}=\text{C})$ vibration of benzene ring emerged at 1593, 1531 and 1471 cm^{-1} . The band at 1361 cm^{-1} belongs to $\delta(\text{—OH})$ from phenol group. The absorptions at 1291 and 1268 cm^{-1} are assigned to the $\nu(\text{C—O})$. Az deposited on $\text{Au-Al}_2\text{O}_x$ film (Fig. 9(B)) also exhibit similar PM-IRRAS spectrum with Al^{3+} -Az complexes on Au film. This phenomenon demonstrates that Az molecules

interacted with Al_2O_x film surface very similarly to Al^{3+} -Az complexes. The band at 945 cm^{-1} is assigned to the stretching vibration of Al—O—Al groups, which belong to the Al_2O_x substrate. This band in Fig. 9 (A) is absent due to Au substrate, instead of Al_2O_x . But this phenomenon suggests that the deposition of Al^{3+} -Az complex did not form a polymeric structure like the bulk lakes, maybe because of the small amount (only on the surface and remove excess complex after 15 min).

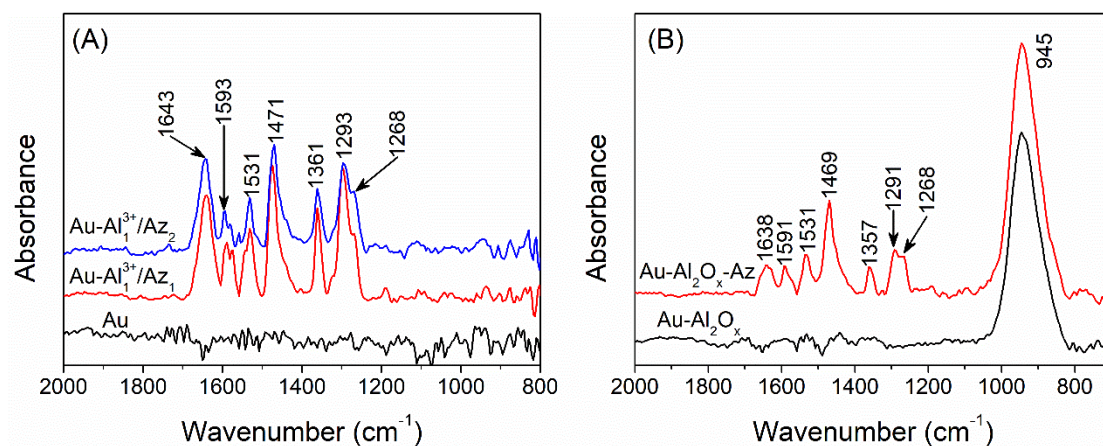


Fig. 9 PM-IRRAS spectra of (A) Al^{3+} -Az complex on Au film and (B) Az on $\text{Au-Al}_2\text{O}_x$ film.

Al—O groups exposed on the surface of Al_2O_x film and most Al^{3+} cations were coordinated by 6 O atoms. Compared with the FTIR spectrum of raw Az, both two phenolic C—O groups (1280 and 1263 cm^{-1}) of Az occurred at larger wavenumbers (1289 and 1267 cm^{-1}) in the hybrid of Az and $\text{Au-Al}_2\text{O}_x$ film. This demonstrates the interaction between the 1,2-dihydroxyl (catechol) and Al—O groups (Fig. 10). Az molecules substituted the coordinated O on the surface, resulting in a conjugative system.

