Tunable Command Layers for Liquid Crystal Alignment


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Abstract: A simple method for the construction of a stable, tunable, self-assembled command layer for liquid crystal display purposes is described. A pyridine-functionalized oligosiloxane spontaneously forms an anisotropic, grooved surface on indium–tin-oxide, enabling it to align liquid crystalline molecules. The pyridine functions act as seeds for the epitaxial growth of stacks of highly ordered zinc phthalocyanines, the height of which can be controlled. These stacks increase the interaction between the surface and the liquid crystal matrix by amplifying the surface ordering into the liquid crystal bulk. By varying the height of the stacks, direct control over the properties of the liquid crystal domains is achieved. These properties can be further tuned by adding to the liquid crystal, micro- and nanomolar concentrations of nitrogen-containing compounds, which are capable of interacting with and dissolving the stacks. The procedures we describe offer possibilities to use such tunable systems in LCD-based sensor devices as well as in solar-cell applications.

1. Introduction

Uniform alignment of liquid crystal domains is usually achieved by rubbing a spin-coated polymer with a velvet cloth, a process which has been applied by industry for many years.1–3 This rubbing can create microscopic grooves and/or an anisotropic distribution of molecular bonds, in this way forming an anisotropic surface on which the nematogenic molecules align themselves, thereby minimizing the surface energy.3,4 In the past decade, several new methods for liquid crystal alignment have been introduced,3,5–7 e.g. photoinduced alignment5,6 and magnetic field alignment.7 Despite this, the attainment of direct and rational control over the molecular makeup of the alignment layer and its interaction with liquid crystalline molecules still poses a challenge.

We recently described a procedure for the preparation of stable, self-assembled alignment layers,8 which makes use of parallel nanometer-sized grooves that are present on commonly used indium–tin-oxide (ITO) surfaces. The size of these so-called “nanogrooves” is amplified over 3 orders of magnitude by the anisotropic grafting of monomer 1 (Chart 1), which spontaneously forms rigid oligomers in situ that self-assemble during layer formation to give aligned grooves of surface-grafted siloxane oligomers (Figure 1). When used in a twisted-nematic liquid crystal display (LCD), these self-assembled surfaces rivaled industrially manufactured displays in terms of interaction energy between the alignment layer and the liquid crystalline matrix.

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Chart 1

Figure 1. (A) AFM micrograph of a spontaneously formed alignment layer from I-Bar is 1 μm. (B) Schematic representation of the molecular order in the alignment layer shown in (A); the aligned siloxane oligomers are indicated in yellow.
In this paper, we report on a modification of this procedure by including the pyridine-functionalized siloxane 2 (Chart 1), which results in the formation of an alignment layer with micrometer-sized grooves that is capable of binding the meta-lated dye zinc-phthalocyanine, ZnPc (Chart 1), by coordination of the pyridine functions to the zinc centers. This coordination initiates the surface-templated epitaxial growth of perpendicular stacks of highly ordered ZnPc, the height of which can be controlled by varying the immersion time of the layer into a solution of ZnPc. In turn, this controlled molecular makeup of the alignment layer allows for rational control over its interaction properties with mesogens, leading to tunable anchoring energies. Moreover, the properties of the alignment layer can be further tuned by adding to the liquid crystalline matrix of a display after its construction, compounds that are capable of interacting with the stacks of dyes (nitrogen ligands 4–9).

2. Experimental Methods

2.1. Synthesis. Unless stated otherwise, all reactions were carried out under Schlenk conditions in a nitrogen atmosphere. Toluene, THF, and hexane were distilled from sodium/benzophenone prior to use. Pyridine was distilled from CaH2. Thionyl chloride was distilled under nitrogen and within 5 min after isolation of the acid chloride by stirring with 10 mL of thionyl chloride at 40 °C. Pyridine-functionalized siloxane 2 (Scheme 1), by including the pyridine-functionalized siloxane 2 (Chart 1), was synthesized in a two-step reaction with the stacks of dyes (nitrogen ligands 4–9). The pyridine-functionalized siloxane 2 was synthesized in a two-step reaction with the stacks of dyes (nitrogen ligands 4–9).

