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M. A. Salim

King Fahd University of Petroleum and Minerals

G. D. Khattak

King Fahd University of Petroleum and Minerals

Petru S. Fodor

Cleveland State University, p.fodor@csuohio.edu

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Lowell E. Wenger

Wayne State University, wenger@physics.wayne.edu

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X-ray photoelectron spectroscopy (XPS) and magnetization studies of iron vanadium phosphate glasses

M.A. Salim , G.D. Khattak , P.S. Fodor , L.E. Wenger

Introduction

Studies of semiconducting oxide glasses containing mixed transition-metal (TM) oxides have been pursued in part because of the detection of mixed exchange pairs in these glasses [1–11]. For example, Bogomolova et al. [5,6] have shown that

a strong exchange interaction exists between Cu^{2+} and V^{4+} ions in the $\text{BaO-V}_2\text{O}_5\text{-CuO}$ and $\text{P}_2\text{O}_5\text{-CaO-V}_2\text{O}_5\text{-CuO}$ glass systems, which leads to the formation of mixed exchange $\text{Cu}^{2+}\text{-V}^{4+}$ pairs. The formation of these mixed clusters also appears to affect the electrical properties of these vanadate and vanadium phosphate glasses as electron hopping between the different TM ions can occur. Furthermore, the inclusion of the TM magnetic species in oxide glasses provides an opportunity to study these exchange pair formations through the interactions between the magnetic

ions. Electron paramagnetic resonance, magnetic susceptibility and Mössbauer spectroscopy have been particularly successful in studying the mixed exchange pair formation $\text{Cu}^{2+}\text{--V}^{4+}$ and $\text{Cu}^{2+}\text{--Mn}^{4+}$ in the $\text{B}_2\text{O}_3\text{--K}_2\text{O}$ glass system [8–10] and of $\text{Fe}^{3+}\text{--V}^{4+}$ in the $\text{BaO--V}_2\text{O}_5$ glass system [7].

In order to provide additional information on the interactions between unlike magnetic ions in vitreous matrices, magnetic susceptibility (χ) measurements in combination with X-ray photoelectron spectroscopy (XPS) and inductively coupled plasma spectroscopy (ICP) have been used to determine the ratio of different valence states of the V and Fe ions in vanadium phosphate glasses containing Fe_2O_3 . In these $\text{V}_2\text{O}_5\text{--P}_2\text{O}_5$ glasses, V_2O_5 is partially replaced by Fe_2O_3 and the possibility exists for magnetic interactions between the conjugate ions V^{4+} ($3d^1$) and Fe^{3+} ($3d^5$) or Fe^{2+} ($3d^6$). The extent and nature of this interaction can correspondingly be deduced from parameters associated with the observed Curie Weiss behavior ($\chi = C/(T - \theta)$) of these glasses.

Experimental procedure

Glass preparation

The glasses were prepared by melting dry mixtures of reagent grade V_2O_5 , P_2O_5 and Fe_2O_3 in alumina crucibles with the batch composition $[(\text{Fe}_2\text{O}_3)_x(\text{V}_2\text{O}_5)_{0.6-x}(\text{P}_2\text{O}_5)_{0.4}]$, where $x = 0.00, 0.10, 0.20, 0.25$ and 0.30 . Since the oxidation and reduction reactions in a glass melt are known to depend on the size of the melt, on the sample geometry, on whether the melt is static or stirred, on

thermal history and on quenching rate, all glass samples were prepared under similar conditions to minimize these factors [12]. Approximately 30 g of chemicals were thoroughly mixed to obtain homogenized batches. The crucible containing the batch mixture was placed in a furnace, heated at 300°C for 1 h and then transferred to an electrically heated melting furnace maintained at 1100°C . The melt was left for about 3.5 h under atmospheric conditions in the furnace during which the melt was occasionally stirred with an alumina rod. The homogenized melt was then cast onto a stainless steel plate mold to form glass buttons and glass rods of approximately 5 mm diameter for XPS measurements. The batch and actual compositions of the glasses studied are listed in Table 1 with the actual compositions determined by ICP spectroscopy.

Magnetization measurements

The temperature-dependent dc magnetic susceptibility was measured using a Quantum Design SQUID magnetometer in a magnetic field of 100.0 Oe over a temperature range 5–300 K. The susceptibility of the sample holder is negligible below 100 K for all samples and less than a 2% correction at the highest temperature for samples with a total susceptibility less than 1×10^{-8} emu/Oe. The overall accuracy of the magnetic measurements is estimated to be approximately 3% due to the uncertainty of the magnetometer calibration.

