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# Encapsulation of Polymers into $MoS_2$ and Metal to Insulator Transition in Metastable $MoS_2$

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A general approach for the encapsulation of a variety of saturated polymers between the layers of $MoS_2$ giving electrically conductive lamellar compounds is reported.

Recently, the area of composite organic/inorganic polymeric materials, mixed at the molecular level, has been receiving considerable research interest. The organic polymers can be saturated or conjugated, while the inorganic components can be three-dimensional (3-D) systems such as zeolites,[1] 2-D layered materials such as clays[2] and metal oxides,[3] and even 1-D materials such as $(Mo_3Se_3^-)_n$ chains.[4] In principle, such systems may exhibit a variety of unique properties arising from the combination of the organic and inorganic components.[5] For example, recently researchers from Toyota prepared new molecular-scale nanocomposites made from saturated polymers (Nylon-6 and other plastics) intercalated in clay layers.[6] These products show extraordinary mechanical strength far greater than that attainable or expected by simply mixing the two components. Along similar lines, we chose $MoS_2$ as the inorganic component for intercalation with a series of saturated polymers and we present here preliminary results on materials obtained from such studies.

Recently, $MoS_2$ was shown to disperse into single layers by reaction of $Li_xMoS_2$ with water.[7] Reprecipitation of the layers in the presence of small molecules results in intercalation compounds.[8] We have exploited this property and used single layer $MoS_2$ in water and polymer solutions to produce novel polymer/$MoS_2$ nanocomposites. $MoS_2$ is attractive for investigation because it is readily available (it occurs naturally) and is already an important material with several practical applications, such as hydrodesulfurization (HDS),[9] solid lubrication[10] and rechargeable batteries.[11] A previous report described the polymerization of styrene in the galleries of $MoS_2$ to yield relatively low molecular mass polystyrene.[12] The approach reported here involves the direct intercalation of polymers and is not limited by the ability to polymerize a monomer after it has been inserted in $MoS_2$. Even if such polymerization can be initiated, the molecular mass of the polymer will be low and in some cases by-products could also be generated which have to be dealt with. Thus, the methodology described here is general and a very large molecular mass can potentially be intercalated.

Treatment of $MoS_2$ with 3 equiv. of $Bu^nLi$ in hexanes under inert atmosphere yields $LiMoS_2$.[13] Reaction of $LiMoS_2$ with water accompanied by sonication, results in a suspension of single layers of $MoS_2$. Addition of a polymer solution to the suspension causes flocculation during which the $MoS_2$ layers sandwich the polymer chains in a remarkably well-ordered manner; eqns. (1), (2) and (3) represent the general synthesis of $(polymer)_xMoS_2$.

$$MoS_2 + BuLi \longrightarrow LiMoS_2 + \tfrac{1}{2}Bu\text{–}Bu \quad (1)$$

$$LiMoS_2 + xH_2O \longrightarrow (MoS_2)_{\text{single layers}} + xLiOH + {}_{(x/2)}H_2 \quad (2)$$

$$(MoS_2)_{\text{single layers}} + x\text{polymer} \xrightarrow{\text{flocculation}} (\text{polymer})_xMoS_2 \quad (3)$$

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**Table 1** Summary of results obtained from the encapsulation of various polymers in $MoS_2$

| Intercalate | Polymer ($M_r$) | *d*-Spacing/Å | Expansion of layers/Å | Solvent | Room temp. cond. (S cm$^{-1}$) | Thermal stability under $O_2$/°C | Thermal stability under $N_2$/°C |
|---|---|---|---|---|---|---|---|
| $(PEO)_{0.92}MoS_2$ | 100000 | 16.3 | 10.1 | $H_2O$ | 0.1 | 225 | 284 |
| $(PEO)_{1.0}MoS_2$ | 5000000 | 14.5 | 8.3 | $H_2O$ | 0.02 | 220 | 269 |
| $(PPG)_{0.5}MoS_2$ | 1000 | 15.4 | 9.2 | $H_2O$ | 0.2 | 200 | 250 |
| $(PVP)_{0.76}MoS_2$ | 10000 | 21.1 | 14.9 | $H_2O$ | 0.003 | 240 | 270 |
| $(MCel)_{0.26}MoS_2$ | 63000 | 20.4 | 14.2 | $H_2O$ | 0.0004 | 210 | 225 |
| $(PEI)_{0.83}MoS_2$ | 55000 | 10.2 | 4.0 | $H_2O$ | 0.005 | 248 | 253 |
| $(Nylon\text{-}6)_{3.6}MoS_2$ | 10000 | 17.5 | 11.3 | Hot $CF_3CH_2OH$ | $<10^{-6}$ | 277 | 320 |
| $[(\text{-}CH_2\text{-})_n]_{3.0}MoS_2$ | Ultra high | 10.3 | 4.1 | Hot decalin | $<10^{-6}$ | 274 | 425 |

