

Fourier Transform Raman Studies of Methyl Red Adsorbed on γ -Alumina and Silica-Alumina

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Fourier transform Raman spectra of methyl red adsorbed on untreated and pretreated γ -alumina and silica-alumina calcined at 900 °C under 1 atm steam flowing were recorded. Spectral analysis shows that the active species adsorbed on γ -alumina was to be deprotonated methyl red, and on silica-alumina to be di-protonated. This indicates that γ -alumina adapted in this work holds Brønsted basicity, and silica-alumina Brønsted acidity. Raman intensities of methyl red on pretreated γ -alumina are about three times stronger than on untreated γ -alumina, while spectral features are unchanged. For silica-alumina, spectral features show modified vibrational characteristics upon surface hydroxylations generated from pretreatment. Consequently, the acidity loss for silica-alumina and the basicity gain for γ -alumina were observed by increasing the surface hydroxyl groups on the catalysts through pretreatment of the steam calcination.

Key Words : Raman, Methyl red, Calcination, γ -Alumina, Silica-alumina

Introduction

The structure of azo dyes has attracted considerable attentions recently due to their wide applicability in the light-induced photoisomerization process, and their potential usage for the reversible optical data storage.¹ Derivatives of 4-amino-*trans*-azobenzene are an important class of dyes for both natural and synthetic fibres. They are also widely used as acid-base indicators and as spectrophotometric probes in chemistry. Such dyes undergo a color change on protonation, due to changes in their molecular and electronic structure. Methyl Red (MR) is a weakly basic azo dye with $pK_{in} = 4.8$, and has been employed as a useful pH indicator in the range 4.4-6.0, in which it changes color from yellow in basic or neutral solution to red in acidic solution.² The chemical structure of MR is shown in Figure 1. MR may be a good probe for reference molecule of acid-base study because the spectral features of neutral, protonated, di-protonated, and deprotonated MR are distinct from one another in the vibrational spectra, and further they have been well studied in the Raman spectroscopy, especially in resonance Raman spectroscopy, so far. Resonance Raman (RR) studies of azobenzene and azonaphthalene derivatives have been reported and the key features of their spectra are well understood. The RR effect occurs when laser excitation is in close proximity to an intense electronic absorption band, in which case there may be a 10^3 – 10^4 fold enhancement of the normal Raman scattering signal.³ Usually, the enhancement factors of Raman band intensities in the resonance Raman scattering are quite selective to the modes and dependent on the electronic structure of molecules. However, advantages of using an excitation wavelength departed from electronic absorption bands, such as 1064 nm laser, are well under-

stood. Employing such excitation wavelength, one could expect to obtain the vibrational characteristics of the sample from nearly normal Raman scattering. RR spectra give richer information for assignments because they have symmetry information of normal modes and afford classifying a number of normal modes into different symmetry blocks.

The oxides of aluminum and silicon are catalytically important materials that have been investigated extensively in order to determine the nature of their adsorption behaviors. Investigations have been concentrated on the several

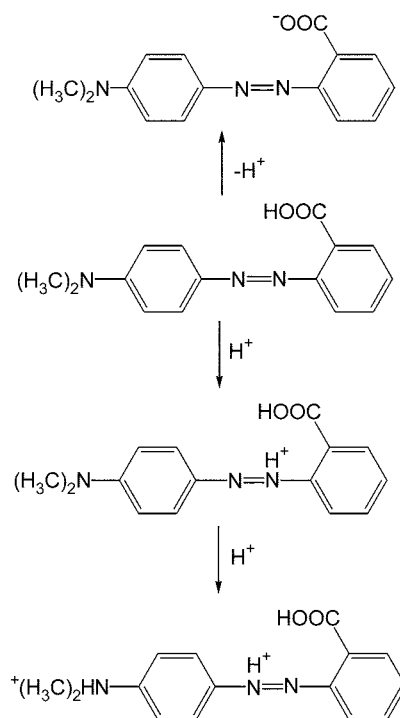


Figure 1. Chemical structures of MR species (from top: MRA, MR, MRH and MRHH).

