Chapter 5

Stability of complex formation between a donor azo dye and an acceptor molecule at the air-water interface

5.1 Introduction

The phenomena of complex formation are seen in various systems in surface science. There are many ways a complex can be formed and their mechanisms and interactions are very different. In biological systems, there have been reports on the complex formation between cholesterol and di-palmitoyl phosphatidyl choline at the air-water interface [1, 2]. Here, the interactions between the polar heads are important which lead to the formation of a condensed complex system. A complex formation in the mixed monolayer of oppositely charged lipid groups (dioctadecyl dimethyl ammonium bromide with sodium hexadecyl sulfate), has been reported [3]. Here, the electrostatic interactions play a dominant role and led to a large reduction in the A/M in the mixed monolayer.

Mixed Langmuir monolayer studies are useful in stabilizing the monolayers with functional properties. For the stabilization one needs to understand the interactions between the monolayer components in the mixed film. There are a few reports in the literature on the mixed monolayer behavior of a dye with a liquid crystalline (LC) material. A mixed monolayer study on the non-amphiphilic dichroic azo dye with an amphiphilic LC molecule, 8CB has been reported [4]. Their studies indicated that the azo dye which possessed both donor and acceptor groups at the ends did not form a stable mixed monolayer. However, an azo dye
with only a donor group at one end and no acceptor group at the other end formed a stable miscible mixed monolayer up to 0.8 MF of 8CB.

A mixed Langmuir monolayer study on liquid crystal molecules in which one component with an electron donor group and the other with an electron acceptor group has been reported [5]. Here the two individual monolayers were unstable in time at constant surface pressure. The equilibrium spreading pressure for the individual components were found to be zero. On the other hand, the mixed monolayer formed a condensed phase exhibiting large reduction in the A/M and was found to be stable with time. Several studies have discussed the importance of dye doped liquid crystals. This chapter deals with the miscibility studies in the mixed monolayer of an azo dye, p-(ethoxy)p-phenylazo phenyl hexanoate(EPPH) with 8CB. There are many reports on azo dye in the literature on the more polar nature of the cis form (3.0 to 3.5 Db) compared with the trans form (0 to 0.5 Db) [6, 7, 8]. This change in the polarity affects the monolayer phases and its rheological properties [9]. Their studies show that, in the cis-form, the monolayer yields an isotropic, Newtonian film which cannot be oriented by surface flows. However, reverting the molecule back to trans form led to the formation of compact monolayer which responds to flow with an anisotropic and non-Newtonian behavior.

5.2 Experiment

The materials EPPH(Eastman Kodak) and 8CB(Aldrich) were obtained commercially. EPPH was recrystallised using ethanol as a solvent and 8CB was used as procured. The isomers of EPPH are shown in Figure 5.1. The structure of EPPH in trans state is shown in Figure 5.1(a). The molecule is rod like with core length of about 9Å. The structure of EPPH in cis state is shown in Figure 5.1(b). Here, the molecule is in bent form with core dimension of 5.5 Å. A change in geometry for the azo dye from a rod shape to bent shape can be induced by photo-illuminating the azo dye with UV or visible radiation. For the EPPH azo dye, the trans(rod shape) form can be converted to cis(bent shape) form by irradiating UV light of 360 nm wavelength. In the cis-form, the length of the molecule decreases but, the effective
area which it occupies increases. To revert back the molecule to trans form one can either use blue light (400 \text{nm}) for faster relaxation or can keep it in dark for slower relaxation. The set up for the study of the surface pressure (\(\pi\) - area per molecule (A/M)) isotherm has been described in detail in chapter-3. Stock solution of 1.5 mM concentration was used to prepare the mixtures. The barriers were compressed at the rate of 1.1 (Å\(^2\)/molecule)/min. The temperature was maintained at 28±0.7 ° C by circulating water. The relative humidity was about 80±5 %. The \(\pi\)-A/M experiments were usually carried out in dark for the mixed monolayer unless otherwise mentioned.