(9) See Supporting Information for further details.
conventionally rubbed polyimide plate as a counter plate.\(^9,13\) In the case of the ZnPc alignment layers, plates with shear-induced alignment were used in anchoring energy measurements. Other LCDs were prepared by using one sample plate and a counter plate which was spin coated with Polyimide Pyralin PI2555 (HD Microsystems) at 5000 rpm for 20 s and baked at 120 °C for 90 min. Mylar spacers of 6 \(\mu\)m were used to separate the plates.\(^8,17,19\) LCDs were filled with 4-n-pentyl-4′-cyanobiphenyl (5CB) in the isotropic phase at 40 °C \((T_{NI} = 35.3 °C)\) and were submitted to a heating—cooling cycle between 20 and 40 °C three times to exclude any memory effects.\(^8,19\) Solutions of nitrogen-containing compounds in 5CB were pulled into the liquid crystal cell via capillary force in 15 min. The concentrations used were as follows: 4 in 5.9 mM and 14 \(\mu\)M, 5 in 61 \(\mu\)M and 139 nM, 6 and 7 in 4 \(\mu\)M and 8 and 9 in 4 mM.

3. Results and Discussion

3.1. Physical Characterization of Alignment Surfaces.

Pyridine-containing alignment layers were constructed by immersing an ITO plate into a solution of 2 and 3 in toluene. When these ITO plates were examined with atomic force microscopy (AFM), a surface containing micrometer-sized grooves was observed, resembling those previously reported by us (Figure 2A).\(^8,9\) After this layer was reacted with a 7.1 \(\mu\)M chloroform solution of ZnPc for 3 h, AFM studies revealed the presence of large objects on the surface with a height of ca. 50 nm (Figure 2A), which we attribute to the presence of stacks of aggregated ZnPc molecules.\(^20\) The height of the stacks could be controlled by varying the immersion time of the ITO plate into the ZnPc solution. The maximum height increased from 25 nm after 30 min to 50 nm after 3 h.\(^9\) Longer immersion times did not yield higher stacks, suggesting that they are in thermodynamic equilibrium with ZnPc in solution. As the critical aggregation concentration of ZnPc was determined to be at least 3 orders of magnitude higher than the concentration used in this experiment,\(^9\) these results imply that the coordination of single molecules of ZnPc from solution to the pyridine functions in the alignment layer results in the surface-templated growth of stacks, which, to our knowledge, is unprecedented for phthalocyanines.\(^21\)

UV—vis experiments clearly showed that 2 coordinates to ZnPc.\(^9\) To investigate whether the binding of ZnPc to the surface could be attributed to interactions other than coordination, e.g. van der Waals forces or \(\pi-\pi\)-interactions, an ITO plate was covered with only 3 from the vapor phase and reacted with a 7.1 \(\mu\)M solution of ZnPc for 3 h, followed by washing with chloroform. AFM and FTIR confirmed the absence of ZnPc.

(stacks) on the surface. Furthermore, experiments showed that under the same experimental conditions, the free-base phthalocyanine did not form aggregates on the pyridine-containing alignment layer and was rinsed off completely with chloroform. These experiments support the idea that coordinative binding of the phthalocyanine is required.

The width of the objects did not display a time dependence. It varied between 25 and 190 nm, which is too large for a single ZnPc stack. Although this width can be partially attributed to AFM-tip effects, it is very likely that a cluster of ZnPc stacks is present, which is anchored to the surface by several pyridine functionalities and held together by van der Waals interactions between adjacent ZnPc stacks (schematically depicted in Figure 2A). The aspect ratio (h/w, as calculated by AFM) of the stacks after 3 h of immersion supports this notion, as it varies from 0.21 to 0.53 for stacks between 45 and 55 nm in height, suggesting the presence of multiple stack types. Aggregation of this type has been observed before by us for similar dye systems, albeit not epitaxially templated by a surface.20,22,23 The distribution of the number of ZnPc aggregates after 3 h of immersion was not constant over the area of the ITO plate, which measured 10 mm × 15 mm. It ranged from 13 to 31 stacks per μm², with an average of 20 stacks per μm², measured over 4 AFM micrographs with a total surface area of 60 μm². Assuming that a stack of 50 nm contains about 150 ZnPc molecules,20 these results imply a ballpark figure for the surface coverage of ZnPc of 10¹¹ molecules per cm², or 10⁻¹³ mol per cm².

