Study on the Third-order Nonlinear Optical Properties of bis(tetraethylammonium) bis(1,3-dithiole-2-thione-4,5-dithiolato)cadium

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ABSTRACT
A dmit²- salt: bis(tetraethylammonium)bis(1,3-dithiole-2-thione-4,5-dithiolato)cadium (CADMIT) was synthesized. The Optical Kerr Effect (OKE) signal of its acetonitrile solution was measured by femtosecond optical Kerr gate technique. Using CS₂ OKE signal as reference signal measured under identical conditions, the third-order optical nonlinear susceptibility, χ(3), of the sample solution was obtained to be about 2.98 × 10⁻¹⁴ esu at the concentration of 1.57 × 10⁻³ M. The second-order hyperpolarizability of its molecular was estimated to be as large as 1.23 × 10⁻³² esu. Its response time was about 195 fs, which is believed to be the contribution from the delocalized electrons.

Keywords: Optical Nonlinearity; Kerr Gate Technology; Response Time

1. Introduction
Recently, nonlinear optical materials have attracted much interest because of their potential in fabricating ultrafast optical switching and processing devices [1-3]. For this purpose, many materials, including semiconductors, polymers, nanomaterials and inorganic materials, have been studied. Among them, π-conjugated organic material has been paid great attention for its high nonlinear optical (NLO) properties and ultrafast response [4]. Especially, the organometallic and coordination material has been attracting great concern, because it has the advantages of architectural flexibility, ease of fabrication and tailoring, and high NLO properties of organics. Simultaneously, it possesses good transmissivity, temporal and thermal stability of inorganic matter. Such material also has strongly enhanced second-order hyperpolarizability by introducing a metal atom.

Currently, the synthesis and characterization of 1,3-dithiole-2-thione-4,5-dithiolate (dmit) complexes and related selenium- and oxygen-substituted isologs have been studied. As a special π-electron delocalization conjugated system, dmit and related ligand complexes have been used in the assembly of highly electrically conducting radical anion salts and charge-transfer complexes. They are generally used as important building blocks for organic, organometallic and coordination complex electrical conductors and superconductors. The π-electron delocalization in conjugated systems can also contribute to the ultrafast response capability and large third-order nonlinearity. Recently, some of these complexes possessing good second-order[5,6] and third-order[7-11] NLO properties have been reported. In this paper, a dmit²- salt: bis(tetraethylammonium)bis(1,3-dithiole-2-thione-4,5-dithiolato)cadium was synthesized. The Optical Kerr Effect (OKE) signal of its acetonitrile solution was measured by femtosecond optical Kerr gate technique.

2. Material Preparation and Experiment Method
The preparation method of CADMIT crystal was a modification according to literature method [12]. The sample solution was prepared using acetonitrile as the solvent whose concentration is 1.57 × 10⁻³ M. The femtosecond optical Kerr gate technique was applied in this experiment whose measurement setup is illustrated in Figure 1. The light source, centered at 800 nm, is a Ti:sapphire femtosecond laser system (Mira 900F, Coherent, USA), which is pumped by a multiline Ar⁺ laser system (Innova 400, Coherent, USA) at 11 W. The repetition rate is 76 MHz and its pulse width is 120 fs. The femtosecond laser beam whose average power was about 0.64 W is split into a probe beam and a pump beam by a beam splitter. The intensity ratio for the probe beam and the pump beam is 1:10. The polarization of the probe beam is carefully adjusted at 45⁰ to the linear polar-
ized pump beam. The pump beam passes through a delay line driven by a step motor. Then it is carefully reflected parallel to the probe beam at an adjacent configuration. The two beams are focused by a convex lens. At the focus of the lens, the two beams overlap each other. This focus would be enclosed within the sample cell of 1 mm in thickness during the measurement. After transmitting through the sample cell, the pump beam is blocked, while the probe one passes an analyzer, of which the transmission axis is strictly perpendicular to that of the probe beam. Finally, the OKE signal was recorded as a function of pump-probe delay by an amplified photodiode and recorded by a digital lock-in amplifier (SR830, Stanford, USA) referenced to the chopper frequency. Both delay stage and lock-in amplifier were controlled by a personal computer.