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We intercalated poly(vinylpyrrolidinone) (PVP), poly(ethylene oxide) (PEO), poly(propylene glycol) (PPG), methyl cellulose (MCel) and poly(ethylenimine) (PEI) by adding solutions in $H_2O$, MeOH or MeOH–$H_2O$–mixture to an aqueous suspension of single layers of $MoS_2$.† Using the non-miscible solvent decalin we also intercalated polyethylene of ultra-high molecular mass into $MoS_2$. Furthermore, we introduced the nylon-6 into $MoS_2$ using $CF_3CH_2OH$ as a solvent for the polymer. All products were thoroughly washed with the appropriate solvent to remove any separate polymer phase.

X-Ray powder diffraction patterns show that all polymer intercalates have layered structures as suggested by the intense (00l) reflections and indicate well-defined mono-or bi-layers of polymers in the gallery space (Fig. 1). The *d*-spacings, interlayer expansions and other properties for all intercalates are tabulated in Table 1. The largest $MoS_2$ layer separations were obtained from PVP and MCel.

The charge transport properties of these materials were determined by four-probe electrical conductivity measurements of pressed pellets. The room temperature conductivity values are listed in Table 1. Variable temperature electrical conductivity measurements for $(PEO)_xMoS_2$ ($M_r$ of polymer: 100000) indicate a surprisingly high conductivity, as shown in Fig. 2(*a*). In the temperature range 50–300 K, the material exhibits weak, thermally activated behaviour. The corresponding thermoelectric power measurements, however, indicate that the material is a p-type metallic conductor, as observed by the very small and positive Seebeck coefficient. A marked feature in both the conductivity and the thermopower data is an abrupt, well-defined transition at *ca.* 14–15 K, see Fig. 2. Below 14 K, the conductivity decreases by six orders of magnitude while the thermopower suddenly discontinues its upward slope and drops to negative values. This type of behaviour is not typical for a normal metal. However, a similar anomalous decrease in conductivity at low temperatures has been reported for certain layered metallic dichalcogenide phases when doped with metal ions, *e.g.* 1T-$Ti_{0.085}Ta_{0.915}S_2$, 1T-$Nb_{0.15}Ta_{0.85}S_2$, 1T-$Hf_{0.085}Ta_{0.915}S_2$.[15] The metal to insulator transition in these systems has been attributed to a charge density wave (CDW); a coupled periodic distortion of the conduction electron density with respect to the crystal lattice.[16] Even though, the nature of the transition in our system is not well understood at the moment, we believe that a similar CDW effect might be operative at low temperature (<14 K), creating an electronic instability. The metallic character of the material is confirmed by the magnetic susceptibility data, which show a well-behaved temperature independent Pauli-like behaviour in the 50–300 K, temperature range.