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points, e.g., the geometry of sites (e.g., octahedral and tetrahedral environments of the highly defective spinel structures of aluminas), the type of bonding between substrate and adsorbate (e.g., physical adsorption whether of nonspecific nature due to dispersion forces or specific due to hydrogen bonding with surface hydroxyls), and the molecular dynamics of the adsorbed species (e.g., molecular diffusion and chemical exchange, which are important, for example, in cracking reactions).⁴ Aluminum oxide is an important technological material widely used in catalytic process. In particular, γ -Al₂O₃ is employed as a catalyst and a supporting material because of its acid-base properties, and its mechanical and high-temperature resistance. Surface properties of γ -Al₂O₃ have been the object of many experimental works using different probe molecules. Water adsorption has received special attentions since the basicity of this solid is generally related to the presence of surface hydroxyl groups. Under atmospheric conditions, the surfaces adsorb water both molecularly and dissociatively, giving rise to surface hydroxyls that persist even at temperatures higher than 1000 °C.⁵ The catalytic properties of amorphous silica-alumina catalysts are usually correlated with the Brønsted acidity that is located on the Al-OH-Si bridging bonds. Structural characteristics and reactivity properties of Al₂O₃/SiO₂ have been studied. Silica deposited on alumina is known to generate Brønsted acidity.⁶ Recently, these metal oxides have been revisited and investigated to improve the catalytic selectivity and kinetics for a specific reaction in the environmental chemistry⁷ and in the nano-particle chemistry.⁸

In this study, we investigate Raman spectra of MR adsorbed on γ -alumina holding basicity and on silica-alumina holding acidity to elucidate active species on the catalyst surface and the steam calcination effects. Raman spectra of neutral, mono-protonated, di-protonated, and deprotonated species of MR were also obtained to compare with the spectra from the MR adsorbed on the solids. It is well recognized that γ -alumina and silica-alumina can display both acidity and basicity, depending on pretreatment. Their acidic and/or basic sites may be classified as Lewis and/or Brønsted types. Steam calcinations of γ -alumina and silica-alumina were employed to investigate the acid-base types through the surface modification. By way of enforcing the hydroxyl groups on the surface of catalysts, it will be shown that the steam calcined silica-alumina decreased in the acidity strength, and the steam calcined γ -alumina increased in the basicity strength, and that γ -alumina adapted in this study has mainly Brønsted basic sites rather than Lewis types, while silica-alumina has mainly Brønsted acidic sites.

Experimental Section

MR was recrystallized from methanol and dried in vacuum. Toluene was dried over calcium chloride and distilled at atmospheric pressure. The protonated MR and deprotonated MR were prepared by mixing an equal or double molarity of HCl solution and an excess of NaOH solution, respectively.

Then dried in vacuum at room temperature. The γ -alumina was purchased of the 0.05 micron from Baikowski International Corporation (Charlotte, North Carolina, USA). The silica-alumina was purchased of silica-alumina catalyst support from Aldrich Chemical Company. Two sets of samples were prepared, containing untreated catalyst and pretreated catalyst calcined for 5 hours where the temperature was controlled at 900 °C under water-steam flow in the atmospheric pressure. MR solution of 150 μ M in toluene was prepared. MR solution of 25 mL was added to each 50 mg sample of γ -alumina and silica-alumina, and then mixtures were allowed to equilibrate for 24 hours. The supernatant solution of each sample was decanted, and then the samples were dried in vacuum at room temperature.

Bruker FRA106 Fourier transform spectrometer was applied in air purge mode to obtain Raman spectrum. It was equipped with 1064 nm cw Nd:YAG laser, a calcium fluoride beam splitter and liquid nitrogen cooled Ge detector for Raman scattering. The laser power was about 0.1 watt at sample. The spectral resolution was set to 4 cm⁻¹ and 100 times accumulated for each run. A sample powder was contained for irradiation with tight packing in ordinary melting point capillary tube (Drummond Scientific Co.).