X-ray photoelectron spectroscopy (XPS)

High-resolution photoelectron spectra were collected on a spectrometer equipped with dual

Table 1
Batch and actual composition (molar fraction) of various vanadium phosphate glasses containing Fe_2O_3

	Batch			Actual (from ICP)		
	P_2O_5	V_2O_5	Fe_2O_3	P_2O_5	V_2O_5	Fe_2O_3
	0.40	0.60	0.00	0.398	0.602	0.000
	0.40	0.50	0.10	0.401	0.502	0.097
	0.40	0.40	0.20	0.407	0.402	0.191
	0.40	0.35	0.25	0.401	0.349	0.250
[A]	0.40	0.30	0.30	0.421	0.301	0.279
[B]	0.40	0.30	0.30	0.398	0.294	0.309

aluminum magnesium anodes using Al K α radiation ($h\nu = 1486.6$ eV) as described elsewhere [13]. The electron analyzer was set at a bandpass energy of 20 eV with the energy scale of the spectrometer calibrated using the core level of Cu 2p $_{3/2}$ (932.4 eV) and the energy separation between Cu 2p $_{3/2}$ and Cu 2p $_{1/2}$ (19.8 eV). The charging of non-conducting glass samples was avoided by flooding the sample with a separate source of low-energy electrons. The energy and intensity of these external electrons were adjusted to obtain the best resolution as judged by the narrowing of the full-width at half-maximum (FWHM) of the photoelectron peaks. At the optimum settings of the neutralizing gun (electron kinetic energy between 5 and 10 eV and electron emission current at the sample between 1 and 5 nA), the position of the adventitious C 1s line was within ± 0.5 eV of 284.6 eV. This peak arises due to hydrocarbon contamination and its binding energy is generally accepted as remaining constant, irrespective of the chemical state of the sample. For the sake of consistency, all energies are reported with respect to the C 1s transition at 284.6 eV. Glass rods of approximately 5 mm diameter were broken by vacuum cleavage inside the preparation chamber of the spectrometer in order to obtain fresh glass surface. The typical time required to collect an XPS spectrum for a sample was about 4 h and the base pressure in the analysis chamber during these measurements was less than 5×10^{-11} mbar.

Results

A typical, low-resolution X-ray photoelectron spectrum for a Fe $_2$ O $_3$ vanadium phosphate glass sample ($x = 0.20$) is shown in Fig. 1. This wide-scan spectrum as well as similar spectra obtained for the other glass compositions was obtained in about 1 h using non-monochromatic Al K α . In addition to the photoelectrons and Auger transitions of the glass constituents, the C 1s transition is evident at 284.6 eV. Since this peak is associated with the hydrocarbon contamination and almost always present on samples introduced from the laboratory environment or from a glove box [14], it is used as an energy reference. The resulting peak

Fig. 1. A low-resolution XPS spectrum from a fractured surface of a Fe $_2$ O $_3$ vanadium phosphate glass with $x = 0.2$ obtained using Al K α radiation ($h\nu = 1486.6$ eV).

positions for the Fe 2p, V 2p, P 2p, P 2s and O 1s core levels based on this assignment are listed in Table 2.

Fig. 2 compares the high-resolution V 2p and O 1s core level spectra for these Fe $_2$ O $_3$ vanadium phosphate glasses with the spectra of V $_2$ O $_5$ and P $_2$ O $_5$ powders. While it is apparent that V 2p $_{3/2}$ and 2p $_{1/2}$ peak positions for the glass samples are essentially the same as that for the V $_2$ O $_5$ powder, the V 2p spectra for the glasses are significantly broader and more asymmetric at the lower energies. Such broadening of the V 2p spectra suggests the existence of some non-equivalent vanadium atoms. Since vanadium can exist in several different valence states [15], each V 2p $_{3/2}$ spectrum was decomposed into two Lorentzian Gaussian peaks with a linear sloping background by means of a least-square fitting program [16] as shown in Fig. 3. The resulting peak positions and the relative areas under the peaks determined from this fitting for each spectra are given in Table 3. The difference between the positions of the two peaks ($\Delta V 2p_{3/2}$) is about 1.1 eV, which is in agreement with the reported difference between the binding energies of V $^{5+}$ (in V $_2$ O $_5$) and V $^{4+}$ (in VO $_2$) [17]. In general, V $^{5+}$ compounds have higher binding energies than vanadium compounds having V ions in a lower oxidation state. Thus the lower binding energy and the higher binding energy peaks are assigned to V $^{4+}$ and V $^{5+}$, respectively, with the relative areas under the peaks representing the relative ratio of V $^{4+}$ and V $^{5+}$ being present.

