# Bistable Property in a Splay Cell with a Chiral Additive

Seo Hern Lee<sup>a\*\*</sup>, Tae Jin Kim<sup>a\*\*</sup>, Jong Lac Lee<sup>a</sup>, Mi Jun Jung<sup>a</sup>, Kyoung-Ho Park<sup>a\*\*</sup>, Gi-Dong Lee<sup>ab\*\*</sup>, Tae-Hoon Yoon<sup>ab\*</sup>, and Jae Chang Kim<sup>ab\*</sup>

#### **Abstract**

In this paper, we propose the bistability the bistability in a chiral-splay nematic liquid crystal cell, which is obtained by adding a chiral additive to a splay cell and introduce a novel switching method for the bistable chiral-splay nematic cell in order to transit the LC director between the non-twisted state with splay and the 180° twist state.

Keywords: bistable LCD, splay cell, bend state

### 1. Introduction

Since it was shown that it was possible to switch nematic liquid crystal cells between two metastable twist states in early 1980s, there has been increasing interest in bistable liquid crystal devices. Up to now, the volume<sup>2-4</sup> and surface switching types<sup>5-6</sup> have been mainly demonstrated as bistable devices. In the case of the volume switching type, there is a serious problem in that the  $\phi$ - $\pi$  and  $\phi$ + $\pi$  twist states are metastable with a short lifetime. Although Wang et al.<sup>7</sup> achieved a longterm bistability by using the multidimensional alignment method to prevent the nucleation of the  $\phi$  twist state, there was still limitations in the application. The surface switching type<sup>8</sup> using the effect of surface anchoring, except for the surface-controlled bistable nematic (SCBN) cell<sup>6</sup> in which standard anchoring layers have been demonstrated, have still the demerit that the manufacturing process for the surface alignment is

troublesome in compared with the conventional liquid crystal cells. In this paper, we propose a bistable chiral-splay nematic(BCSN) liquid crystal device using a new horizontal switching, that can overcome the weak points of the volume and surface switching bistable devices. This device is similar to SCBN in that the two bistable states are the 0° state and the 180° twist state. However, unlike SCBN that uses the breaking of surface anchoring conditions using high voltage with short pulse width, the proposed bistable cell can achieve the metastable 180° twist state with a low voltage of only 5 V under the experimented sample size of 3×3 cm² or 2 V/µm under the normal pixel size of 100×100 µm². Hence, the bistable mechanism of this device is irrelevant to the breaking of surface anchoring.

## 2. Transition Process

Fig. 1 illustrates the transition process of the proposed chiral-splay cell. When no voltage was applied, the splay cell was in a parallel state, where the molecules were all aligned with the rubbing direction. In such geometry, when the voltage above  $V_c$  is applied to the cell, it would transform in to a bend state through the middle state with reverse tilt-like domains. The transition time from the splay state to the bend state depends on the applied voltage, pulse duration, and LC materials. For example, in the case of our sample cells of  $3\times3$  cm<sup>2</sup>, in

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<sup>\*</sup> Member, KIDS.; \*\* Student Member, KIDS.

Corresponding Author: Seo Hern Lee

a. Department of Electronics Engineering, Pusan National University, Pusan 609-735, Korea.

b. Research Institute of Computer, Information and Communication, Pusan National University, Pusan 609-735, Korea.

**E-mail**: jckim@pusan.ac.kr **Tel**: +51 510-2375 **Fax**: +51 515-5190

order to transform the splay state to the bend state it is necessary to apply a voltage of 12 V square wave of 1 kHz for about 12 seconds. In the case of the voltage of about 5 V, it takes up about 40 seconds to transform the splay state to the bend state. In general, once the applied voltage is above 4 V, we can obtain a bend state by controling the pulse duration. After the bend state is generated, it returns to the splay state through the 180° twist state during the voltage-off state. While the retention time of the twist state is few seconds in the pure splay cell without a chiral additive, chiral additive increases the retention time above about ten time in the splay cell blended a chiral additive with a cell thickness over pitch (d/p) of 0.1. However, if d/p ratio is above approximately 0.25, careful treatment is needed since the initial state with no voltage may be the twist state. The twist state is more stable as d/p ratio increases. As a result, since the voltage for transforming the twist state to the splay state can be increased in the switching of the splay cell with chiral additive, it is important to optimize d/p ratio.