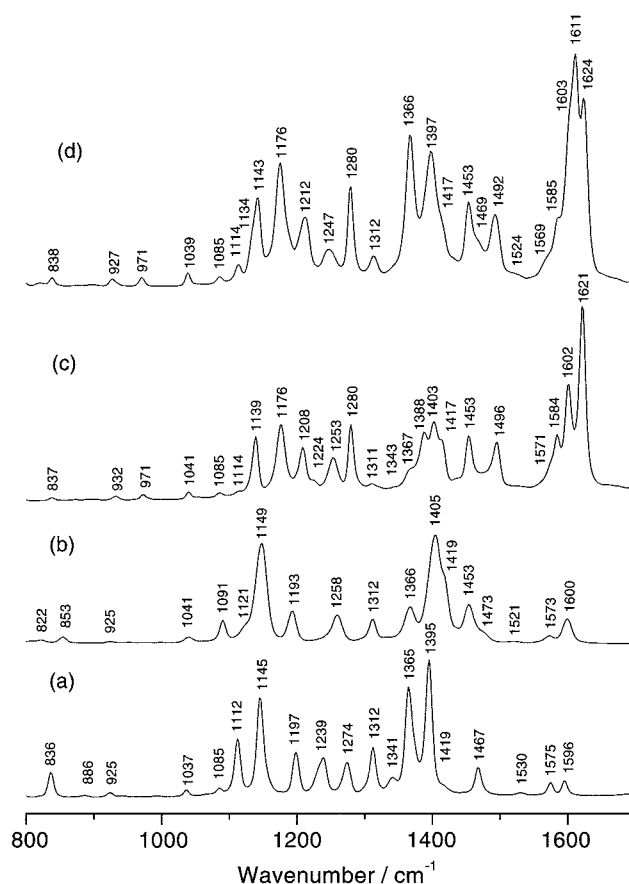


Figure 2. FT-Raman Spectra of Methyl Red (a), deprotonated Methyl Red (b), mono-protonated Methyl Red (c), and di-protonated Methyl Red (d). Excitation: 1064 nm Nd:YAG cw Laser, 4 cm⁻¹ resolution, 100 Scans coadded for each. All samples are in powder forms dried in vacuum.

Results and Discussion

Fourier transform Raman spectra in the mid-range of neutral (symbol: MR), deprotonated (symbol: MRA), mono-protonated (symbol: MRH), and di-protonated (symbol: MRHH) species of MR in the powder state are shown in Figure 2 and the band wavenumbers and their tentative assignments are given in Table 1, Table 2 and Table 3, respectively. These spectra are very similar to the resonance Raman spectra reported previously using 457.9 nm or 514.5 nm excitation except Raman intensity features.^{9,10,11} The spectrum of MR, Figure 2(a), shows a strong characteristic

Table 1. FT-Raman and resonance Raman bands and assignments for Methyl Red

FT Raman in Solid	Resonance Raman		Assignments
	in CH ₂ Cl ₂ ^a	in Methanol ^b	
836	836	836	δ_p (Ring-C)
886			δ_p (Ring-H)
925	923	922	ν_s (CNC)
1037		1040	ν (Ring Breathing)
1085		1086	δ (CH ₃ Rocking)
1112	1115	1118	δ_p (Ring-H)
1145	1150	1148	ν (C=N=), δ_p (Ring-H)
1197	1197	1198	ν (=N-C), δ_p (Ring-C)
1239	1240	1246	δ_p (Ring-H)
1274	1280	1282	δ_p (Ring-H)
1312	1313	1314	ν (amine N-C)
1341			δ (C-O-H)
1365	1368	1368	ν (Ring-C)
1395	1401	1402	ν (N=N)
1419			δ_s (CH ₃)
1467	1472	1460	ν (Ring-C)
1530			ν (Ring-C)
1575	1577	1576	ν (Ring-C)
1596	1602	1603	ν (Ring-C)