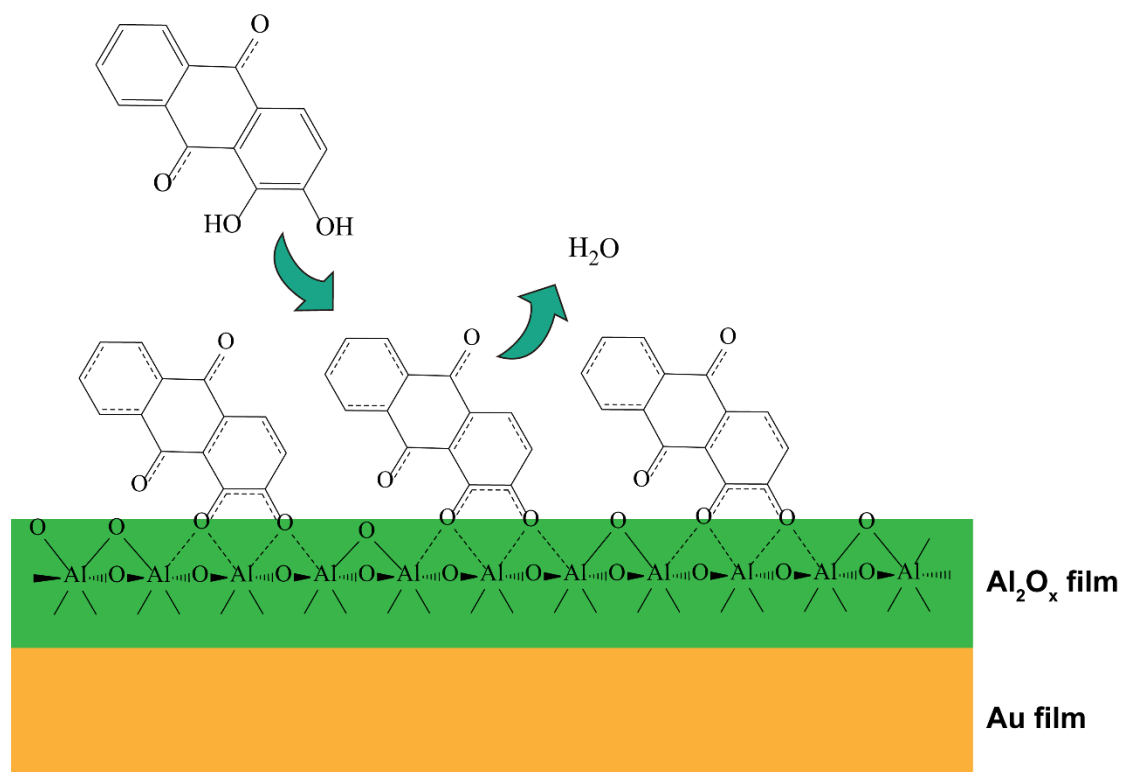


Fig. 10 Schematically interpretive diagram of Az dye on the Al_2O_x film.

3.4 Photostability

The above results demonstrated that Az deposited on the Au- Al_2O_x film surface showed similar interaction between Al^{3+} and Az in bulk lakes. Hence, the “film lake” can be employed to evaluate the photostability and degradation pathway. The Au- Al_2O_x -Az complex was exposed under UV light to evaluate the light resistance (Fig. 11). There are three changes after UV-light exposure: (i) Some new bands appeared at 1705, 1430, 1416, 1378 and 1102 cm^{-1} under UV light and their intensity increased with the exposure time; (ii) The bands at 1359 and 1327 cm^{-1} , corresponding to $\delta(\text{—OH})$ from phenol group decreased with the exposure time and finally disappeared; (iii) Some bands shifted to lower wavenumbers or decreased the

intensity with the UV light exposure time increasing, i.e., a) the band at 1644 cm^{-1} , referring to $\nu(\text{C}=\text{O})$ from Az, shifted to 1628 cm^{-1} , b) the band of $\nu(\text{C}=\text{C})$ from benzene ring at 1594 cm^{-1} shifted to 1573 cm^{-1} and the absorption at 1534 and 1469 cm^{-1} decreased, c) the bands of $\nu(\text{C}-\text{O})$ at C2 (1292 cm^{-1}) and C1 (1264 cm^{-1}) positions decreased their intensity with the UV light exposure, and finally the signal of C2 $\nu(\text{C}-\text{O})$ disappeared, d) the absorption bands corresponding to phenol $\delta(\text{-OH})$ (1359 and 1327 cm^{-1}) decreased the intensity and finally disappeared after 264-h UV light aging. We noticed that the absorption belonging to Al—O—Al group (945 cm^{-1}) remained.

The new band at 1705 cm^{-1} indicated the presence of C=O group under UV light aging. Considering the decrease and even disappearance of absorption bands referring to $\nu(\text{C}-\text{O})$, it can be concluded that C—O groups from Az molecules were oxidized to C=O groups (ketone or carboxylic acid). The decreasing intensity or even disappearance of absorption bands referring to C1 and C2 —OH (phenol) groups suggested the oxidation reaction happened to the phenol groups under UV light.