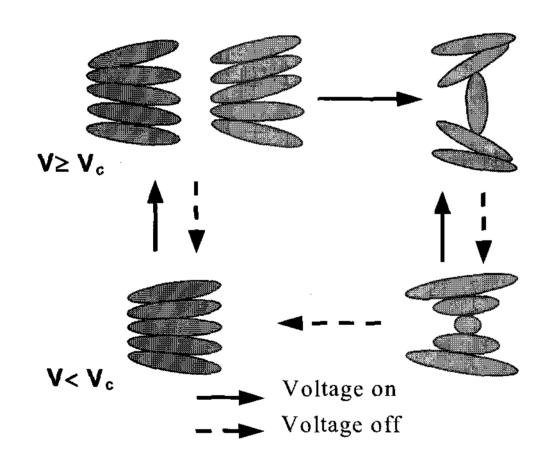


Fig. 1. The transition process of the BCSN LC cell.

#### 3. Transition Mechanism in 180° Twist State

By numerically solving the Ericken-Leslie hydrodynamics equations with a natural twist term of a nematic sample of thickness d aligned parallel along the x axis, we can understand how this BCSN cell switches from the initial  $0^{\circ}$  splay state to the metastable 180 (twist state. The hydrodynamics equations may be expressed by four equations 9, which is composed of the two equations for shear force(() across the liquid crystal cell and the other two balance equations of elastic, viscous, and electric torques. By using these equations and through simple calculations, we obtain:

$$V'_{,} = \frac{(\sigma_{x_{1}}\gamma_{1} - T_{13}\lambda_{1} - T_{14}\lambda_{2})(T_{22}\gamma_{1} - T_{23}T_{32} - T_{24}T_{42})}{\xi_{1}}$$

$$- \frac{(\sigma_{x_{1}}\gamma_{1} - T_{23}\lambda_{1} - T_{24}\lambda_{2})(T_{22}\gamma_{1} - T_{23}T_{32} - T_{24}T_{42})}{\xi_{1}}$$

$$V'_{,} = \frac{(\sigma_{x_{1}}\gamma_{1} - T_{13}\lambda_{1} - T_{14}\lambda_{2})(T_{22}\gamma_{1} - T_{23}T_{32} - T_{24}T_{42})}{\xi_{1}}$$

$$- \frac{(\sigma_{x_{1}}\gamma_{1} - T_{23}\lambda_{1} - T_{34}\lambda_{2})(T_{22}\gamma_{1} - T_{23}T_{32} - T_{24}T_{42})}{\xi_{1}}$$

$$= (\gamma_{1}T_{11} - T_{13}T_{31} - T_{14}T_{41})(T_{22}\gamma_{1} - T_{23}T_{32} - T_{24}T_{42}),$$

$$- (\gamma_{1}T_{21} - T_{23}T_{31} - T_{24}T_{41})(T_{12}\gamma_{1} - T_{13}T_{32} - T_{14}T_{42})$$
and 
$$\xi = (\gamma_{1}T_{12} - T_{13}T_{32} - T_{14}T_{42})(T_{11}\gamma_{1} - T_{13}T_{31} - T_{14}T_{41})$$

$$- (\gamma_{1}T_{22} - T_{23}T_{32} - T_{24}T_{42})(T_{11}\gamma_{1} - T_{13}T_{31} - T_{14}T_{41})$$

$$- (\gamma_{1}T_{22} - T_{23}T_{32} - T_{24}T_{42})(T_{11}\gamma_{1} - T_{13}T_{31} - T_{14}T_{41})$$

The ('s are the shear forces across the cell and Vx

Table 1. The simulation parameters of the liquid crystal

Parameter	Value	Parameter	Value
K <sub>11</sub>	9.5×10 <sup>-7</sup> dyne	$\alpha_1$	$6.5 \pm 4$
$K_{22}$	$5.1\times10^{-7}$ dyne	$\alpha_2$	$-77.5 \pm 1.6$
$\mathbf{K}_{33}$	$11.5 \times 10^{-7}$ dyne	$\alpha_3$	$-1.2 \pm 0.1$
$n_{o}$	$1.4993(\lambda = 589 \text{nm})$	$\alpha_4$	83.2 ±1.4
$n_{\rm e}$	$1.6140(\lambda = 589 \text{nm})$	$\alpha_5$	$46.3 \pm 4.5$
$\epsilon_{ m o}$	3.7	$\alpha_6$	$-34.4 \pm 2.2$
$\epsilon_{ m e}$	7.9		

and Vy represent the flow velocities parallel to the x and y axes in the surfaces of the cell. All of the T's and ('s are functions of ((tilt angle)) and ((twist angle)). They are given in the appendix of Ref. 9. With these equations, we can find the influence of flow effects in transforming the bend state to the twist state in the field-off state.

