PS-11.02.13 X-RAY REFLECTIVITY STUDY OF POLYMERIC THIN FILMS. By Lichen Wang and Earle Ryba, Department of Materials Science and Engineering, The Pennsylvania State University, University Park, PA 16802

Recently, neutron and X-ray reflectivity have emerged as powerful tools for the investigation of the surface, interface behavior of polymeric materials. We have used the X-ray reflectivity technique to study various types of polymeric thin films. The density profile for a very thin polyethylene film on a gold substrate, determined through non-linear model fitting of the reflectivity data, will be presented. The density of the film is significantly higher near the substrate, where the substrate surface appears to have induced an enhanced crystallinity. Near the air/polyethylene interface, the density is slightly higher than that in the bulk of the film. The crystallinity near this interface appears to be greatly affected by the surface tension. It is interesting that the roughness near both of these interfaces derived from the model fitting of the reflectivity curve is smaller than what we expect. Support from the Eastman Kodak Company is gratefully acknowledged.


The ability to obtain structural information with sub-Angstrom resolution on Langmuir-Blodgett (LB) model membranes using long period x-ray standing waves (XSW) has been demonstrated previously (Bedzyk et al., Science, 1988, 241, 1788-1791; Phys. Rev. Lett., 1989, 62, 1376-1379). In the present study, we wished to determine 1) if the variable period XSW generated by an x-ray mirror during total external reflection could be used to locate precisely and accurately a heavy atom layer positioned several hundred Angstroms or higher above the mirror surface which would prevail in a lost of biologically relevant model systems, and 2) if the thermally induced phase transitions occurring in LB films are sensitive to the number and identity of the lamellae in the membrane stack. The sample series examined in addressing the first question consisted of an octadecylthiol coated gold mirror on top of which a variable number (0, 2, 8, 14) of di-tricosenoic acid (ωTA) bilayers followed by a single, upper bilayer of zinc arachidate was deposited by the LB technique. With 14 bilayers of ωTA, zinc-to-gold surface separations were close to 1,000 Å. The XSW measurements showed that the zinc Kα fluorescence yield profiles from the sample set are in excellent agreement with the calculated XSW electric field distribution (Wang et al., Nature, 1991, 354, 377-380). Further, the numerical fitting of the data reveals that Angstrom precision can be achieved in determining both zinc atom layer mean position and width above the gold mirror surface. The two samples chosen for the thermal stability study of LB films were identical to those described above and incorporated 0 and 2 bilayers of ωTA, respectively.

Variable period XSW measurements provided precise positional information on the zinc layer mean position and width and were used to track the collapse of the heavy atom layer during thermotropic phase transitions. The two samples showed quite disparate pretransitional rearrangements, transition temperatures and apparent cooperativity and final, high temperature zinc distribution. Further, the temperature-induced change observed was not reversed upon cooling to and subsequent storage at room temperature. The results of these experiments demonstrate clearly that the XSW field is well defined at close to a thousand Angstroms above the mirror surface and that the XSW method is well suited for determining the position of layered heavy atoms with a precision of Angstroms while the heavy atom layer is up to 1,000 Å or even higher above the mirror surface. Additionally, the quality of these data allows the enormous potential of XSW as structural probes in membranes and in thin film related phenomena.

PS-11.02.15 STRUCTURE OF MOLYBDENUM SULPHIDE THIN FILMS

N. Mattner*, W. Pischke*, S. Doyle**

*) Institut für Festkörper- und Werkstoffforschung Dresden e.V., Postfach, 01171 Dresden, FRG

**) Technische Hochschule Darmstadt, FB Materialwissenschaft

Thin films of MoS2 (150 nm - 600 nm in thickness) were produced by Hf-magnetron sputtering onto silicon wafers. The structure of the films has been investigated as a function of temperature using X-ray diffraction and transmission electron microscopy. Fig. 1 shows the obtained diffraction patterns as a function of temperature. The X-ray patterns indicate that there is a continuous development of the structure rather than transition from amorphous to crystalline state.

The crystal structure of hexagonal MoS2 (space group P63/mmc) is characterized by a stacking of Mo and S layers. The initial state of the films is characterized by a random stacking of S-Mo-S layers forming two-dimensional lattice in the a-b plane. The extension in c-direction of the stacks is around 2 nanometres. Thermal treatment leads to an increase in grain size to around 20 nm at 900°C.
11-Surfaces, Interfaces and Thin Films

Parallel to the crystal growth a reduction of the fraction of stacking faults is observed indicated by the decrease of the c-lattice parameter and by the appearance of (hk1) reflections. Differences in the initial film thickness influence the structural state of the untreated material but they disappear during heat treatment.