^afrom reference 11. ^bfrom reference 2

Table 2. FT-Raman and resonance Raman bands and assignments for Methyl Red Anion

FT Raman in Solid	Resonance Raman in KOH Solution ^a	Assignments
822		δ_p (Ring-C)
853		δ_p (Ring-H)
925	924	δ_s (CNC)
1041		ν (Ring Breathing)
1091	1090	δ (CH ₃ Rocking)
1121		δ_p (Ring-H)
1149	1150	ν (C=N=), δ_p (Ring-H)
1193	1196	ν (=N-C)
1258	1266	ν (=N-C), δ_{as} (CNC)
1312	1316	ν (amine N-C)
1366	1372	ν (Ring-C)
1405	1410	ν (N=N)
1419	1420	δ_s (CH ₃)
1453	1456	ν (Ring-C)
1473	1474	ν (Ring-C)
1521		ν (Ring-C)
1573	1580	ν (Ring-C)
1600	1602	ν (Ring-C)

^afrom reference 11

Table 3. Resonance Raman and FT-Raman bands and assignments for protonated MR derivatives, MRH and MRHH

Resonance Raman ^a of MRH in H ₂ O	FT Raman of MRH in Solid	FT Raman of MRHH in Solid	Assignments
840	837	838	δ_p (Ring-C)
926	932	927	δ_s (C-N-C)
972	971	971	δ_p (Ring-C)
	1041	1039	ν (Ring Breathing)
1084	1085	1085	δ (CH ₃ Rocking)
	1114	1114	δ_p (Ring-H)
		1134(sh)	
1144	1139	1143	δ_p (Ring-H)
1180	1176	1176	δ_p (Ring-H)
1192	1208	1212	δ_p (Ring-H)
1220	1224		
1250	1253	1247	ν (N=N), ν (=N-C)
1284	1280	1280	ν (=N-C), ν (N=N)
	1311	1312	ν (amine N-C)
	1343	(vw)	δ (C-O-H)
1366	1367	1366	ν (Ring-C)
1376	1388		ν (Ring-C)
1406	1403	1397	ν (C=N=)
	1417	1417	δ_s (CH ₃)
1456	1453	1453	ν (Ring-C)
1496	1496	1492	ν (Ring-C)
		1524	
	1571	1569	ν (Ring-C)
1588	1584	1585	ν (Ring-C)
1606	1602	1603	ν (Ring-C)
		1611	ν (Ring-C)
1620	1621	1624	ν (Ring-C)

^afrom reference 11

N=N stretching band at 1395 cm⁻¹ which is down-shifted 6 cm⁻¹ to the resonance Raman band in solution previously reported.¹¹ A weak band at 1419 cm⁻¹ is assigned to the CH₃ symmetric deformation. Five bands at 1365, 1467, 1530, 1575 and 1596 cm⁻¹ are the ring stretching modes attributed to two substituted benzene rings in MR. On comparison with the spectra of MR, the spectrum of MRA, Figure 2(b), shows a strong characteristic N=N stretching band at 1405 cm⁻¹ which is up-shifted 10 cm⁻¹. The intensity of 1366 cm⁻¹ band attributed to the stretching of ring, ν (Ring-C) is decreased. It appears that the localized ring conjugation and N=N stretching in MRA species are rather stronger than neutral MR. The spectra of MRH and MRHH, Figure 2(c) and Figure 2(d), are markedly different from those of the MR and MRA.

Very strong bands at 1621 cm⁻¹ for MRH and 1611 and 1624 cm⁻¹ for MRHH are assigned to the stretching of ring vibration, ν (Ring-C) confirming the existence of quinonoid character of the ring system in the protonated species. The intensities of the peaks (1450 cm⁻¹ ~ 1600 cm⁻¹) attributed to the ring stretching vibration (Wilson modes 8a, 8b, 19a and 19b) are strong in protonated species comparing with those of the MR or MRA. The peak attributed to the ring breathing mode is present at 1041 cm⁻¹ in the spectra of protonated and anionic species, which are 4 cm⁻¹ up-shifted to 1037 cm⁻¹ in MR. The bands at 1253 and 1280 cm⁻¹ for MRH and 1247 and 1280 cm⁻¹ for MRHH assigned to