Previous investigation[57] proved that Az was bleached by H_2O_2 , which oxidized Az molecules from the phenol groups into ketones and finally the opening-ring reaction occurred, resulting in the generation of C=O and COOH groups. The emergence of absorption at 1102 cm^{-1} indicated the formation of Al—OH group. NMR results have demonstrated that Az anions, including Az^- and Az^{2-} , coordinated with Al^{3+} via catechol group. Hence, after long-time UV light exposure, the coordinated Az derivatives dissociated from the film surface due to the oxidation of Az derivatives

(see Fig. 12). Finally, the coordination of Az derivatives was substituted by $-OH$ group from humidity, or left the defect. The oxidation of phenol groups resulted in the change of conjugative status of Az derivatives, leading to the shifts of $\nu(C=C)$ from benzene ring to lower wavenumbers.

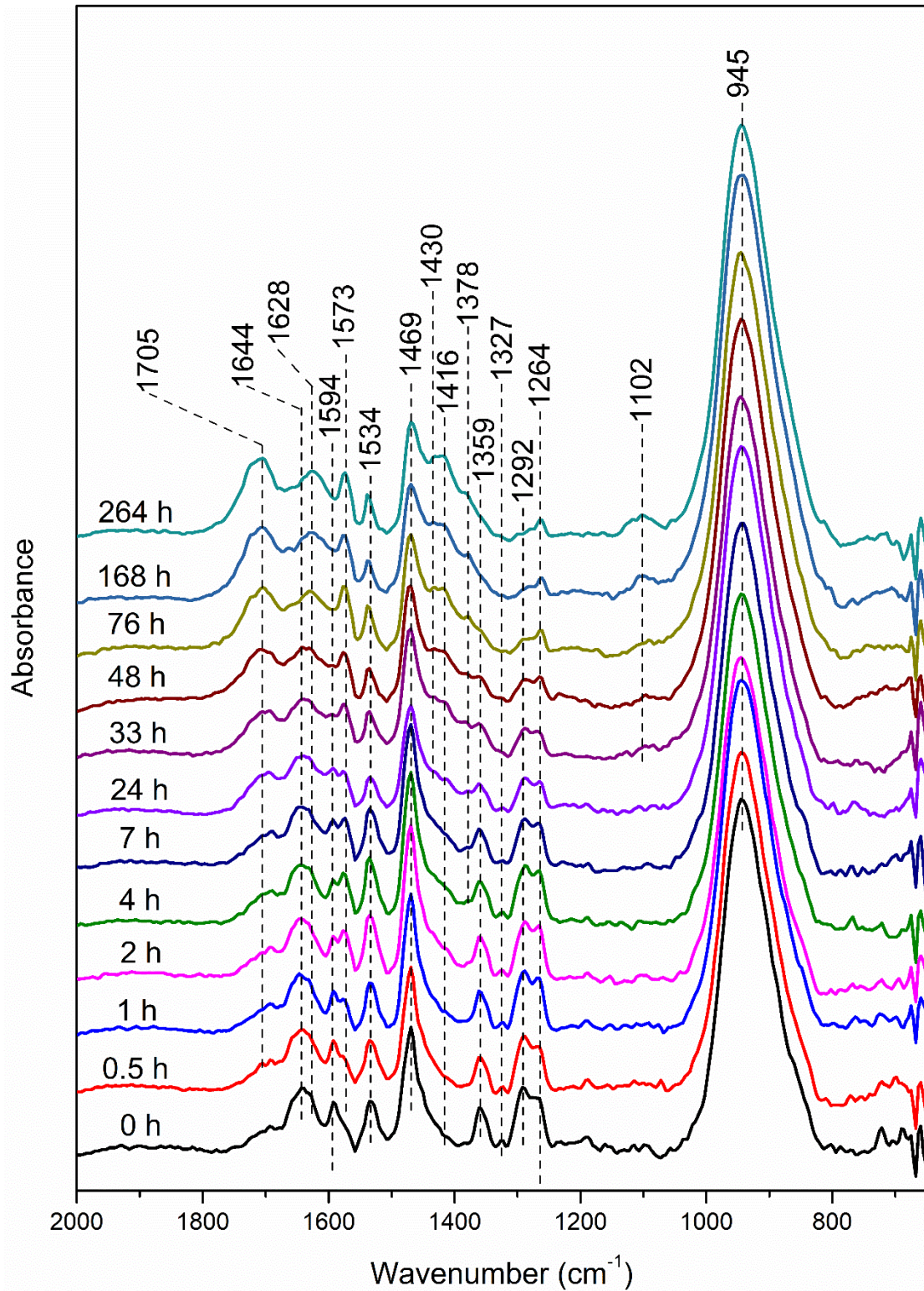


Fig. 11 PM-IRRAS spectra of UV light-aged Au-Al₂O_x-Az complex.