We have carried out epifluorescence microscopy on this system wherein a fluorescent dye, NBDHDA, of about 1% molar concentration in the monolayer was used. The details are given in chapter-3. Brewster angle microscopy (BAM) technique was used for the observation of the monolayer phases. Mini-BAM with a laser of wavelength 632 \text{nm} was incident on the water surface at Brewster angle (53°) and the reflected light from the interface was allowed to fall on the detector. The presence of the monolayer alters the Brewster angle.
condition leading to a change in reflectivity. This change in the reflectivity will give rise to a contrast in the BAM images. The pixel dimensions of the CCD detector was nearly square and was of 8.3 $\mu$m x 8.2 $\mu$m. The field of view was 6.4 x 4.8 mm$^2$ and the resolution was of the order of 200 $\mu$m. The monolayer imaged under BAM with a large field of view gives information on the homogeneity and phase coexistence.

5.2.1 Setup for Photo-induced conformational change

The experiments were conducted in a dark room to avoid stray light which can affect the dye. The schematic diagram of the setup used for observing the textures under BAM for the azo monolayer and their mixtures is shown in Figure 5.2. To study the effect of photo-

![Figure 5.2: A schematic diagram of the setup for photo-illumination and BAM microscopy observations of the monolayer. 1.) Nima 611M trough, 2.) Barriers, 3.) Water, 4.) Photo-active azo monolayer, 5.) Mercury arc lamp(100 W) 6.) Replaceable filters for UV(peaked at 360 nm) and visible excitations(above 420 nm) 7.) Aluminium reflectors and 8.) Brewster angle microscope(Mini-BAM).](image)

illumination on the mixed monolayer we used a standalone Mercury arc lamp of 100 W. Filter(Schott) of 1 mm thickness was used to allow the 360 nm radiation. To revert back to trans form filter which allow 440 nm radiation was used. The intensities at 360 nm and 440 nm were 0.5 mW/cm$^2$ and 0.5 mW/cm$^2$ respectively. The lamp was placed some 15 cm away
from the monolayer to reduce heating effects. Aluminium covers were used as reflectors.

5.3 Results

The surface pressure(\(\pi\))-area per molecule(\(A/M\)) isotherm for EPPH-8CB mixed monolayer is shown in Figure 5.3. The individual EPPH monolayer exhibited a plateau region below an \(A/M\) of 30 \(\text{Å}^2\) where the surface pressure was less than 0.1 mN/m. Upon compression, the surface pressure increased steeply. The monolayer collapsed at a surface pressure of 15.2 mN/m with a limiting area(\(A_0\)) of 26 \(\text{Å}^2\). The isotherm for 8CB was in agreement with earlier reports [10, 11]. For the mixed monolayers, there was a drastic reduction in the \(A_0\) value. This was maximum at 0.5 MF of EPPH in 8CB. For all the compositions of EPPH, the isotherms exhibited only one collapse pressure indicating a good miscibility.

![Figure 5.3: Surface pressure(\(\pi\)) - area per molecule(\(A/M\)) isotherm for EPPH-8CB mixed monolayer at various mole fractions(MF) of EPPH in 8CB at \(t = 28^\circ C\).](image)

93
Another interesting result was the nature of the collapse in the isotherms. For the case of EPPH monolayer, after the collapse, the surface pressure drastically falls down. For the case of 8CB monolayer, the collapse was followed by a plateau region which is referred as “plateau” type collapse. This behavior is referred as “spike” type collapse. For the mixed monolayer, “plateau” type collapse behavior was found up to 0.7 MF of EPPH in 8CB. The “spike” behavior of collapse was seen at higher concentrations of EPPH. The collapse pressure is an important parameter in judging the miscibility and the stability for the mixed Langmuir monolayer. Figure 5.4 shows the variation of collapse pressure for EPPH-8CB mixed monolayer with increasing MF of EPPH in 8CB. It can be seen that the collapse pressure value increases with MF of EPPH and reaches a maximum at 0.7 MF and decreases at higher MF. Interestingly, we find that the magnitudes of the collapse pressure for the mixed monolayer were higher than that of the individual monolayers.