Fig. 1: Diffraction pattern of a 450 nm thick MoS2 film at T = 35 (lower), 100, 200, 300, 400, 500, 600, 700, 800, 100°C (upper)

In conclusion, we have found the PtCo alloy diffraction at PtCo multilayers. The appearance of PtCo alloy did not destroy the perpendicular magnetic anisotropy which is believed to be mainly induced by interface anisotropy, the influence of interface atoms on the anisotropy can be further studied.

PS-11.02.16 THE STRUCTURE AND MAGNETIC ANISOTROPY OF PtCo MULTILAYERS By Zhi-hui Jiang, Chang-lin Kuo, De-fang Shen, Rong-fa Guo, Tian-shen Shi

Recently, PtCo multilayer has stimulated the interests of many researchers due to its potential as a high-density magnetooptical storage material. Generally, such a candidate material must have a large Faraday or Kerr effect at short wavelength, a perpendicular anisotropy and a relatively large coercivity at room temperature. PtCo multilayers happen to satisfy all these conditions.

In our experiments, PtCo multilayers were prepared using dc magnetron sputtering on silicon substrates. The structures were determined by X-ray diffraction and the magnetic properties were measured by Kerr angle hysteresis loops. In table 1 are listed the sample parameters and their magnetic properties.

The low angle X-ray diffraction pattern, as shown in Fig. 1, clearly confirmed the existence of multilayer structure of sample 1. The bilayer thickness deduced from Bragg formula agreed well with the sample parameter. A simple simulation of high angle x-ray diffraction pattern was made. At any circumstances, the angle of 111 satellite peak was higher than the angle of (111) satellite peak, which is not in contradiction to the real pattern as shown in Fig. 2. We contribute this phenomena to the appearance of PtCo alloy at the interfaces, whose (111) peak overlaps with the Bragg peak of multilayers and (200) peak is at the same position of n=+1 satellite peak (C.-t. Lin and G.L. Gorman, Appl. Phys. Lett., 1972, 61(13), 1600). In sample 2, when Co layer was much thicker, the multilayer diffraction overwhelmed the PtCo alloy diffraction and the relative heights of these two peaks reversed.

Fig. 1: Low angle X-ray diffraction pattern of sample one

Fig. 2: High angle X-ray diffraction pattern of sample one

In the present letter we report the x-ray topographic examinations of InGaAs/GaAs superlattices. The specimens used in the experiments were grown by molecular beam epitaxy (MBE) on GaAs substrates. The minimal structure of the SLs samples were 150 periods of 70 Å In0.7Ga0.3As and 250 Å GaAs with about 1 µm GaAs buffer layer and a 3 µm capping layer, respectively. In order to eliminate the influence of the capping layer on the reflection topographs a sample (denoted A) was etched to remove its capping layer. The synchrotron radiation experiments on sample A were performed at 4W1A beam line of the Beijing Synchrotron Radiation Facilities (BSRF).

The actual structure of the specimens were determined by the simulations of experimentally measured rocking curves based on the dynamic diffraction theory for deformed crystals (Z. Hu, M. F. Cui and C. C. P. Phys. Rev. B, 1999, 41, 9303-9304). Percentage relaxations of the two component layers of SLs were particularly found from the simulations.

224 reflection topographs were taken at the zero order peak of the SLS and the substrate peak, respectively. They were characterized as orthogonal strain parallel to the [110] and [110] directions, respectively. X-ray topographs were also taken under incoherent transmission conditions using Cu Kα radiation (µ=2.17, where µ is the linear absorption constant and t is the sample thickness). In 220 or 220 anomalous transmission topographs shown in Fig. 1 the striations parallel to [110] or [110] disappeared, respectively. According to the invisibility criteria of dislocations, both the striations parallel to [110] or [110] were edge-type dislocations with Burger's vectors in [110] or [110] direction in the [110] growth plane.

It is interesting to see that in addition to the striations in the (001) growth plane other striations (see region C of Fig. 1.) which obeyed the same extinction law were observed on the cleavage planes of sample A. The stereoscopic topographs appear evidence that the multi-striations distribute over the SLS and...