Polarized FTIR measurements on the CH-stretching vibrations of the alkyl tails of the stacks showed that their ordering is clustered in rather large domains of several hundred square micrometers but that the domain orientations differ from place to place on the surface, as indicated by a phase shift in their respective polarization angle-dependent absorption curves (Figure 3A). Remarkably, FTIR measurements showed that the application of a shear force, i.e., the unidirectional rinsing of the plates with chloroform, caused this phase shift to disappear, indicating a similar orientation of the alkyl tails over the entire sample plate (Figure 3B,C), offering a second degree of control over the surface ordering. From these data, it is difficult to determine an absolute angular direction or conformation of the alkyl tails on the surface. However, the remarkable angle dependence of the absorption profiles suggest a (partial) rearrangement of the alkyl tails upon unidirectional rinsing. The FTIR experiments also provided information on the orientation of the phthalocyanine rings in the stacks. Both the order parameter, S, and the average tilt angle of the aromatic rings of the phthalocyanine dye molecules with respect to the surface, Ψ, can be calculated from the polarized FTIR measurements. From eight spots on one plate, an order parameter S of 0.22 ± 0.04 and an angle Ψ of 51.4 ± 1.1° were calculated (the ordering is schematically depicted in Figure 2A). Interestingly, S and Ψ remained the same with or without application of shear force, which implies that rinsing had little effect on the angle between the aromatic plane of the ZnPc molecules in the stack and the surface and that the aggregates are firmly linked to the surface.

3.2. Effects on Liquid Crystal Alignment. When the alignment layer, prepared by the deposition of 2 and 3 but without exposure to the ZnPc solution, was used in the
construction of a parallel liquid crystal cell with 5CB as the mesogen, alignment of the mesogen due to the groove structure was observed with domain sizes in the order of several tens of micrometers (Figure 2B). Remarkably, when the plates containing the tallest ZnPc stacks were tested, the domain size increased to hundreds of micrometers, indicating that the alignment capability had increased as a result of the presence of the stacks (Figure 2C). The directors of the liquid crystal domains were not uniformly aligned, which can be attributed to the clustering of the alkyl tails of the ZnPc stacks, causing adjacent domains to be oriented differently, which was supported by the polarized FTIR experiments (vide supra). However, when the alignment surfaces that had been subjected to a shear force were used in the construction of the liquid crystal cells, the domain boundaries of the liquid crystalline compound disappeared completely (Figure 2C, inset). This implies the formation of uniform alignment throughout the entire liquid crystal cell, extending over long distances up to 1.2 cm, which was also supported by the FTIR experiments on the shear aligned surfaces (vide supra).