![Figure 5.4: Variation of the collapse pressure with increasing mole fraction(MF) of EPPH in 8CB.](image)

We have carried out Brewster angle microscopy(BAM) studies on the mixed monolayer
of EPPH and 8CB for various compositions under dark conditions. The BAM images obtained for EPPH monolayer are shown in Figure 5.5. At large A/M, corresponding to the zero surface pressure region, the monolayer appeared stiff with irregularly shaped islands separated by voids. The islands exhibited grainy textures(Figure 5.5(a)). The voids decreased progressively upon compression(Figure 5.5(b)). They disappeared completely at an A/M of 23.5 Å² corresponding to the steep region of the isotherm (Figure 5.5(c)). On further compression, the monolayer collapsed to 3D crystals(Figure 5.5(d) and 5.5(e)). The BAM images for the mixed monolayer at 0.7 MF of EPPH in 8CB are shown in Figure 5.6. The textures differs drastically from that of the EPPH monolayer alone. Here, the textures were homogeneous with less defects and voids(Figure 5.6(a) and 5.6(b)). At lower A/M, the im-

Figure 5.5: Brewster angle microscopy(BAM) images for EPPH at different A/M. Figure (a) and (b) show the presence of irregularly shaped islands with voids between them. These islands are rough and possess grainy texture. Figure(c) shows the disappearance of voids. Figure(d) shows the occurrence of bright spots indicating the onset of nucleation. Figure(e) shows the presence of 3D crystals in the collapsed state. Scale of each image is 6.4 x 4.8 mm².
Figure 5.6: BAM images for 0.7 MF of EPPH in 8CB. Figures (a) and (b) show the presence of homogeneously covered monolayer. Figures (c) and (d) show the collapsed state of the mixed monolayer. Here the bright spots indicate 3D crystals which coexists with multilayers in the background. Scale of each image is 6.4 x 4.8 mm$^2$.

Images show the nucleation of small crystallites which appeared as bright spots (Figures 5.6(c) and 5.6(d)). The BAM images taken for the mixed monolayer of 0.5 MF of EPPH in 8CB are shown in Figure 5.7. The images were similar to that of 0.7 MF of EPPH. At very low A/M, an uniform and homogeneously covered film was observed(Figure 5.7(a)). At lower A/M, they exhibited similar textures (Figure 5.7(b) and 5.7(c)). In the collapsed state, a few bright dots were observed in the mixed monolayer at 0.7 MF of EPPH in 8CB. The bright dots were much less. We found that with the increasing presence of 8CB uniform and homogeneous texture was formed. The BAM images for the mixed monolayer at 0.3 MF of EPPH
Figure 5.7: BAM images for 0.5 MF of EPPH in 8CB. Figures(a) to (c) exhibit a homogeneous and uniformly covered monolayer. Figure(d) shows the collapsed state of the mixed monolayer. Here the texture is predominantly uniform with a few bright spots (3D crystals). Scale of each image is 6.4 x 4.8 mm².

in 8CB are shown in Figure 5.8. In this case also, the mixed monolayer appeared uniform and homogeneous (Figure 5.8(a)). In the collapse stated, appearance of very small bright dots were seen (Figure 5.8(b)). These brighter domains are multilayers and were found to grow more (Figures 5.8(c) and 5.8(d)). The varying intensity of the multilayer domains indicate varying thickness. The BAM images for pure 8CB are shown in Figure 5.9. The coexisting G (dark region) + L₁ (bright region) phase is shown in Figure 5.9(a). With compression, L₁ phase appeared and the G domains diminished in size (Figure 5.9(b)). Above the collapse, the appearance of three layer (D₁) domains was observed as small grains (Figure 5.9(c)).
Figure 5.8: BAM images for 0.3 MF of EPPH in 8CB. Figure(a) shows the predominantly present condensed monolayer which appeared uniform and homogeneous. Figure(b) shows the collapsed state indicating the onset of small bright dots. Figures (c) and (d) show domains of varying intensities (multilayers). Scale of each images is 6.4 x 4.8 mm$^2$.

With compression, the fusion of $D_1$ domains are seen (Figures 5.9(d) to 5.9(f)). At lower $A/M$, a nearly uniform coverage of the $D_1$ phase was seen (Figure 5.9(g)). At still lower $A/M$, the images show the presence of multilayers ($D_2$) which were much brighter than the $D_1$ domains. These observations are in accordance with the earlier BAM and epifluorescence microscopic techniques on 8CB monolayer [10, 11, 12].