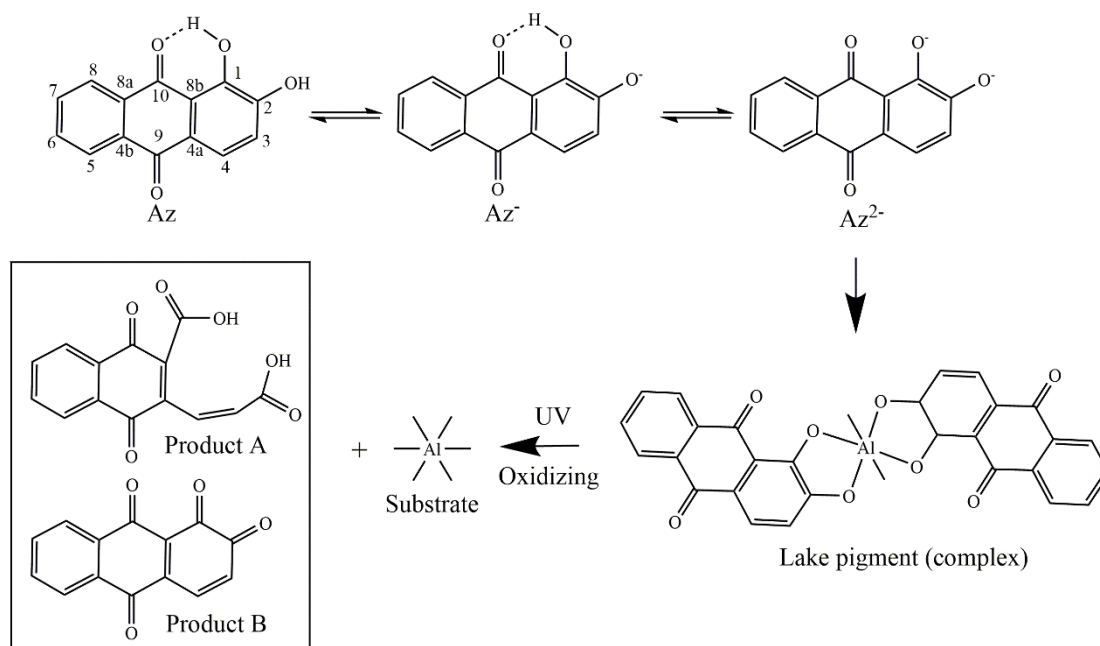


Fig. 12 The degradation pathway of Az in the lake pigments under UV light exposure. Az ionizes in water and donates protons, generating two derivatives, i.e., Az^- and Az^{2-} . The derivatives connect with Al^{3+} , forming complexes. The Az derivatives degraded exposed under UV light by oxidizing the C—O(H) into COOH or C=O group, breaking the conjugative connection between Az derivatives and Al^{3+} .

4 Conclusion

In this work, we investigated the structure and chromatic properties of non-stoichiometric Al^{3+} -Az complexes in bulk and the interaction between Az and Al_2O_x film, including the degradation under UV light exposure; this was made possible by combining complementary characterization techniques, and adapting samples for high sensitive measurements of reactivity. Several conclusions can be drawn. (i) Az lake pigment can be synthesized at a very low concentration of Az (even with the $R = 200$). The decrease of R value only effected the lightness of lakes,

without obvious influence on the a^* and b^* values, i.e. of the color. (ii) Al^{3+} formed complexes with Az derivatives by reacting with the 1,2-dihydroxyl (catechol) groups, and not with the 1-hydroxy-9-keto groups. Al^{3+} showed major hexa-coordination with minor penta-coordination and tetra-coordination structures. More important, the coordination between Az derivatives and Al^{3+} resulted in a conjugative system. (iii) The basic reagents, K_2CO_3 and KOH, did not affect the color of lake pigments. But the addition of KOH gave rise to more coordinated $-OH$ groups. (iv) Az derivatives formed conjugative complexes on the Al_2O_x film surface, similar with the bulk lakes. (v) The degradation pathway of Az derivatives on Al_2O_x film demonstrated the oxidation of catechol groups to carbonyl, resulting in dissociation of the conjugative system. This work presented a new insight into the structure and degradation of Az lake pigments. The fading of lake pigments is due to the oxidation of $C-O(H)$ groups from Az derivatives. Hence, adding some reductants or something are more sensitive to be photoexcited may be beneficial for the conservation and restoration of artworks.

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