Spatially tunable laser emission in dye-doped photonic liquid crystals

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A spatially tunable laser emission of the dye-doped cholesteric liquid crystal (CLC) cell using a one-dimensional temperature gradient is demonstrated. The photoexcitation of dye-doped CLC device using a frequency-doubled pulsed Nd: yttrium–aluminium–garnet laser gives rise to laser emission in the yellow-red spectral range. The lasing wavelength is widely tunable from 577 to 670 nm by shifting the position of the dye-doped CLC cell with respect to the pumping beam. The lowest excitation energy and maximum lasing efficiency occur at λ ~ 605 nm which corresponds to the peak fluorescence emission of the dye. © 2006 American Institute of Physics.

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A spatially tunable laser was obtained by filling the CLC cell. A spatially turnable laser was obtained by filling the CLC sample, we can easily obtain a spatially tunable CLC cell. Different from the abovementioned spatially tunable methods, our CLC cell contains only one mixture. The spatial tunability originates from the temperature dependent CLC PBG. The lasing emission wavelength can be tuned from 577 to 670 nm by changing the temperature from 23 to 40 °C. The temperature of the dye-doped CLC cell was monitored by a temperature controller. Figure 1 shows the shift in the PBGs of the dye-doped CLC cells with the increased temperature. The average wavelength (λ_ave) is defined as λ_ave=(λ_long+λ_short)/2, where λ_long and λ_short stand for the wavelength at the long and short reflection band edges, respectively. As the temperature of the CLC cell increases, the selective reflection band of CLC shifts toward a shorter wavelength. The reason is that at room temperature, the maximum solubility of ZLI-811 chiral agent in BL-006 is ~ 25 wt%. However, our sample contains 34 wt% ZLI-811 which is beyond the maximum solubility. Therefore, a portion of the chiral ZLI-811 cannot be dissolved in the LC host and is separated out. As the temperature increases, the solubility of the chiral agent increases which, in turn, reduces the helical pitch length. As a result, the PBG shifts towards a shorter wavelength as the temperature increases. At T>40 °C, nearly all the chiral dopants have been dissolved and, therefore, no more blue-shift occurs as the temperature is further increased.

Based on the temperature dependent optical properties of the CLC sample, we can easily obtain a spatially tunable CLC PBG by generating a 1D temperature gradient across the CLC cell. To achieve this goal, we simply placed one side of the CLC cell on a heating stage and left the other side in the air. By raising the temperature of the heating stage over room temperature, the 1D temperature gradient was formed. Figure 1(b) shows the image of a CLC cell with gradient temperature at T~50 °C. The reflected colors spread from blue to red as the position gets farther away from the heat source.

An important application of the tunable PBG in the CLC system is for tuning the laser oscillation. To test the laser emission properties, a second-harmonic Q-switched Nd:YAG
pulsed laser (from Continuum) at $\lambda=532$ nm with vertical linear polarization was used to excite the dye-doped CLC sample, as Fig. 2 depicts. The pulse width and repetition rate are 6 ns and 1 Hz, respectively. The reason that we chose 1 Hz repetition rate is to avoid sample heating and degradation. A beam splitter was used to divide the incoming laser into two beams: one was sent to a laser energy meter (Ophir) for monitoring the pumping pulse energy and the other used as the excitation beam. A linear polarizer and a quarter-wave plate were used to convert the linear polarization into right-handed circular polarization to avoid the reflection by the CLC PBG. A lens with 15 cm focal length focused the incident beam to a small spot of $\sim160 \mu$m diameter at the sample. One side of the sample was placed on a heating stage ($T=50 \, ^\circ\text{C}$) and the other was left in the air. The heating stage was mounted on a translational stage. The output laser emission in the forward direction of the sample was collected by a lens to a fiber-optics based universal serial bus (USB) spectrometer ($0.04 \, \text{nm resolution; USB HR2000, Ocean Optics}$).

In Figs. 3(a) and 3(b), we plot the normalized laser emission and the corresponding transmission spectra of the dye-doped CLC cells with various PBGs at different sample positions. Positions 1–5 correspond to $T \sim 30.6, 28.8, 27.6, 26.8,$ and $26.0 \, ^\circ\text{C}$, respectively. The gray line represents the dye’s fluorescence spectrum. The laser emission occurs at band edge.
between 577 and 670 nm. This behavior stems from the competition between the optical loss and gain in the CLC system. Generally, the laser emission can be generated through the feedback amplification only when the optical gain inside the medium is larger than the loss. For the CLC cell, since the internal distributed feedback of the CLC PBG provides the amplification of the optical gain traveling in the CLC cell, the laser emission can be obtained when the amplification of the optical gain traveling in the CLC cell is sufficient to overcome the losses in the medium. Typically, the optical gain spectrum has the same spectral appearance as the fluorescence spectrum of the dye. When the CLC PBG is falling within the fluorescence spectrum of the doped dye, the optical emission can be effectively obtained by the internal distributed feedback of the CLC PBG. Therefore, we can obtain laser emission in this range. However, when the CLC PBG is scarcely overlapped with the fluorescence spectrum of the doped dye, the amplification of the optical emission provided by the internal distributed feedback of the CLC PBG cannot yet overcome the optical losses. Thus, no lasing can be observed. It was also observed that the laser emission always appears at the long edge of the CLC PBG because the long edge of the CLC PBG needs lower energy than the short edge to get laser emission.

In addition, we also investigated the dependence of the lasing emission intensity at different wavelengths on the excitation energy. Figure 4 shows the results we obtained from the dye-doped CLC cell with a PBG at different spectral position by adjusting the position of the dye-doped CLC with respect to the pump beam. For any lasing wavelengths, the emission intensity is significantly enhanced as the excitation energy exceeds the threshold. However, the threshold excitation energy and lasing efficiency strongly depend on the lasing wavelength, as Fig. 4 shows. When the lasing wavelength is at λ=605 nm, the lasing efficiency reaches the maximum (~0.5%) and threshold excitation energy is the lowest (~4.7 μJ/pulse). If the lasing wavelength is away from 605 nm, higher threshold excitation energy is required and the lasing efficiency is decreased. We know that laser emission can be obtained only when the optical gain overcomes the losses in the medium and larger optical gain makes the generation of the laser emission easier because the optical emission is more effectively obtained by the feedback effect. As stated above, the optical gain spectrum relies on the fluorescence spectrum of the dye. Since the DCM dye exhibits its maximum fluorescence at λ ~ 605 nm, the lowest threshold excitation energy of ~4.7 μJ/pulse and the highest lasing efficiency occur for the dye-doped CLC sample when the long edge of the PBG is at 605 nm. When the long edge of the CLC PBG overlaps with the tail of the fluorescence band, higher threshold energy is required for exciting more optical emission to overcome the loss and finally generating the laser feedback effect. Consequently, the lasing efficiency is decreased.

In conclusion, we have demonstrated a spatially tunable laser emission by generating a 1D temperature gradient on a thermally sensitive dye-doped CLC cell. The lasing wavelength is tunable from 577 to 670 nm by changing the spatial position of the dye-doped CLC cell. The lowest excitation energy and maximum lasing efficiency are obtained at λ ~ 605 nm since the DCM dye has the maximum fluorescence emission at around 605 nm.