We have calculated thermodynamic excess functions for the mixed monolayer of EPPH and 8CB. For an ideal case of either complete miscibility or complete immiscibility, the law
Figure 5.9: BAM images for 8CB. Figure(a) shows the coexisting $G + L_1$ phase. Figure(b) shows the homogeneously covered $L_1$ phase. In Figure(c), the nucleation of the three layer $D_1$ domains from the $L_1$ phase is seen. Figures (d) to (f) show progressive fusion of the $D_1$ domains leading to an uniform coverage(Figure(g)). Figure(h) shows the occurrence of many bright spots which represent the multilayer $D_2$ domains coexisting with $D_1$ domains. Scale of each image is 6.4 mm x 4.8 mm$^2$.

The rule of additivity of areas is given by,

$$A_{id} = A_{EPPH}X_{EPPH} + A_{8CB}X_{8CB}$$  \hspace{1cm} (5.1)$$

where, $A_{id}$ is the ideal A/M, $A_{EPPH}$ and $A_{8CB}$ are the A/M for the pure monolayer components, $X_{EPPH}$ and $X_{8CB}$ are the mole fractions of EPPH and 8CB respectively. The $A_{id}$
calculated using the values obtained for individual monolayer was compared with the experimentally determined A/M($A_{12}$) for the mixed monolayer. A deviation from the ideal behavior indicates the presence of interactions between the monolayer components in the mixture. The variation of $A_{id}$ and $A_{12}$ with MF of EPPH in 8CB at different surface pressures are shown in Figure 5.10. A marked deviation in $A_{12}$ from $A_{id}$ was seen for the mixed monolayer. The excess or deficit of area per molecule ($A_{exc}$) for the mixture from that of the ideal case will indicate the type of interactions. This is given by,

$$A_{exc} = A_{12} - A_{id}$$  \hspace{1cm} (5.2)

The $A_{exc}$ being positive indicates that the interactions are repulsive and the $A_{exc}$ being negative indicates that the interactions are attractive. The variation of $A_{exc}$ with increasing MF of EPPH in 8CB at different surface pressures are shown in Figure 5.11. We find that the $A_{exc}$ is negative throughout the EPPH composition. The maximum condensation was found
to occur(-13 Å²) at 0.5 MF of EPPH in 8CB. This indicated that the molecules in the mixed monolayer are compactly packed. The degree of stability for the mixed monolayer was ana-

Figure 5.11: Plot of the excess A/M, A_{exc} with mole fraction(MF) of EPPH in 8CB. The maximum condensation is seen at 0.5 MF of EPPH in 8CB.

lyzed by calculating the excess Gibbs free energy. The excess Gibbs free energy, ΔG_{exc}, for the mixed monolayers at constant surface pressures was obtained by integrating the excess area, A_{exc}, over surface pressure [13]. This is given by,

$$\Delta G_{exc} = N_a \int_{\pi^*}^{\pi} A_{exc} d\pi$$

(5.3)

Here, \(\pi^*\) is the surface pressure at which the two components of the mixed monolayer behave ideally(usually taken as zero) and \(N_a\) is the Avogadro number. The variation of the excess Gibbs free energy(\(\Delta G_{exc}\)) with increasing MF of EPPH in 8CB at different surface pressures is shown in Figure 5.12. The \(\Delta G_{exc}\) is found to be negative throughout the composition and is minimum at 0.5 MF of EPPH in 8CB. This indicated that the mixed monolayer was stable at all compositions and the maximum degree of stability occurs at 0.5 MF for \(\Delta G_{exc}\) value of about -275 J/mol.
Figure 5.12: Variation of the excess Gibbs free energy, $\Delta G_{exc}$ with increasing mole fraction (MF) of EPPH in 8CB at different surface pressures. At 0.5 MF of EPPH in 8CB, a pronounced minima is seen for different surface pressures.