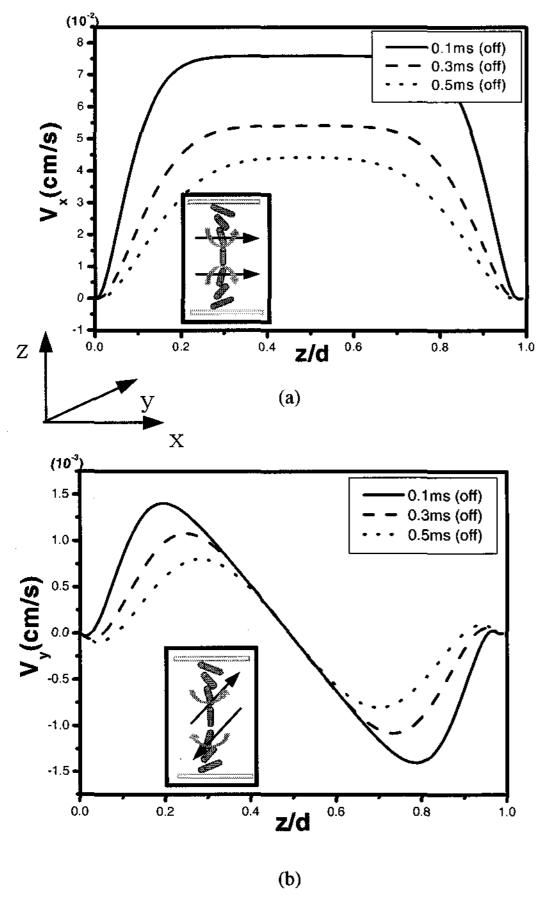


Fig. 2. The calculated flow velocities and the illustrations of the relaxed directors (the inset) during the voltage-off state: (a) Component  $V_x$  in the x direction and (b) component  $V_y$  in the y direction of the flow velocity.

Fig. 2 shows the calculated flow velocities after applying the voltage of a 15V to a test cell of d/p=0.1, cell with thickness of 6.4  $\mu$ m and pretilt angle of 5° for the bend state, and subsequently, disconnecting the voltage source. The parameters used in the simulation are shown in Table 1. Due to the lack of the Leslie coefficients of ZLI-1557, these coefficients were taken from the values of MBBA<sup>10</sup>. There is an obvious influence of flow velocities in the x-direction and y-direction in the relaxation. The velocity component  $V_x$  in the x-direction was the same at the both sides of the cell.

As shown in the inset of Fig. 2(a), the torque

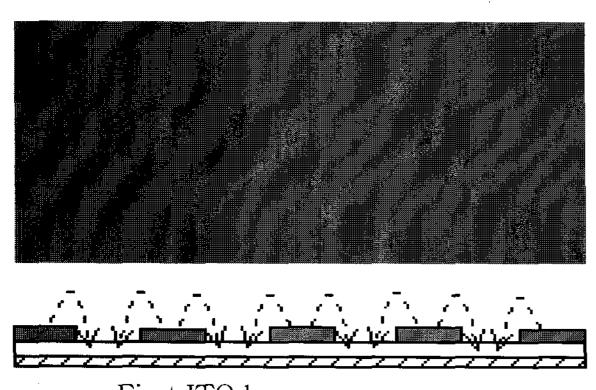
induced by the flow relaxes the liquid crystal directors in the x-direction. The velocity component  $V_y$  in the ydirection plays an important role in forming the 180° twist state. As shown in Fig. 2(b), the flow in the ydirection is reversed symmetrically about the midplain of the cell, which is known as the backflow effect. Although its magnitude is smaller than that of  $V_x$ , the effect of the flow is enough to induce relaxation of the directors symmetrically about the midplain in the y direction as shown in the inset of Fig. 2(b). As a result, the twist state is generated due to the flow-induced viscous torque in the y-direction. For comparison, we performed a computer simulation with the same conditions, except that rubbing was done in an antiparallel direction, which is called a homogeneous state. The calculated results showed that the role of velocity components  $V_x$  in the x-direction and  $V_y$  in the y-direction was inverted unlike a splay cell. That is, in a homogeneous cell, the velocity  $V_x$  in the x-direction was reversed symmetrically about the midplain in the xdirection, and  $V_y$  in the y-direction was the same on the both sides of the cell (not shown). Therefore, we can inter that the flow-induced viscous torque in the xdirection relaxes the liquid crystal directors in the same direction and the twist state is not generated due to velocity components  $V_{y}$  with the same direction on the both sides of the cell.