Table 2

Peak positions (in eV) for the core levels Fe 2p, V 2p, P 2p, P 2s and O 1s relative to the C 1s peak (284.6 eV) and the FWHM for the O 1s level

x	Fe 2p _{3/2}	Fe 2p _{1/2}	V 2p _{3/2}	V 2p _{1/2}	P 2p	P 2s	O 1s	FWHM (O 1s)
0.00			517.16	524.56	133.61	190.95	530.91	1.70
0.10	712.06	725.41	517.17	524.56	133.58	191.03	531.19	3.03
0.20	711.77	725.37	517.25	524.72	133.56	191.02	531.18	2.38
0.25	711.65	725.25	517.32	524.67	133.32	190.98	531.17	2.12
0.30	712.08	725.49	517.51	524.76	133.46	191.03	531.40	2.10
P ₂ O ₅					135.37	192.87	[d] 531.88 [b] 533.59	2.06 (0.397) 2.22 (0.603)
V ₂ O ₅			517.38	524.88			530.32	
Fe ₂ O ₃	710.80	724.18						

The uncertainty in the peak positions is ± 0.10 eV and in FWHM is ± 0.20 eV. The two O 1s peaks for P₂O₅ represent the double-bonded oxygen sites [d] and the bridging oxygen sites [b] with the numbers in parentheses representing the relative areas under each peak.

Fig. 2. High-resolution O 1s and V 2p spectra for the Fe₂O₃ vanadium phosphate glasses, V₂O₅ and P₂O₅ powders.

It is also evident in Fig. 2 that the O 1s peak for P₂O₅ is very asymmetric and, in fact, consists of two overlapping peaks. It is well known that orthorhombic P₂O₅ is based on an assembly of discrete molecules consisting of four PO₄ tetrahedra with three oxygen atoms being shared by the three

Fig. 3. A high-resolution V 2p_{3/2} spectrum (points) for a Fe₂O₃ vanadium phosphate glass ($x = 0.2$) and the resulting peaks from a least-squares fitting routine (solid lines).

neighboring tetrahedra and the other being double-bonded to the phosphorus atom [18]. From charge density considerations, the non-bridging atom peak will be shifted towards a lower binding energy with respect to the bridging oxygen atom peak [19]. Thus after decomposing the O 1s spectra into two peaks, the lower binding peak at 531.9 eV is associated with the non-bridging oxygen (P=O bonds) and the higher energy peak at 533.6 eV is associated with the bridging oxygen (P–O–P bonds) for the P₂O₅ powder. In addition, the ratio of the areas under the P=O and P–O–P peaks should reflect the relative ratio of 2:3 for non-bridging oxygen atoms to bridging oxygen atom for every P₂O₅ molecule. The resulting determi-

Table 3

The peak positions (in eV) resulting from the curve fitting of the V 2p_{3/2} and Fe 2p_{3/2} peaks for the various glass samples

x	V 2p _{3/2}	Fe 2p _{3/2}
0.00	516.1 (0.165) 517.3 (0.835)	
0.10	516.6 (0.518) 517.8 (0.482)	711.9 (0.634) 714.2 (0.366)
0.20	516.7 (0.439) 517.7 (0.561)	711.8 (0.541) 713.3 (0.459)
0.25	516.7 (0.415) 517.7 (0.585)	711.7 (0.564) 713.9 (0.436)
0.30	516.8 (0.398) 517.9 (0.602)	711.6 (0.524) 714.0 (0.476)

The relative areas under each peak are in parentheses. The uncertainty in the peak positions is ± 0.1 eV.

nation of the relative areas as shown in Table 2 is in agreement with this prediction.

In comparison to the P₂O₅ powder spectrum, the spectra for the vanadium phosphate glass samples and the V₂O₅ powder are much more symmetric without any discernible evidence for more than a single peak. Furthermore, the FWHM of the O 1s spectra for the glasses are comparable to the FWHM deduced from fitting either of the two O 1s peaks for the P₂O₅ powder as shown in Table 2. Thus the data suggest that a 'single' type of oxygen bond dominates the local oxygen structure in the vanadium phosphate glasses even though the O 1s spectra for the glasses are broader than that for the V₂O₅ powder.

Fig. 4 shows the high-resolution Fe 2p spectra for the glass samples as well as the spectrum for Fe₂O₃ powder. Although the Fe 2p_{3/2} and 2p_{1/2} peak positions for the glass samples are approximately 1.2 eV higher in energy than that for the Fe₂O₃ powder, the binding energies are in good agreement with those reported for iron phosphate glasses [20]. Furthermore, since Fe can exist in either the Fe²⁺ or Fe³⁺ state, the Fe 2p_{3/2} peaks for the glass samples are also decomposed into two separate peaks as shown in Fig. 5. From this fitting routine, two peaks with average values of 711.7 and 713.8 eV are found (see Table 3) and are assigned to Fe²⁺ and Fe³⁺, respectively. This assignment is based on the binding energy of

Fig. 4. High-resolution Fe 2p_{3/2} spectra for the Fe₂O₃ vanadium phosphate glasses and Fe₂O₃ powder.