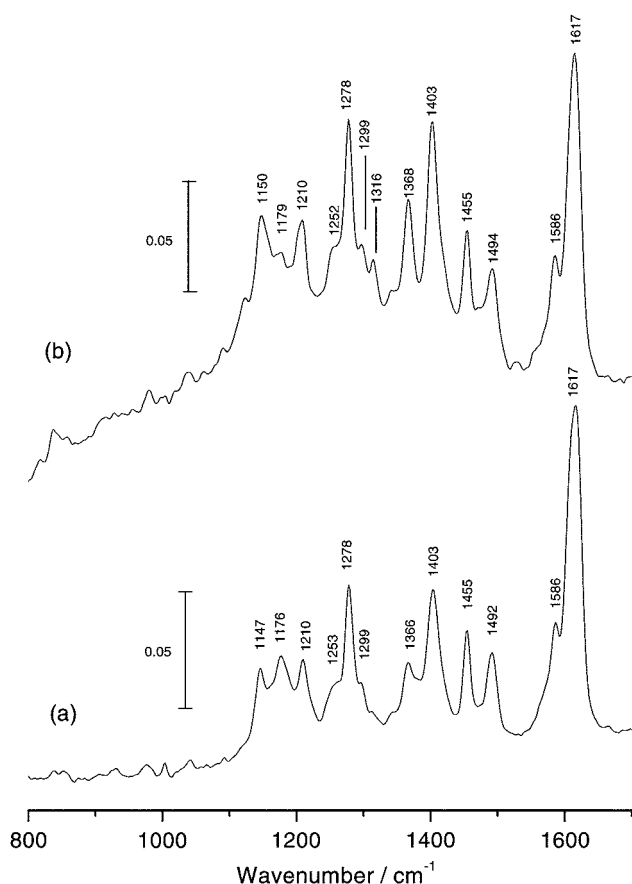


Figure 3. FT-Raman Spectra of Methyl Red Adsorbed on untreated silica-alumina (a) and on pretreated silica-alumina (b). Experimental conditions are the same as in Figure 2.

stretchings of both $\nu(\text{N}=\text{N})$ and $\nu(\text{=N-C})$ have shown that MR species is protonated to nitrogen atom of azo group. The double bond character between N and N is weaker and has some degrees of the single bond characteristics. Though the spectra of MRH and MRHH are looking nearly the same, MRHH has rather strong azo-dye character than MRH.^{10,11,12} This feature could be seen from the recovered strong bands at 1366 and 1397 cm^{-1} in Figure 2(d) which were observed strong in neutral MR, Figure 1(a), but not strong enough either in MRH, Figure 2(c) or MRA, Figure 2(b). These recovered features due to di-protonation in less 1400 cm^{-1} range for MRHH may show ideas of any conformational changes (*cis*-to-*trans*, or vice versa) induced by adsorption of MR on the colloid surface,¹³ or any structural changes (quinonoid-to-azo form, or vice versa) due to the inclusion of azonium group into cyclodextrin.⁹

The spectra of MR species adsorbed on untreated silica-alumina are shown in Figure 3(a) and the band wavenumbers and tentative assignments are given in Table 4. The spectra of MR species adsorbed on silica-alumina quite resemble those obtained for protonated MR, Figure 2(c). Very strong band at 1617 cm^{-1} is assigned to the $\nu(\text{Ring-C})$ vibration of the quinonoid structure adopted by protonated MR. The peaks in the region 1450 cm^{-1} to 1600 cm^{-1} are attributed to the ring stretching vibration (Wilson modes 8a, 8b, 19a and

Table 4. FT-Raman bands and assignments for MR species adsorbed on silica-alumina and γ -alumina