The static compressional elastic modulus ($|E|$) is a measure of the monolayer resistance towards change in area. In general, it provides information on the rigidity of the mixed monolayer system. According to Davies and Rideal [14], the monolayer phases can be classified based on the magnitude of $|E|$ (Table 5.1). It is calculated from the $\pi - A/M$ isotherm using,

$$|E| = (A/M)(d\pi/d(A/M))$$  \hspace{1cm} (5.4)

Higher value of $|E|$ implies that the system possess low in-plane elasticity. The variation

<p>| Table 5.1: Magnitude of compressional elastic modulus, $|E|$, in different monolayer phases |
|-----------------------------------------------|</p>
<table>
<thead>
<tr>
<th>Phase</th>
<th>Clean Surface</th>
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<td>E</td>
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of compressional elastic modulus, $|E|$ with surface pressure, $\pi$ is shown in Figure 5.13 for EPPH-8CB mixed monolayer. Here, we find that for the mixed monolayer the value of $|E|$ above 0.5 MF of EPPH in 8CB is higher when compared with that of individual monolayer. The individual monolayers of EPPH and 8CB yielded $|E|$ values of 35 mN/m and 100 mN/m at a surface pressure of 5 mN/m respectively. The $|E|$ exhibited a maximum value of about 250 mN/m at 0.7 MF of EPPH and at a surface pressure of 12 mN/m. This indicated a higher rigidity for the mixed monolayer.

The temporal behavior of the monolayers were analyzed by monitoring the kinetics through area relaxation and surface pressure relaxation in a dark environment. In the area relaxation method, the surface pressure is fixed. To maintain the surface pressure, the barriers need to move correspondingly to adjust the area. Monitoring the variation of area with time provides information on the material loss at the A-W interface. The evolution of area

Figure 5.13: Variation of compressional elastic modulus, $|E|$, with surface pressure at different mole fraction(MF) of EPPH in 8CB.
with time for different EPHH compositions is shown in Figure 5.14. For the case of EPHH monolayer at a surface pressure of 5 mN/m, the $A_n$ decreases gradually. The normalized area($A_n$) is defined at a given surface pressure($\pi_0$) as the ratio of the $A/M$, $A$, at a given time to the initial $A/M$, $A_0$. At higher surface pressures(10 mN/m) decrease in $A_n$ was rapid for EPHH monolayer. However, for the case of 0.5 MF of EPHH and 0.7 MF of EPHH in 8CB, the decrease in $A_n$ were negligible (at $\pi_0$ values of 5 mN/m and 10 mN/m).

Figure 5.14: Variation of the normalized area, $A_n$ with increasing time at a given surface pressure $\pi_0$ at different mole fractions(MF) of EPHH in 8CB.

The kinetics of the surface pressure variation with time at given area per molecule was studied for the monolayers. In this method, we have used a compression rate of 2.8 (Å$^2$/molecule)/min. to reach the desired surface pressure. This method did not perturb the monolayer since there was no barrier movement. The analysis of the variation of $\pi$ with
time provides information on the kinetics of reorganisation of the molecules in the monolayer and on the kinetics of collapse of the monolayer [3]. The $\pi_n$ is defined as the ratio of the surface pressure($\pi$) which varies with time from the initial surface pressure($\pi_0$) attained at a given area per molecule. The change in normalized surface pressure, $\pi_n$, with time for different compositions are shown in Figure 5.15. For EPPH monolayer, the surface pressure falls gradually from the initial $\pi$ of 5 mN/m. For an initial $\pi$ of 10 mN/m, the fall in $\pi$ was rapid. Also the plot showed a bump around 7.5 minutes. In general, there is a fall in the surface pressure as a function of time. However, this is less for the case of 0.5 MF of EPPH in 8CB.

![Figure 5.15](image_url)

Figure 5.15: Variation of the normalized surface pressure, $\pi_n$, with increasing time and at different mole fractions(MF) of EPPH in 8CB.

We have probed the photo stability of the monolayers by alternately illuminating them
at given area per molecule with UV and visible radiation. When illuminated with UV radiation, the EPPH molecules, which have a photo-active azo moiety changes from trans to cis isomer in the mixed monolayer. On illuminating with visible light, it reverts back to trans isomer. These transformations from cis to trans and trans to cis leads to change in the surface pressure. The variation in surface pressure for EPPH monolayer on alternately illuminating with UV and visible radiation with time is shown in Figure 5.16. The monolayer when illuminated with UV radiation leads to a sudden jump in the surface pressure from a value of about 4.5 mN/m to 11 mN/m. However, when illuminated with visible radiation instead of UV radiation the surface pressure drops to about 2 mN/m. Repeated cycles of alternately UV and visible illumination yields a pattern as shown in Figure 5.16. We find, in general, for repeated cycles a decrease in the peak surface pressure for UV illumination. Correspondingly the dip(decrease from the initial surface pressure during repeated cycles of illumination) arising due to visible illumination also decreases with repeated cycles.