Fig. 5. A high-resolution Fe 2p_{3/2} spectrum (points) for a Fe₂O₃ vanadium phosphate glass ($x = 0.1$) and the resulting peaks from a least-squares fitting routine (solid lines).

Fe(III)₂O₃ being about 1.2 eV higher than that for Fe(II)O [21] and the strong agreement with the Fe 2p_{3/2} peak positions reported for Fe²⁺ and Fe³⁺ in the iron phosphate glasses [20]. The fitting routine further suggests that the ratio of the relative Fe³⁺

to Fe^{2+} areas increases with increasing iron content in these Fe_2O_3 vanadium phosphate glasses.

The variation in the binding energies of the P 2p and P 2s electrons has also been examined. Figs. 6 and 7 compare high-resolution XPS spectra of the P 2p and P 2s core levels for the glasses to that of P_2O_5 powder. Each spectrum appears to consist of a single peak whose width remains essentially unchanged; however, both the P 2p and P 2s peaks for the glasses shift by about 1.9 eV to lower energies as shown in Table 2. Also it should be noted that the binding energies of the P 2p and P 2s electrons for all glass samples is essentially independent of the Fe content.

The magnetic susceptibility results for the glasses are shown in Figs. 8 and 9 as plots of inverse mass magnetic susceptibility, χ^{-1} , as a function of the temperature, T . The data appear to follow a Curie Weiss behavior ($\chi = C/(T - \theta)$) over most of the temperature range. From the slope and intercept, the Curie constant, C_{expt} and paramagnetic Curie temperature, θ (K), are de-

Fig. 6. High-resolution P 2p spectra for the Fe_2O_3 vanadium phosphate glasses and P_2O_5 powder.

Fig. 7. High-resolution P 2s spectra for the Fe_2O_3 vanadium phosphate glasses and P_2O_5 powder.

Fig. 8. The inverse of mass magnetic susceptibility as a function of temperature for the Fe_2O_3 vanadium phosphate glasses in a magnetic field of 100 Oe.

termined for each of the glass samples (see Table 4). Both the Curie constants and the magnitudes of the paramagnetic Curie temperature increase with

Fig. 9. The inverse of mass magnetic susceptibility as a function of temperature for the Fe_2O_3 vanadium phosphate glasses for temperatures below 100 K. The solid lines represent Curie Weiss fits to the high-temperature data.

increasing Fe content in the glasses. Since the paramagnetic Curie temperature, θ , is a rough measure of the strength of the interaction between the magnetic ions in the sample, a higher value implies stronger interaction and/or more ions participating in the interaction. The negative values for the paramagnetic Curie temperatures in-

dicate that the magnetic interaction is predominately antiferromagnetic in these Fe_2O_3 vanadium phosphate glasses.

Discussion

As described in the preceding section, XPS provides information of the short-range structure in these glasses. For example, the O 1s spectrum for P_2O_5 in Fig. 2 was found to consist of two overlapping peaks centered at 533.6 and 531.9 eV with a ratio of relative areas of nearly 3:2. This is understood in terms of the higher binding energy being associated with the six bridging oxygen atoms (P–O–P type of bonds) in the P_4O_{10} structure and the lower binding energy with the four double-bonded oxygen atoms (P=O bonds). Following similar arguments for a difference in the binding energies between bridging and non-bridging oxygen sites, the spectrum for V_2O_5 powder should have two peaks as four of the oxygen atoms in the V_2O_5 crystal structure are double-bonded oxygen (V=O bonds) while only one is a bridging oxygen (V–O–V bond). The narrowness and symmetry of the O 1s peak for V_2O_5 powder suggest that the energy difference between V=O and V–O–V bonds

Table 4

Magnetic susceptibility results for the vanadium phosphate glass samples and comparison to calculated paramagnetic Curie constants based on ICP and XPS results to experimental results

	Molar fraction (nominal)			C_{expt} (emu K/g Oe)	θ_{expt} (K)			
	Fe_2O_3	V_2O_5	P_2O_5					
[A] [B]	0.00	0.60	0.40	7.35×10^{-4}	–0.97			
	0.10	0.50	0.40	6.09×10^{-3}	–21.4			
	0.20	0.40	0.40	1.03×10^{-2}	–40.4			
	0.25	0.35	0.40	1.35×10^{-2}	–57.9			
	0.30	0.30	0.40	1.53×10^{-2}	–63.7			
	0.30	0.30	0.40	1.61×10^{-2}	–69.5			
	Molar fraction (from ICP)			Ion fraction (from XPS)		MW (g/mol)	C_{calc} (emu K/g Oe)	C_{expt} (emu K/g Oe)
	Fe_2O_3	V_2O_5	P_2O_5	$\text{Fe}^{3+}/\text{Fe}_{\text{total}}$	$\text{V}^{5+}/\text{V}_{\text{total}}$			
	0.000	0.602	0.398	0.000	0.766	163.591	6.97×10^{-4}	7.35×10^{-4}
	0.097	0.502	0.401	0.366	0.482	158.570	6.10×10^{-3}	6.09×10^{-3}
	0.191	0.402	0.407	0.459	0.561	156.912	1.06×10^{-2}	1.03×10^{-2}
	0.250	0.349	0.401	0.436	0.585	155.746	1.34×10^{-2}	1.35×10^{-2}
	0.279	0.302	0.421	0.476	0.602	154.979	1.50×10^{-2}	1.53×10^{-2}
	0.309	0.294	0.398	0.476	0.602	154.849	1.65×10^{-2}	1.61×10^{-2}