## 4. Horizontal Switching Between 180° Twist State and 0° Splay State

To confirm the validity of the mechanism, a splay cell filled with liquid crystal ZLI-1557(E. Merck) was fabricated with spacer thickness of 4.2 µm. The indiumtin-oxide (ITO) glass was used as the substrates. The polyimide SE-3140(Nissan Chemicals Co.) was coated on the bottom and top glass substrates, and rubbing was done in parallel direction. The pretilt angle of SE-3140 has been known to be about 5°. The S-811 was used as chiral additive in order to obtain the metastable 180° twist state in a splay cell. A He-Ne laser with wavelength 632.8nm was used as the light source.

To switch the proposed chiral-splay cell, we need to use a special electrode. Fig. 3 shows an electrode structure of the bottom substrate to switch from the metastable 180° twist state to a 0° splay state. By a

vertical switching between ITO layer of the top substrate and the second ITO layer of the bottom substrate, we can transform a 0° splay state to a bend state, and then obtain a twist state during the voltage-off state. By the horizontal switching between the first ITO layer and the second ITO layer of the bottom substrate, we can transform the 180° twist state to the 0° splay state. Fig. 4 shows the equipotential lines that are calculated by 2DIMMOS in the horizontal switching on the used electrode. By the electric field in the horizontal direction, it can be transformed from the 180° twist state to the 0° splay state. Fig. 5 shows the driving pulses used to switch the BCSN cell and the transmittance as a function of time. In Fig. 5,  $V_{st}$  and  $V_{ts}$  represent the vertical voltage for the transition from the 0° splay state to the metastable 180° twist state, and the horizontal voltage for the reverse process, respectively. Although, it can be switched into the twist state with a low voltage of only 5 V. We used a high voltage of  $V_{st}=20 \text{ V}$  and  $V_{ts}=20 \text{ V}$  for a faster switching. The driving pulse width was 150 ms for  $V_{st}$  and 700 ms for  $V_{ts}$ , respectively.



First ITO layer

Passivation layer

Second ITO layer

**Fig. 3.** The electrode structure for the switching from the metastable 180° twist state to the stable 0° state.

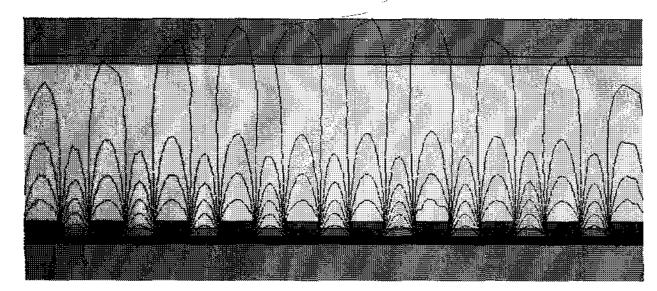


Fig. 4. The calculated equipotential lines in the horizontal switching on the used electrode

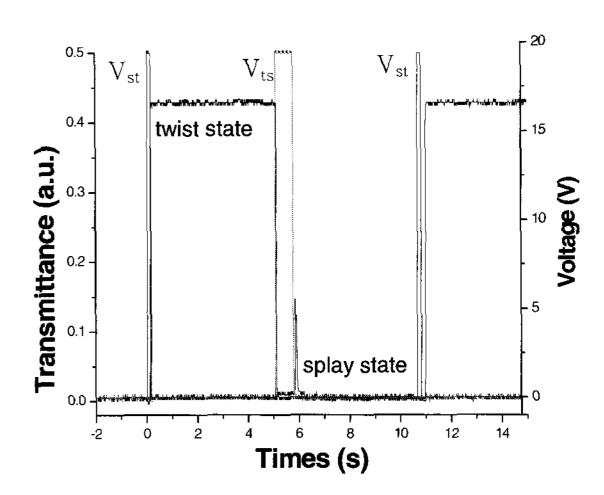


Fig. 5. The driving pulses used to switch the BCSN cell, and the transmittance

### 5. Conclusions

In summary, we have investigated the bistable property in a splay cell with a small amount of chiral additive, and proposed a new way of switching between the two states of the bistable chiral-splay cell. In the case of proposed chiral-splay cell, unlike the conventional BTN, bistable property is easily obtained in both the large cell gap and the low voltage. By optimizing the various LC parameters, we expecte to obtain a noble BTN device with superior optical as well as electrical properties.

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