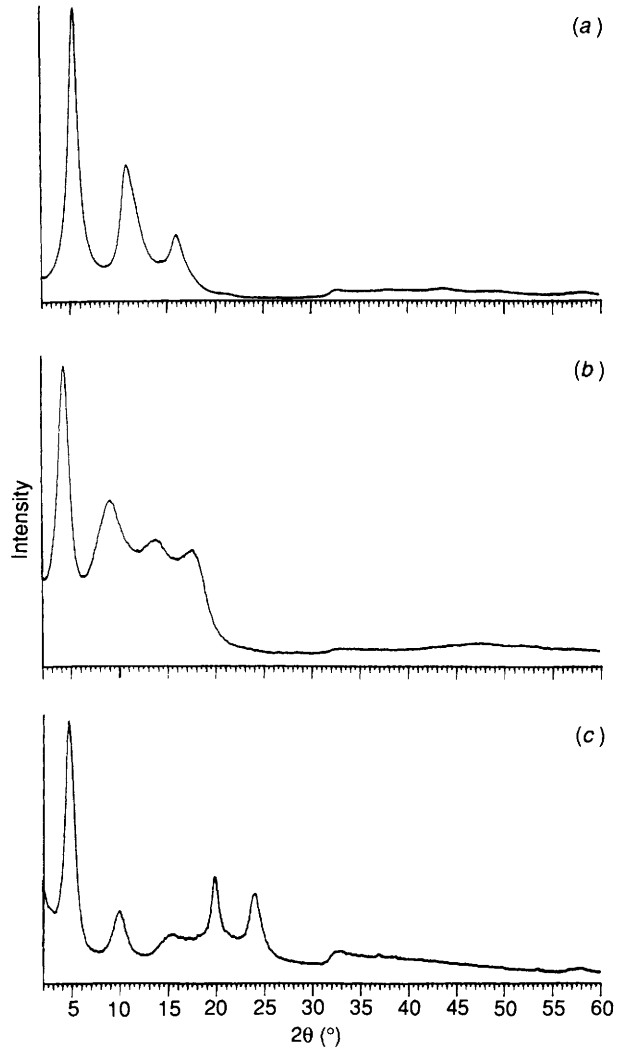


**Fig. 1** XRD patterns of (*a*) $(PEO)_{0.92}MoS_2$, $M_r$ of PEO = 100000, (*b*) $(MCel)_{0.26}MoS_2$ and (*c*) $(nylon\text{-}6)_{3.6}MoS_2$

The metallic character of $(PEO)_xMoS_2$ is explained by considering the structure of $MoS_2$ in this material. Pristine 2H-$MoS_2$ undergoes a structural transformation upon interca-

† Contrary to the report of earlier workers,[8] we found that this particular technique does not necessarily require the use of a water-immiscible liquid system. We have also successfully been able to intercalate neutral polyaniline dissolved in *N*-methylpyrrolidinone (a water-miscible solvent), when added to an aqueous suspension of single molecular $MoS_2$ layers.[14]

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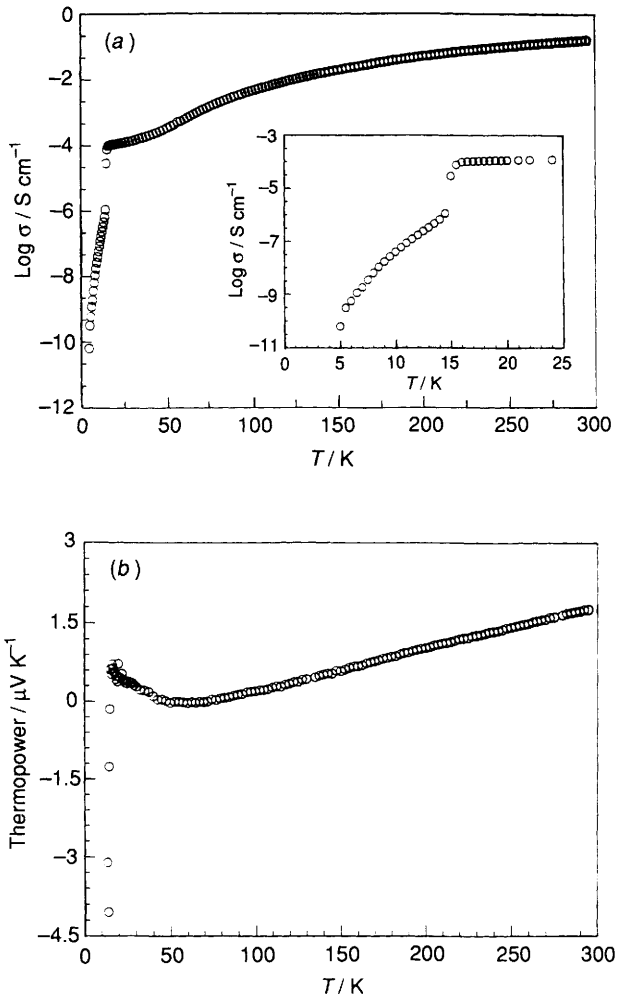


**Fig. 2** (*a*): Four-probe variable temperature electrical conductivity data for polycrystalline pellets of $(PEO)_{0.92}MoS_2$. The inset graph shows an expanded view of the low temperature data to highlight the metal to insulator transition. (*b*) Variable thermopower data of $(PEO)_{0.92}MoS_2$[$M_r$ of PEO = 100000]. The data were corrected for the contribution of the Au contacts employed during the measurements.