may be too small to be observed, especially since the relative weight of the two types of bonds is 4:1. Similarly, no discernible second peak is observed in the O 1s spectrum for the Fe-free vanadium phosphate $[(V_2O_5)_{0.6}(P_2O_5)_{0.4}]$ glass sample, even though it is broader than that for V_2O_5 . Assuming that the local structure at this vanadium concentration maintains the P_4O_{10} structure, then the following types of bonds should exist with the corresponding relative weightings:

$$P=O \quad 4(1 - n_v) = 1.60, \text{ where } n_v = 0.60,$$

$$P-O-P \quad 8(1 - n_v)^2 - 2(1 - n_v) = 0.48,$$

$$P-O-V \quad 16n_v(1 - n_v) = 3.84,$$

$$V-O-V \quad 8(n_v)^2 - 2(n_v) = 1.68,$$

$$V=O \quad 4n_v = 2.40.$$

Thus less than 5% of the total number of possible oxygen bonds should be bridging oxygen bonds (P–O–P) in this highly concentrated vanadium phosphate glass and the remaining 95% should be non-bridging oxygen atoms (P=O, P–O–V, V–O–V and V=O) whose photoelectrons would give rise to the observed single O 1s peak. In principle, this O 1s peak could be decomposed into four smaller peaks representing each of these four types of non-bridging oxygen atoms; however, it would require a minimum of eight fitting parameters. Furthermore, the separation of the peaks might be indistinguishable as a previous XPS study on alkali phosphate glasses [22] showed that contributions in the O 1s spectra from the double-bonded oxygen atoms (P=O) and the ‘non-bridging’ oxygen atoms (P–O–Na) had an energy difference on the order of 0.1 eV or less. Thus, it is doubtful that any meaningful insight into the local structure would be gained from a four-peak fitting analysis, especially considering the narrowness of the observed O 1s peak. Instead, it is more reasonable to hypothesize that all of the non-bridging oxygen atoms have nearly the same binding energies. The addition of Fe_2O_3 into the vanadium phosphate glass structure will result in non-bridging Fe–O–P and Fe–O–V bonds being formed and replacing some of the non-bridging P–O–V, V–O–V and V=O bonds with little effect on the relative ratio of the P–O–P bonds to non-

bridging oxygen bonds. Thus a single O 1s peak should remain with minimal shift in the peak position (binding energy of the photoelectrons) as a function of increasing Fe_2O_3 concentration.

The 1.9 eV shifts to lower binding energies of the P 2p and P 2s electrons for the vanadium phosphate glasses are mainly due to a difference in the molecular environment surrounding the P atom between the P_2O_5 and Fe vanadium phosphate glass structures. While the next-nearest neighbors in P_2O_5 are all P atoms, there is a more than 50% probability that a V (or Fe) atom will be the next-nearest neighbor in these highly concentrated phosphate glasses. Since the electronegativities of V and Fe are lower than that of P (=2.1), V and Fe have less of an affinity for electrons than P. Thus the electron density at the P atom increases with the the presence of V/Fe atoms and leads to a decrease in P 2p and P 2s binding energies in these glasses. Furthermore, since the ratio of the V/Fe (~0.60) to P (~0.40) remains nearly constant for all glass samples, the P 2p and P 2s binding energies should show little variation with Fe content as observed in the XPS spectra.

In comparing binding energies of the $2p_{3/2}$ and $2p_{1/2}$ electrons of V and Fe in the phosphate glasses to standard TM oxide powders (see Table 2), the V 2p peaks for the glass samples and metal oxides are almost at the same positions while the Fe 2p peaks for the glass samples shift by approximately 1.2 eV towards higher binding energies. These shifts can result not only from a change in the molecular environment from V_2O_5 and Fe_2O_3 , but also from a reduction of the formal oxidation state of V and Fe in the glass structures [23]. Following a similar explanation to that given in the preceding paragraph, V and Fe atoms in the glass structures will be surrounded by a large number of P atoms instead of V or Fe atoms at the next-nearest neighbor positions. Thus the electron density of the V or Fe atoms would be smaller while their 2p binding energies would be greater. In contrast, binding energies typically decrease with a reduction in the oxidation state of a metal atom, e.g., $V^{5+} \rightarrow V^{4+}$ and $Fe^{3+} \rightarrow Fe^{2+}$. Consequently the change in the molecular environment must be the dominant mechanism responsible for

the shifts in the Fe 2p energies and for keeping the V 2p levels unchanged.