Figure 5.16: Variation in the surface pressure($\pi$) on alternately illuminating with UV and visible radiation for EPPH monolayer. Here the initial surface pressure was 4.5 mN/m.
We have carried out photo-stability tests on the mixed monolayer. The variation of the surface pressure with time on alternately illuminating with UV and visible radiation is shown in Figure 5.17. For 0.7 MF of EPPH, the variation in surface pressure with time upon alternately illuminating with UV and visible radiation showed a similar trend as that of the pure EPPH. This shows that even in the mixture, the transformation from trans to cis occurred indicating a weaker association between the components forming the complex. This trend is shown in Figure 5.17(a).
Figure 5.17: Variation of the surface pressure($\pi$) in the mixed monolayer at different mole fraction(MF) of EPPH in 8CB on alternate UV and visible illumination.
Similar trend was seen for 0.5 MF of EPPH but with less variation in peak surface pressure and dip (decrease from the initial surface pressure during repeated cycles of illumination) values on alternately illuminating with UV and visible radiation. This is shown in Figure 5.17(b). This showed that the dissociation of complex occurs even in the mixed monolayer. To confirm the weak complex formation we have undertaken a direct BAM imaging during the alternate UV and visible irradiation on the pure and mixed monolayer for EPPH-8CB system. Figure 5.18 shows the texture of the EPPH monolayer changing with time upon illumination. The initial grainy islands transformed to bright dots when it was UV illuminated (Figure 5.18(a)). Less bright dots were observed upon visible illumination (Figure 5.18(b)). In general, upon UV illumination, more bright dots were seen to nucleate (Figure 5.18(c)). The repeated runs led to increase in the growth of crystallites even in visible radiation. The in-situ BAM imaging during alternate UV-visible illumination for 0.5 MF of EPPH in 8CB at a surface pressure of 5 mN/m is shown in Figure 5.19. At this surface pressure, reversible demixing and mixing occurred during alternate UV and visible illumination. We have undertaken in-situ BAM imaging during repeated UV and visible illumination. This is carried out for 0.5 MF of EPPH in 8CB at a higher surface pressure of 14 mN/m (Figure 5.20). At this surface pressure phase separation of 8CB rich phase and

Figure 5.18: Brewster angle microscope images for EPPH monolayer during alternate illumination of ultra-violet (UV) and visible (V) radiation. Figure (a) shows the presence of bright dots appearing with the grainy texture in the background when illuminated using UV radiation. Figure (b) shows the presence of grainy islands with decrease in the number of bright dots when illuminated using visible radiation. Figure (c) shows the further increase of bright dots when illuminated using UV radiation. Scale of each image is 6.4 x 4.8 mm².
Figure 5.19: Brewster angle microscopy images for 0.5 MF of EPPH in 8CB at 5 mN/m with alternate UV and visible(V) irradiation. Figure(a) shows the initial homogeneous and uniform texture. Figure(b) shows the appearance of voids under UV illumination. Figures (c) to (e) show the disappearance of voids on illuminating with visible light. Figures (f) and (g) show the textural change under the second cycle of UV illumination. Here a phase separation to 8CB rich phase and EPPH rich phase was observed. Figures (h) to (j) show the evolution from the immiscible phase to a homogeneous and uniform phase indicating a miscibility under visible illumination. Figures (k) and (i) show the phase separation occurring during UV illumination in the third cycle. Figures (m) to (o) show the evolution from the phase separated to miscible monolayer during visible illumination. Scale of each image is 6.4 x 4.8 mm².
EPPH rich phase was observed. This separated phases were not miscible fully during visible irradiation even for longer durations.