MR/Silica-Alumina		MR/ γ -Alumina		Assignments
Untreated	Calcined	Untreated	Calcined	
(974) ^a	(970) ^a			$\delta_p(\text{Ring-C})$
		1093	1094	$\delta(\text{CH}_3 \text{ Rocking})$
1147 (1145)	1150 (1146)	1150	1149	$\delta_p(\text{Ring-H})$
1176 (1172)	1179 (1172)			$\delta_p(\text{Ring-H})$
sh (1187)	sh (1188)			
		1193	1195	$\nu(\text{=N-C})$
1210 (1218)	1210 (1216)			$\delta_p(\text{Ring-H})$
1253 (1246)	1252 (1247)			$\nu(\text{N=N}), \nu(\text{=N-C})$
		1262	1264	$\nu(\text{=N-C}), \delta_{as}(\text{CNC})$
				$\nu(\text{=N-C}), \nu(\text{N=N})$
1278 (1274)	1278 (1275)			
1299 (1294)	1299 (1295)			
weak ()	1316 ()	1314	1314	$\nu(\text{amine N-C})$
1366 (1364)	1368 (1363)	1368	1368	$\nu(\text{Ring-C})$
1403 (1400)	1403 (1399)			$\delta(\text{C-N=})$
		1411	1411	$\nu(\text{N=N})$
		1422	1421	$\delta_s(\text{CH}_3)$
1455 (1452)	1455 (1452)	1459	1459	$\nu(\text{Ring-C})$
1492 (1490)	1494 (1489)			$\nu(\text{Ring-C})$
1586 (1576)	1586 (1582)			$\nu(\text{Ring-C})$
		1602	1602	$\nu(\text{Ring-C})$
1617 (1615)	1617 (1615)			$\nu(\text{Ring-C})$

^aThe values in the parentheses are resonance Raman bands from reference 14.

19b). Like protonation of MR, the substrate holds surface sites which can donate protons to adsorbates adsorbed on the silica-alumina surfaces, resulting in protonation of adsorbed MR species. Consequently, the examination of the spectra reveals that the silica-alumina surfaces function as the Brønsted acidic sites. The spectra of MR adsorbed on calcined silica-alumina are shown in Figure 3(b) and the band wavenumbers and assignments are given in Table 4. The spectrum Figure 3(b) represents the properties of Brønsted acid and Brønsted base. On comparison with the spectra of MR adsorbed on untreated silica-alumina, major differences between them are weaker intensity of a band at 1179 cm^{-1} and stronger intensities of bands at 1316 cm^{-1} and 1368 cm^{-1} . These features are also observed in Figure 2(b) and Figure 2(c). For MRA shown in Figure 2(b), we can deduce strong delocalization through azo $\text{N}=\text{N}$ bond conjugated to two ring systems, but for MRH in Figure 2(c) a lot weaker $\text{N}=\text{N}$ bonding character and quinonoid structural characters in the ring systems. Characteristic features for MRH in resonance Raman or surface enhanced Raman were known to be strong two bands at 1176 and 1620 cm^{-1} , which was also verified in Figure 2(c). The other notable difference in Figure 3(b), rather stronger intensity of a band at 1368 cm^{-1} could mean relatively higher degrees of azo-dye character in the molecular system. It is evident from these spectra that the proportion of Brønsted acid site on calcined silica-alumina is smaller than on the untreated silica-alumina. But Brønsted base sites are partly formed on the calcined silica-alumina surfaces due to adsorption of hydroxyl ions during steam calcination, instead.

In different from the spectra of MR species adsorbed on silica-alumina, the spectra of MR species adsorbed on γ