Ideally, magnetic susceptibility measurements should be able to provide an independent verification of the presence and amount of the various magnetic ions in these vanadium phosphate glass samples. However, the presence of three different magnetic ions, i.e., V^{4+} ($p_{\text{eff}} = 1.8 \mu_B$), Fe^{3+} ($p_{\text{eff}} = 5.9 \mu_B$) and Fe^{2+} ($p_{\text{eff}} = 5.4 \mu_B$), results in an overabundance of independent variables such that the magnetic susceptibility data cannot be further analyzed without additional constraints. By using the ratios of V^{4+}/V_{total} , $Fe^{3+}/Fe_{\text{total}}$ and $Fe^{2+}/Fe_{\text{total}}$ as determined from the XPS analyses and the molar fraction of each constituent as measured from ICP, a comparison between the expected (calculated) value of the Curie constant (see Appendix A) and the experimentally measured value for each sample can be made. As shown in Table 4, the calculated Curie constants are within 3% of the experimental values for all vanadium phosphate glass samples containing Fe ($x > 0$) and within 7% for the Fe_2O_3 -free glass sample ($x = 0$). Thus the strong correlation between the calculated and experimental Curie constants confirms the presence of the three magnetic ions, V^{4+} , Fe^{3+} and Fe^{2+} , in these vanadium phosphate glasses. Furthermore, the fact that the paramagnetic Curie temperatures, θ , for all glass samples are non-zero and negative indicates that there are exchange effects between the magnetic spins with the interaction being antiferromagnetic in nature. This is consistent with previous investigations on similar systems [24,25] reporting that the antiferromagnetic interactions are present in magnetic TM oxide glasses. Assuming that the

local structure of the glass is P_4O_{10} , the possibility exists for the cluster formation of dimers (two P sites being occupied by magnetic ions) and closed-trimers (three sites being occupied by magnetic ions in a closed triangle) with a ‘non-bridging’ oxygen atom between the magnetic ions. It is well known that such oxygen intermediaries provide an indirect exchange mechanism responsible for the antiferromagnetic ordering in magnetic insulators [26].

The observation of the paramagnetic Curie temperature, θ , becoming more negative with increasing Fe_2O_3 content in these glass samples implies that the interaction strength between the magnetic ions becomes greater and/or that more ions are participating in the interaction as $\theta \propto J \cdot z \cdot S$, where J is the interaction strength, z the number of magnetic nearest neighbors and S is the magnetic spin. Assuming a random probability for the magnetic ion substitution in the P_4O_{10} structure, 17% of the P_4O_{10} structures would not have any magnetic ions, 38% would contain a single magnetic ion, 32% a dimer, 12% a closed-ring trimer, and less 2% would have all P sites replaced with magnetic ions for the $x = 0.10$ glass sample (see Table 5). With increasing Fe content in the vanadium phosphate glasses, the number of dimers, trimers and ‘quads’ (all P sites occupied by magnetic ions) would increase while the number of the other two possible cluster configurations would decrease. Thus the average value of z and S would increase with increasing Fe_2O_3 content. Additionally, one could expect the exchange overlap and hence the effective interaction strength J to increase if the local P_4O_{10} structure has any contraction in the lattice spacing between the P

Table 5
Probability distribution of a given magnetic cluster type in the P_4O_{10} structure

Cluster type	Probability	Nominal molar fraction of Fe_2O_3					
		$x = 0.0$	0.10	0.20	0.25	[A] 0.30	[B] 0.30
No magnetic ions	$(1 - q)^4$	0.545	0.171	0.160	0.134	0.130	0.109
Single	$4q(1 - q)^3$	0.357	0.380	0.372	0.350	0.346	0.322
Dimer	$6q^2(1 - q)^2$	0.088	0.316	0.324	0.343	0.345	0.359
Closed-trimer	$4q^3(1 - q)$	0.010	0.117	0.126	0.149	0.153	0.178
‘Quad’	q^4	0.000	0.016	0.018	0.024	0.025	0.033

$q = N_{Fe} + N_V V^{4+} / V_{\text{total}}$ (= the total molar fraction of magnetic ions) where N_{Fe} and N_V are the molar fractions of Fe and V determined from ICP analyses of the glass samples.

sites with the substitution of more Fe on the P sites.