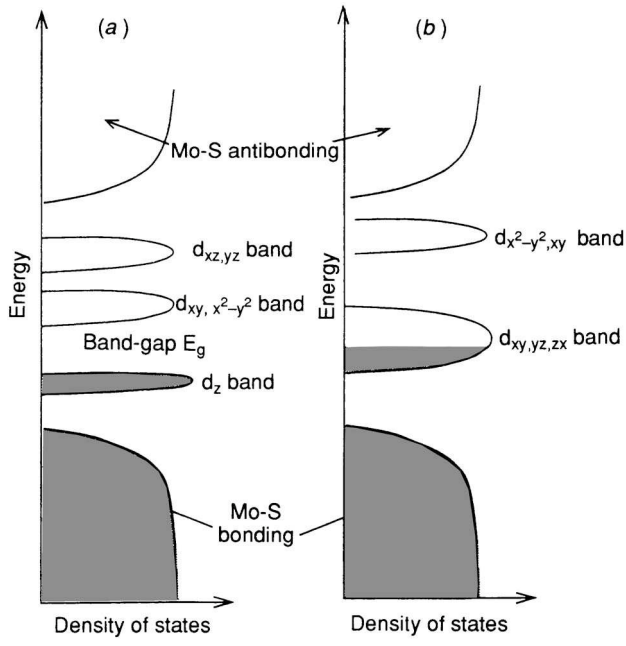


**Fig. 3** (*a*): Qualitative band diagram of the $D_{3h}$-$MoS_2$ slab. (*b*) Qualitative band diagram of the $O_h$-$MoS_2$. Shaded bands are filled. Adapted from refs. 18 and 19.

lation with lithium in which the coordination of the $Mo^{3+}$ atom becomes octahedral from trigonal prismatic.[17] The band diagram of 2H-$MoS_2$ indicates that it is a semiconductor.[18] In 2H-$MoS_2$ each layer will be referred to as $D_{3h}$-$MoS_2$. Upon reaction of $LiMoS_2$ with water, single $MoS_2$ layers form by rapid oxidation which leaves the $Mo^{4+}$ atom trapped in an octahedral coordination, thereby stabilizing a metastable structure for an $MoS_2$ layer.[17b] In this material each layer will be referred to as $O_h$-$MoS_2$. Fig. 3 shows qualitative band structure diagrams associated with the $D_{3h}$-$MoS_2$ and $O_h$-$MoS_2$ layers. The trigonal prismatic modification develops a band gap between the filled $d_{z^2}$ and empty $d_{x^2-y^2,xy}$ band. In the octahedral modification the degenerate $d_{xy,yz,xz}$ orbitals form a single band populated by two electrons producing a metallic system.[19] Recently, Wypych and Schöllhorn reported the preparation of the metastable 1T-$MoS_2$ which was proposed to contain $O_h$-$MoS_2$ layers.[20] This phase was shown to be metallic, by its Pauli paramagnetism, and thermopower measurements. Differential scanning calorimetry (DSC) studies on our restacked $O_h$-$MoS_2$ layers show that the layers transform to the more stable $D_{3h}$ form at *ca.* 100 °C, by observation of an exotherm at that particular temperature. This is in good agreement with Schöllhorn's 1T-$MoS_2$, which was reported to show an exothermic transition at *ca.* 95 °C. It appears that the material produced by restacking the dispersed $MoS_2$ layers and 1T-$MoS_2$ are one and the same product. Interestingly, the conversion temperature varies depending on the polymer involved and the associated d-interlayer spacing. The $(PVP)_{0.76}MoS_2$ shows the highest conversion temperature ($T_c$ *ca.* 177 °C) while plain restacked $MoS_2$ the lowest ($T_c$ *ca.* 94 °C).

Upon aging, metallic 1T-$MoS_2$ reverts to the more stable semiconductive 2H-phase.[17b] This is consistent with our observations that an aged $(PEO)_xMoS_2$ sample shows three orders of magnitude lower conductivity. However, the transformation from $O_h$ to $D_{3h}$ is very slow and seems to take months to complete at room temperature. On the other hand, an increase in temperature, greatly accelerated this transformation. This is reflected in the observed low conductivities of $(\text{nylon-6})_{3.6}MoS_2$ and $(\text{polyethylene})_{3.0}MoS_2$ which were synthesized by refluxing the respective reaction mixtures.

The high electrical conductivities of $(PEO)_xMoS_2$ and $(PPG)_xMoS_2$ with respective room temperature values of 0.1 and 0.2 S $cm^{-1}$, are among the highest reported among polymer/host nanocomposites.[3] These materials can offer two advantages as cathodes in solid state high energy density lithium batteries. First, the encapsulated polymers when complexed with Li would provide the desired solid electrolyte[21] and second, the high conductivity of the layers should not necessitate the addition of conductive additives (*e.g.* graphite, carbon black). The exfoliation/reprecipitation property of $MoS_2$ has been exploited to produce novel nanocomposites lamellar materials. This process appears to be general and should apply to a large variety of other soluble polymers including conductive polymers *via* soluble precursor routes such as poly(*p*-phenylene vinylene) and poly(*p*-phenylene).[22]

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