Figure 5.20: BAM images of 0.5 mole fraction(MF) of EPPH in 8CB at 14 mN/m with alternate UV and visible(V) radiation. The initial texture was uniform and homogeneous. Figures (a) to (c) show the appearance of grainy texture during UV illumination. Figures (d) to (h) show development of streaks and less bright domains during visible radiation in the first cycle. Figures (i) to (k) show the bright fluidic domains appearing from the less bright texture during UV illumination. Figures (l) to (o) show the transformation of bright texture to less bright texture during visible radiation in the second cycle. The pattern did not change back to uniform and homogeneous film even after leaving overnight. Scale of each image is 6.4 x 4.8 mm$^2$. 
5.4 Discussions

The experiments on EPPH monolayer and EPPH - 8CB mixed monolayer were carried out in dark environment. The studies on the behavior of the pure and mixed monolayer were carried out using surface manometry, collapse pressure behavior, BAM imaging, thermodynamic excess function calculations and temporal stability. The $\pi$-A/M isotherm for the mixed monolayer exhibited a single collapse pressure for all EPPH compositions. This indicated a good miscibility between the EPPH and 8CB molecules. The collapse pressures for the mixed monolayer were higher in the range of 0.3 to 0.7 MF of EPPH in 8CB as compared to the collapse pressures for the individual monolayer (4.7 mN/m for 8CB and 15.1 mN/m for EPPH). The maximum collapse pressure occurred at 0.7 MF of EPPH in 8CB and was 25 mN/m. This indicated a much better stability of the mixed monolayer. The EPPH monolayer exhibited a spike after the collapse where the surface pressure decreased rapidly. The 8CB monolayer exhibited a plateau region after the collapse where the surface pressure was constant. For the mixed monolayer, the nature of collapse projects a transition from the spike to a plateau. Upto 0.7 MF of EPPH, the $\pi$-A/M isotherm exhibited a plateau. The spike nature of collapse occurred for higher concentrations of EPPH. Such collapse behavior has been reported in systems with dissociated polar heads [15]. Recent studies in cholesterol and phospholipid system indicated that the mechanism of collapse was similar to that of liquid crystalline compounds at the air-water interface [16]. Though the interactions are quite different, we found a transition from a “spike” to “plateau” in our system. There was a drastic reduction in the limiting area per molecule ($A_0$) value for the mixed film and the reduction was maximum at 0.5 MF. This indicated a highly condensed mixed monolayer.

The Brewster angle microscopy imaging on pure EPPH indicated large irregularly shaped islands with grainy textures separated by voids in between them (Figures 5.5(a) and 5.5(b)). We observed the nucleation and subsequent growth of crystallites even before the collapse for EPPH monolayer (Figure 5.5(d)). The occurrence of bright 3D crystals indicated the collapse (Figure 5.5(e)). The presence of 8CB in EPPH stabilized the mixed monolayer. The
Brewster angle microscopy images at a composition of 0.3, 0.5 and 0.7 MF of EPPH in 8CB indicated an uniform bright and homogeneous textures till collapse. In all these cases, there were less occurrences of bright 3D crystals after the collapse. This indicated that the presence of 8CB inhibits the crystal formation in EPPH tending to form multilayers.

Our thermodynamic excess function analysis indicated a deviation from the ideality in the $A/M$ for the mixed monolayer. This showed the presence of interactions in the mixed monolayer. The excess $A/M$ calculated was negative indicating an attractive interactions. The maximum degree of condensation was at 0.5 MF of EPPH. The reduction in the $A/M$ of about $-13 \text{ Å}^2$ was seen at 0.5 MF which is an indication for higher degree of condensation. The components in the mixed monolayer tend to form a compact monolayer which was more stable at air-water interface. The excess Gibbs free energy shows a value of about $-275 \text{ J/mol}$ at 0.5 MF of EPPH in 8CB indicating a stable complex formation between the components in the mixed monolayer [7].

The analysis of the $|E|$ reveals the rigidity for the mixed monolayer. From the magnitude of $|E|$, we infer that the mixed monolayer is a condensed monolayer. Based on these analysis and our BAM studies we conclude that this is likely to be a variant of $L_2$ phase. A weak charge transfer interactions between the donor EPPH and acceptor 8CB molecules might have played a role in the formation of the complex at the A-W interface. Due to the quenching of the fluorescent dye epifluorescence studies on these systems was not possible. The quenching might be due to the complex formation of EPPH with fluorescent dye which is an electron acceptor.