The enhanced alignment capability of the layer could be affected by introducing compounds into the cells that are capable of disturbing the stacks of dye molecules, e.g. nitrogen donor ligands 4 and 5 or amino acids 6 and 7 (Chart 1). By pulling micromolar concentrations of these ligands through the cells, a wave front was seen traversing the cell, leaving a nonaligning layer behind (Figure 2D). This effect is ascribed to the coordination of the nitrogen-containing molecules to the zinc metal centers in the dyes24–26 followed by the dissolution of the stacks. AFM indicated that after the addition of the nitrogen donor compounds, the stacks of ZnPc on the surface had disappeared (Figure 2A), which was confirmed by FTIR which also revealed the presence of molecularly dissolved ZnPc in the mesogenic medium. To test the selectivity of the effect with respect to nitrogen-donor compounds, benzene, cyclohexanone, and toluene were pulled through liquid crystal cells in millimolar concentrations. Polarizing microscopy revealed that the quality of the liquid crystal alignment was not affected, supporting the notion that coordinative dissolution of the ZnPc stacks is responsible for the loss of liquid crystal alignment observed in the other experiments. Furthermore, 5CB itself, which is capable of coordinating metals with its cyano group and is present in a high concentration, did not influence the alignment quality, suggesting that only more strongly binding amines are capable of dissolving the ZnPc stacks. In addition, the introduction of the sterically hindered 2,6-di-tert-butylpyridine 8 in millimolar concentrations, which is not capable of coordinating to zinc and hence of disrupting the stacks, did not have any adverse effects on the alignment properties of the layer, further supporting the idea that the dissolution of ZnPc stacks from the surface is responsible for the loss of alignment quality. Remarkably, when the chiral bidentate amine (1R,2R)-1,2-diphenylethane-1,2-diamine 9, in a concentration of 4 mM, was used to disrupt the uniform alignment of a shear-aligned cell, polarizing microscopy showed that a cholesteric liquid crystal phase was formed, embedded in a normal nematic phase (Figure 4).27 Blank experiments indicated that 9 by itself is not capable of inducing a cholesteric phase inside 5CB. As such, the occurrence of this phase has to be ascribed to the coordination complex of 9 and ZnPc. The ZnPc aggregates are not uniformly distributed over the surface, which implies that, immediately upon their dissolution, there are large local differences in the concentration of the coordination complex. It is therefore likely that the cholesteric droplets only appear in regions where enough ZnPc has been dissolved into the 5CB. These experiments show that the ZnPc-modified alignment layer can be easily influenced by even micromolar concentrations of nitrogen-donor ligands and respond distinctly upon the addition of a varying concentration of different amines.

These above-described effects were also reflected in the anchoring energy, which is a property of alignment layers that quantifies the interaction strength between the layer and the mesogenic molecules on top of it.8,9,13–19 For the alignment layer composed of 2 and 3 and using the common mesogen 5CB in the cell, this energy was measured to be 9.0 × 10^{-6} J m^{-2}. Upon immersion of the plate into the ZnPc dye solution, this value increased rapidly and leveled off at 4.8 × 10^{-5} J m^{-2} (Figure 3D), showing that the anchoring energy, like the stack height, could be controlled by varying the immersion time. Furthermore, the final value of the anchoring energy is a factor of 2–3 higher than that reported for rubbed polyimide alignment layers currently employed by industry.3,19 The increase in anchoring is attributed to the superior molecular interaction of the ZnPc stacks with the bulk of the liquid crystal, compared to that of conventional alignment layers: the ZnPc stacks extend 50 nm into the bulk of the 5CB, whereas in the case of a conventional alignment layer, only interaction with the first few layers of liquid crystalline molecules takes place. Experiments also indicated that the anchoring energy could be tuned. The addition of nitrogen-donor compounds 4–7 in micromolar concentrations induced a complete loss of anchoring (vide supra). Even the addition of nanomolar concentrations of these compounds had a strong effect. For instance, 4 gave a complete loss of anchoring energy upon addition in nanomolar concentrations, whereas 5 reduced it by a factor of 2.5 (Figure 3D). These combined results indicate that the properties of an alignment layer can be rationally manipulated.

4. Conclusions

In conclusion, we have presented a simple procedure for the construction of a stable and tunable alignment layer for mesogens, which is based on hierarchical assembly processes. The interaction of a spontaneously formed pyridine-containing

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alignment layer with single ZnPc dye molecules in solution results in the surface-templated growth of perpendicular stacks of which the size can be controlled by varying the immersion time. This control over the stack size is directly translated into control over the alignment properties of the alignment layer. Moreover, the introduction of compounds that interact with and destroy the stacks offers a second degree of control over the properties of the alignment layers, which have exhibited stability under ambient conditions for periods exceeding 18 months. The procedure presented here offers possibilities to develop very sensitive LCD-based sensor systems\textsuperscript{28–31} for the detection of, for example, proteins, nucleic acids, and other biomolecules that can bind to ZnPc stacks via nitrogen donor ligands, as well as systems that rely on ordered dye assemblies for optimal performance, such as photovoltaic devices.\textsuperscript{32}

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**Supporting Information Available:** Oligomer characterization, additional physical characterization of the alignment layer and aggregates, spectral data on the binding of 2 to ZnPc, and determination of the critical aggregation concentration of ZnPc. This material is available free of charge via the Internet at http://pubs.acs.org.

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