Finally, the deviation from the Curie Weiss behavior below ~ 15 K can be explained on the basis of the local P_4O_{10} structures consisting of single magnetic ions, dimers, etc. In the low-temperature regime ($T \ll J$), the magnetic susceptibilities for antiferromagnetic dimers and 'quads' would vanish to zero while that for the trimers would remain finite but small. However, the susceptibility of the non-interacting, single magnetic ions continues to increase as $1/T$. Consequently, the paramagnetic contribution of these non-interacting magnetic ions to the susceptibility will become the most dominant contribution and cause the downturn observed in the temperature dependence of the inverse susceptibility below 15 K.

Conclusions

By combining studies of the XPS spectra for O 1s, P 2p, P 2s, Fe 2p and V 2p core levels of the Fe_2O_3 vanadium phosphate glasses with magnetic susceptibility measurements, the local structure of these glasses and the nature of the interactions between the magnetic ions have been investigated. The O 1s spectra for the glasses show a single, symmetric peak which results from nearly 95% of the oxygen sites in the local P_4O_{10} structure being occupied by non-bridging oxygen atoms. The shifts in the binding energies of the Fe 2p (1.2 eV increase), the V 2p (no change) and both the P 2p and P 2s (1.9 eV decrease) core levels for these glasses in comparison those for their respective powders (Fe_2O_3 , V_2O_3 and P_2O_5) are explained in terms of changes in their electron densities as a result of differences in the next-nearest neighbor environment between the local structures in the glasses and powders. In addition, the V $2p_{3/2}$ and Fe $2p_{3/2}$ spectra for the glass samples are sufficiently broaden such that two peaks are fitted to the data. These peaks are associated with the presence of V^{5+} and V^{4+} and Fe^{3+} and Fe^{2+} , respectively, and the relative area under each peak reflects the relative amount of each ion. Approximately 40% of the iron is found

to be in the Fe^{3+} state and 50% of the vanadium in the V^{5+} state in the Fe_2O_3 vanadium phosphate glass samples. The presence and amount of each of these magnetic ions (V^{4+} , Fe^{3+} and Fe^{2+}) determined from the XPS results are in quantitative agreement with the analysis from the Curie Weiss behavior ($\chi = C/(T - \theta)$) in the magnetic susceptibility data on the glasses. The Curie constants, C_{calc} , calculated from the XPS and ICP measurements are within 3% of the experimentally measured values for the Fe-containing vanadium phosphate glasses. The negative paramagnetic Curie temperatures, θ , indicate that the interaction between the magnetic ions is antiferromagnetic in nature. Qualitative features in the magnetic susceptibility results including the deviation from the Curie Weiss behavior below 15 K and more negative paramagnetic Curie temperatures with increasing Fe content can be satisfactorily explained by considering the formation of magnetic dimers, trimers, etc., with the introduction of the magnetic ions onto the P sites of the P_4O_{10} structure. Furthermore, this provides a straightforward explanation for the indirect exchange coupling between $V^{4+}-Fe^{3+}$, $V^{4+}-Fe^{2+}$, $Fe^{3+}-Fe^{2+}$, etc., that give rise to the antiferromagnetic interactions.

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Appendix A

To calculate the Curie constants from the ratios of V^{4+}/V_{total} , Fe^{3+}/Fe_{total} and Fe^{2+}/Fe_{total} as determined from the XPS results, we will define

$r = V^{5+}/V_{total}$	ratio of number V^{5+} ions to total number of V ions from XPS
$(1 - r) = V^{4+}/V_{total}$	ratio of number V^{4+} ions to total number of V ions from XPS

$r = V^{5+}/V_{\text{total}}$	ratio of number V^{5+} ions to total number of V ions from XPS
$(1 - r) = V^{4+}/V_{\text{total}}$	ratio of number V^{4+} ions to total number of V ions from XPS
$f = \text{Fe}^{3+}/\text{Fe}_{\text{total}}$	ratio of number Fe^{3+} ions to total number of Fe ions from XPS
$(1 - f) = \text{Fe}^{2+}/\text{Fe}_{\text{total}}$	ratio of number Fe^{2+} ions to total number of Fe ions from XPS
N_V	molar fraction of vanadium atoms from ICP
N_{Fe}	molar fraction of iron atoms from ICP
N_P	molar fraction of phosphorus atoms from ICP
C_{calc}	calculated Curie constant (emu K/Oe g)