The temporal stability for the EPPH monolayer and mixed monolayer were investigated under dark conditions. Drastic reduction of normalized area($A_n$) with time for EPPH was observed when compared to that of mixed monolayer. This indicated that the EPPH molecules tend to nucleate and form crystals with time even at low surface pressure. This is also supported by our BAM imaging for EPPH monolayer(Figure 5.5). The presence of 8CB with EPPH stabilized the mixed monolayer and there was negligible reduction in the $A_n$ with time (Figure 5.14). The analysis on the variation of $A_n$ with time indicated the stability and
low nucleation rate for the mixed film when compared with EPPH alone. The analysis on the variation of $\pi_n$ with time for the EPPH monolayer and mixed monolayer indicated that there was some reorganisation of the molecules in the monolayer of EPPH which resulted in the appearance of bump. Stable monolayer are the ones whose dissolution rates in the subphase are very small when compared with the compression speed. The monolayer tends to reorganize when the compression is stopped towards a state where the free energy is minimum. This reorganization should be faster than the dissolution rates. The magnitude of reorganisation depends on compression and increases with increase in compression rate.

Our experiments on monitoring the surface pressure variation during alternate illumination of UV and visible radiation indicated a decreasing trend in the peak and dip values in the surface pressure for EPPH. This was due to the formation of 3D crystals in EPPH as indicated by our in-situ BAM imaging (Figure 5.18). However, in the mixtures at 0.5 MF and 0.7 MF of EPPH in 8CB, the fall in the peak value and the dip value of surface pressures upon alternate UV and visible illumination was less when compared with EPPH monolayer(Figure 5.17). The peak and the dip in surface pressure values from the initial surface pressure were similar for EPPH, 0.5 MF of EPPH and 0.7 MF of EPPH mixtures. This shows that the complex formation was not very strong and can be dissociated by UV illumination where the azo dye transformed from the trans state to cis state.

We have also carried out Brewster angle microscopy imaging to probe the stability of the complex by photo illumination for EPPH-8CB mixed monolayer system. Though the mixed monolayer was condensed as observed in BAM(under dark conditions), the photo-illumination caused dissociation of complex. This indicated a weaker complex formation between the EPPH and 8CB molecules. Our experiments on 0.5 MF of EPPH in 8CB indicated that at low surface pressure(5 mN/m) the mixed monolayer phase separated during UV illumination and was miscible during visible radiation(Figure 5.19). The experiments carried out at higher surface pressure(14 mN/m) for 0.5 MF of EPPH in 8CB indicated that the system phase separated permanently(Figure 5.20).

Raduge et. al. have reported [17] experiments and simulations on an azo system which
provides vital information on the macroscopic change in wetting at the molecular level. Their experiments clearly demonstrate that the photo-transformation of the azo dye into cis and trans isomers causes change in the orientation of the static dipoles with respect to the surface normal, thus leading to the surface energy difference.

In bulk systems for liquid crystalline mixtures, the stability of the mesophase can be enhanced or a new mesophase can be induced. Acceptor-donor molecules form a class of charge-transfer complexes which results in the induced mesophase. Here, the presence of specific polar ends whose tendency is to withdraw electrons or donate electrons towards the rigid core make it either electron deficient or electron rich [18, 19]. Other phenomena such as enhanced birefringence, polar diffraction gratings [20, 21], dye induced orientation of the liquid crystalline molecules [22], optical recording [23] and optical switching [24] are some of the novel applications of these photo-active systems. Photo-induced wetting and their controllability by proper design of the photo-active compounds yield better stability and durability. These developments are promising for patterned growth and photo-lithography applications [17], [25].

We have carried out surface manometry, kinetic studies on surface pressure and area, compressional elasticity modulus, Brewster angle microscopy and thermodynamic studies on the EPPH - 8CB mixed monolayer. Based on these studies, we conclude that the mixed monolayer is stable when compared to individual molecules. The presence of 8CB in EPPH inhibits the formation of crystallites in EPPH and tend to form multilayers above the collapse. The condensed area and the increase in collapse pressure for the mixed monolayer reveals the formation of the complex. The stability of the complex was checked with photo-illumination. The azo moiety of EPPH transforms from trans to cis upon illumination in EPPH-8CB mixed system. On illumination with visible light, the reverse transformation from cis to trans occurred in the mixed system. Based on BAM studies for mixed system, at low surface pressure, reversible demixing and mixing occurred during alternate UV and visible illumination. However, phase separation was permanently seen at higher surface pressures.
Bibliography