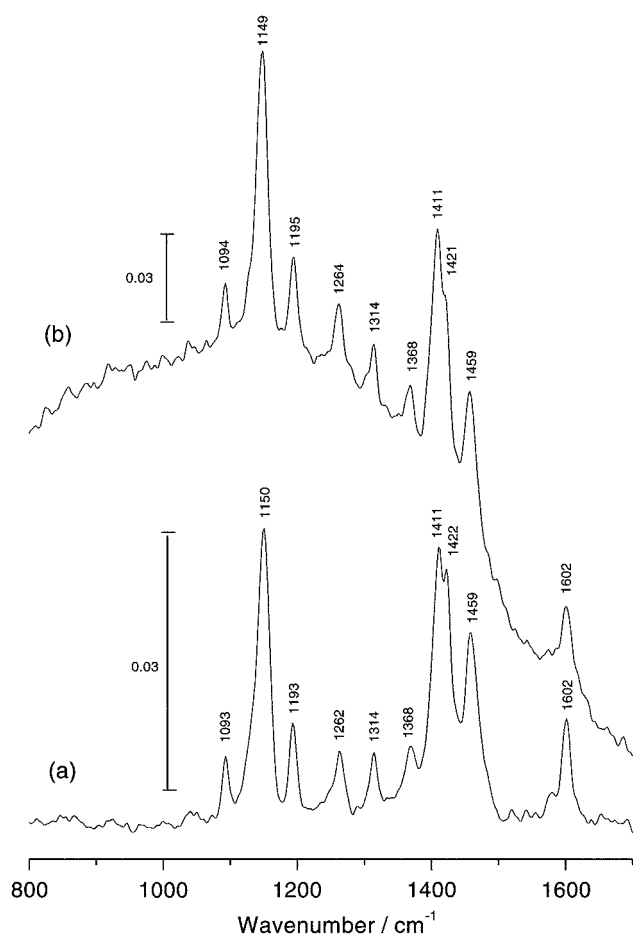


Figure 4. FT-Raman Spectra of Methyl Red Adsorbed on untreated γ -alumina (a) and on pretreated γ -alumina (b). Experimental conditions are the same as in Figure 2.

alumina closely resemble the spectra obtained for deprotonated MR. The spectra of MR species adsorbed on untreated γ -alumina are shown in Figure 4(a) and the band wavenumbers and tentative assignments¹⁴ are given in Table 4. For MR species adsorbed on γ -alumina the strong bands are at 1150 cm^{-1} attributed to δ_p (Ring-H) and ν (C-N=), and 1411 cm^{-1} assigned to the ν (N=N) vibration. On comparison with the spectrum of MR, the spectra of MR species adsorbed on γ -alumina have shown that the intensity of the band attributed to ν (Ring-C) is weaker and the intensity of the band attributed to ν (N=N) stronger. It appears that MR species adsorbed on γ -alumina is strongly conjugated further than in neutral MR. Usually alumina surfaces used at the majority of the catalytic reaction had been known to possess acidic sites, but in this adsorption study of MR it proves from spectral analysis that γ -alumina surfaces holds basic sites. Like deprotonation of MR, the sites deprotonate MR species adsorbed which could accept protons from adsorbates on γ -alumina surfaces. In this case, γ -alumina surfaces play a role as Brönsted basic sites. The spectra of MR adsorbed on calcined γ -alumina are shown in Figure 4(b) and the band wavenumbers and assignments are given in Table 4. The spectral features of Figure 4(a) and Figure 4(b) are nearly the same except spectral intensity. The intensity of MR species

adsorbed on calcined γ -alumina is much higher about three times than that of MR adsorbed on untreated γ -alumina. This observation can be explained as increased population of basic active sites in the calcined γ -alumina surfaces resulting from increased hydroxyl groups generated during steam calcination.

Conclusions

The active species adsorbed on γ -alumina was proved to be deprotonated methyl red, and on silica-alumina to be di-protonated methyl red. It is evident from the spectra analysis that hydroxyl groups are generated on the surfaces of oxides during steam calcination. Silica-alumina surface has Brönsted acidic site, and γ -alumina surface Brönsted basic site. Steam calcinations of γ -alumina and silica-alumina were employed to investigate the acid-base types through the surface modification. By way of enforcing the hydroxyl groups on the surface of catalysts, it is shown that the steam calcined silica-alumina decreased in the acidity strength, and the steam calcined γ -alumina increased in the basicity strength, and that γ -alumina adapted in this study has mainly Brönsted basic sites rather than Lewis types, while silica-alumina has mainly Brönsted acidic sites.

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