3. Results and Discussions

The molecular structure of CADMIT is illustrated in Figure 2 and the absorption spectrum of its acetonitrile solution is in Figure 3. Its absorption coefficient at 800 nm is very small and suggest that 800 nm is far from its resonant band. And the absorption around 400 nm is also not strong which indicates its two-photon absorption at 800 nm has small effect on our measurement result. In the OKE experiment, CS2 was used as reference. Since the sample is prepared as a solution, the measured third-order NLO susceptibility, $\chi^{(3)}$, was the combination of the response from both the sample solute and solvent. Therefore in the experiment, we measured the OKE signals of CS2, the solvent, and sample solution, consecutively. With the solvent signal was removed from the sample solution signal, we obtained the pure signal of sample. Then we used the following equation to calculate its third-order NLO susceptibility $\chi^{(3)}$ [13-15].

$$\chi^{(3)} = \frac{I_S}{I_R} \left( \frac{n_S}{n_R} \right)^2$$

(1)

The subscripts S and R represent the sample of CADMIT and reference sample CS2. I indicates the intensity of OKE and n is the refractive index. The concentration of the sample is very low in the solvent, so we use the refractive index of acetonitrile as the index of solution, which is 1.34. The n for CS2 is 1.62. The third order nonlinear susceptibility of CS2 is estimated to be $1 \times 10^{-13}$ esu in femtosecond time scale [16]. Using the equation, we can get the third-order NLO susceptibility $\chi^{(3)}$ of sample directly by measuring the OKE signal intensity of both sample and reference under identical conditions. The OKE signal of the sample solution was measured and illustrated in Figure 4, in which the small contribution from the solvent was subtracted. Using Eq. (1) and the measured signals of CS2 and sample solution, the third-order susceptibility $\chi^{(3)}$ of sample solution was obtained to be $2.98 \times 10^{-14}$ esu at the concentration of 1.57 $\times 10^{-3}$ M. The second-order hyperpolarizability, $\gamma$, of the sample molecule may be estimated through the equation [17,18]

$$\gamma = \frac{\chi^{(3)}}{(NL)}$$

(2)

where $N$ is the concentration of the solution and $L$ is the local field correction factor which is defined as $\left( n^2 + 2/3 \right)$ (n is the refractive index of solution). By Eq. (2), we may calculate $\gamma$ of CADMIT as $1.23 \times 10^{-32}$ esu using the measured $\chi^{(3)} = 2.98 \times 10^{-14}$ esu.

From Figure 4, we obtained the response time of bis(tetraethylammonium)bis(1,3-dithiole-2-thione-4,5-dithiolato)cadium to be about 195 fs, which is commonly accepted to be the contribution from the transient motion of the $\pi$-conjugated-electron distribution. Normally, the time domain response of the sample should contain several processes of different time scales. Using the femtosecond OKE measurement system, some subpicosecond
processes can be distinguished. The third-order NLO response time induced by delocalized electrons is believed to be $10^{-14} - 10^{-15}$ s. While the response time induced by molecular reorientation is $10^{-11} - 10^{-12}$ s and that induced by density change is $10^{-8} - 10^{-9}$ s.

4. Conclusions

CADMIT was synthesized by making some modifications to the literature method. The absorption spectrum of its acetonitrile solution was measured, which shows that the salt has good transmittance from 600 nm to infrared band. Its third-order optical nonlinearity $\chi^{(3)}$ was studied by a transient OKE measurement and was about $2.98 \times 10^{-14}$ esu at the concentration of $1.57 \times 10^{-3}$ M. Its third-order optical nonlinearity response time was obtained and be about 195 fs, which is commonly accepted to be the contribution from the transient motion of the π-conjugated-electron distribution. Its second-order hyperpolarizability, as large as $1.23 \times 10^{-32}$ esu, was estimated. These features indicate CADMIT is a potential photonics material in future.

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REFERENCES


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