Assuming V only exists either as V^{5+} ($p_{\text{eff}} = 0$) or V^{4+} ($p_{\text{eff}} = 1.8 \mu_B$) and since Fe exists as either Fe^{3+} ($p_{\text{eff}} = 5.9 \mu_B$) or Fe^{2+} ($p_{\text{eff}} = 5.4 \mu_B$), the Curie constant C_{calc} is given by

$$C_{\text{calc}} = \left\{ fN_{\text{Fe}} \frac{p_{\text{eff}}^2(\text{Fe}^{3+})}{3k_B} + (1 - f)N_{\text{Fe}} \frac{p_{\text{eff}}^2(\text{Fe}^{2+})}{3k_B} + (1 - r)N_V \frac{p_{\text{eff}}^2(V^{4+})}{3k_B} \right\} \frac{N_A}{MW},$$

where N_A is Avogadro's number and MW is the molecular weight of the glass sample. The molecular weight is determined from the following expression:

$$\begin{aligned} MW = & fN_{\text{Fe}}MW_{\text{Fe}_2\text{O}_3} + (1 - f)N_{\text{Fe}}MW_{\text{Fe}_2\text{O}_2} \\ & + rN_VMW_{\text{V}_2\text{O}_5} + (1 - r)N_VMW_{\text{V}_2\text{O}_4} \\ & + N_PMW_{\text{P}_2\text{O}_5}. \end{aligned} \quad (\text{A.1})$$

References

- [1] G.F. Lynch, M. Sayer, J. Phys. C 6 (1973) 3661.
- [2] M. Sayer, G.F. Lynch, J. Phys. C 6 (1973) 3774.

- [3] A.K. Bandyopadhyay, J.O. Isard, J. Phys. D 10 (1977) L99.
- [4] L.D. Bogomolova, J. Non-Cryst. Solids 30 (1979) 379.
- [5] L.D. Bogomolova, M.P. Glassova, J. Non-Cryst. Solids 37 (1980) 423.
- [6] L.D. Bogomolova, V.A. Jachkin, M.P. Glassova, S.N. Spasibkina, J. Phys. 42 (1981) C4-993.
- [7] L.D. Bogomolova, M.P. Glassova, O.E. Dubatovko, S.I. Reiman, S.N. Spasibkina, J. Non-Cryst. Solids 58 (1983) 71.
- [8] O. Cozar, I. Ardelean, Physica 27 (1982) 41.
- [9] O. Cozar, I. Ardelean, Gh. Ilonca, Solid State Commun. 44 (1982) 809.
- [10] I. Ardelean, O. Cozar, Gh. Ilonca, J. Non-Cryst. Solids 68 (1984) 33.
- [11] G.D. Khattak, M.A. Salim, L.E. Wenger, A.H. Gilani, J. Non-Cryst. Solids 262 (2000) 66.
- [12] E. Cular, J. Non-Cryst. Solids 223 (1998) 47.
- [13] E.E. Khawaja, Z. Hussain, M.S. Jazzar, O.B. Dabbousi, J. Non-Cryst. Solids 93 (1987) 45.
- [14] G.E. Mullenberg (Ed.), Handbook of X-ray Photoelectron Spectroscopy, Perkin Elmer, Norwalk, CT, 1970.
- [15] C.R. Bamford, Color Generation and Control in Glass, Elsevier, Amsterdam, 1977.
- [16] A. Proctor, P.M.A. Sherwood, Anal. Chem. 52 (1980) 2315.
- [17] J.F. Moulder, W.F. Stickle, P.E. Sobol, K.D. Bomben, Handbook of X-ray Photoelectron Spectroscopy, Perkin Elmer, Eden Prairie, MN, 1992.
- [18] J. Wong, C.A. Angell, Glass Structure by Spectroscopy, Marcel Dekker, New York, 1976.
- [19] B.M.J. Smets, D.M. Krol, Phys. Chem. Glasses 25 (1984) 113.
- [20] R.K. Brow, C.M. Arens, X. Yu, E. Day, Phys. Chem. Glasses 35 (1994) 132.
- [21] C.D. Wagner, W.M. Riggs, L.E. Davis, J.F. Moulder, in: G.E. Mullenberg (Ed.), Handbook of X-ray Photoelectron Spectroscopy, Perkin Elmer, Eden Prairie, MN, 1979.
- [22] R. Bruckner, H.-U. Chun, H. Goretzki, M. Sammet, J. Non-Cryst. Solids 42 (1980) 49.
- [23] D. Briggs, J.C. Riviere, in: D. Briggs, M.P. Seah (Eds.), Practical Surface Analysis by Auger and X-ray Photoelectron Spectroscopy, Wiley, Chichester, 1988, p. 119.
- [24] C.S. Sunandana, K.S. Rao, Phys. Stat. Sol. A 90 (1958) 681.
- [25] I.G. Austine, E.J. Garbett, in: P.G. LeComber, J. Mart (Eds.), Electronic and Structural Properties of Amorphous Semiconductors, Academic Press, London, 1973.
- [26] K.P. Sinha, N. Kumar (Eds.), Interactions in Magnetically Ordered Solids, Oxford Press, Oxford, 1980.