EFFECT OF pH ON THE GROWTH AND CHARACTERIZATION OF GLYCINE SODIUM CHLORIDE (GSC) SINGLE CRYSTAL

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ABSTRACT

Single crystals of semi-organic non-linear optical Glycine Sodium Chloride (GSC) have been successfully grown from three different pH (1.1, 6.0, and 10.8) solutions by evaporation solution growth. In this report we bring out the influence of pH on the optical, structural and NLO properties of the grown crystals. The grown crystals have been subjected to powder X-ray diffraction studies to identify the crystalline nature. Single crystal X-ray diffractometer was utilized to measure the cell parameters and morphology of the grown crystals. The FTIR spectra taken for the crystals grown at different pH values show variations in the peak intensity. The NLO property is found to be varying with the change in the pH values.

Keywords: re-crystallization, seed crystals, semi-organic, nonlinear optical materials.

INTRODUCTION

Non-linear optical materials (NLO) have wide applications in the area of laser technology, optical communication and in storage technology. The high nonlinearity makes it possible for organic crystals to double the frequency of Ga-Al-As diode lasers for generating blue light which is an important coherent light source. However, the short comings of the aromatic crystals such as poor physico-chemical stability, low hardness and cleavage tendency hinder their device applications¹⁻³. In order to keep the merits and overcome the shortcomings of organic materials some new classes of NLO crystals such as LAP, GSC, GSB, GSN and KDP (semi-organic complex crystals) to combine the advantages of inorganic crystals such as good stability, with the advantages of organic crystals such as high nonlinearity, are of special interest. Among the semi-organic complex NLO crystals the nonlinearity of GSC is relatively high (GSC). GSC belongs to the monoclinic system with the molecular formula H₃NCICH₂COONa. The cell dimensions are a = $5.1061A^0$, b = $11.9765A^0$, c = 5.4652A0 and $\beta = 111.78^0$ The SHG intensity of GSC is 0.5 times that of KDP. In the present study, the effect of pH on the growth and properties, such as NLO, structure of crystal, and optical are studied.

EXPERIMENTAL

Preparation of GSC solution at different pH values:

The commercial reagent of Glycine (purity 99%) procured from Merck company was purified by the repeated recrystallization using hot deionised water solution by slow cooling. At pH 6.0, essentially the glycine is in the zwitterionic form. Glycine and Sodium Chloride in the equimolar ratio 1:1 were taken in a beaker containing double distilled Millipore water at room temperature and the saturation solution is brought to pH 6.0. The reaction takes place based on the following equation and the salt is formed.

$$H_2N - CH - COOH + Na Cl --- \rightarrow H_3N - CH_2 - COONa$$

The purity of the synthesized salt was further increased by successive recrystallization process. The saturated GSC solutions of different pH (1.1, 6.0, and 10.8) have been prepared using doubly

recrystallised salt. The solution was filtered using a sintered glass filter of 1u porosity. The solution is transferred to three Petri dishes and covered by a porous paper and kept for slow evaporation process.

Growth of crystals:

Seed perfection and selection of solvents are very important for growing single crystals of high quality ^{6,7}.Defects in seed crystal could cause spurious crystallization and flaws during the crystal growth. To ensure good quality crystals, a seed was obtained by spontaneous nucleation in supersaturated solution of GSC at room temperature. Transparent and good quality seeds were selected for the growth. The grown crystals were harvested after a typical growth period of 30 days.

Structure and morphology:

In order to confirm the crystallinity of the grown crystals, powder X-ray diffraction (XRD) pattern has been recorded using a Rich Seifert Diffactometer (Model 2002) using Cu K α ($\lambda = 1.5418$ Ű)radiation The samples were scanned over the angle range 10-50° at the rate of 0.05° /min. The changes in the unit cell parameters and the morphology of the grown crystals have been observed from single crystal XRD analysis using ENRAF NONIUS CAD4-F (FR 590) single crystal diffractometer. (Table-1).

FTIR spectral studies (fig-1,2,3):

The infrared spectra were taken using Brukker IFS66V FTIR spectrometerby KBr pellet technique to confirm the presence of different organic groups along with the inorganic materials presence.(Table-2)

XRF studies:

The X-ray Fluorescence is taken for the samples to study the abundance of Na and Cl to be present in the crystals.

NLO efficiency :

The comprehensive analysis of second order non-linearity of GSC crystals, grown from different pH was performed by Kurtz powder method⁸. The single crystals of GSC grown from different pH values were irradiated by an incident radiation (1064 nm)of pulse width 8ns and pulse energy of 10-800mJ from a Qswitched quanta RAY GCR Nd:YAG laser. KDP was used for calibrating the SHG intensity. A concave mirror, collimated and focused onto the monochromator slit collected the SHG emitted from the crystal sample. The output power of the crystal was measured using OPHIR power meter model DG with power head model OPHIR 30A. The error in the measured SHG was typically about 5-10%. The NLO property of the crystal was confirmed from the estimation of strong green radiation of the crystal. The results of SHG efficiency of GSC are summarized in table -4. The GSC crystals grown at different pH values showed very good stability under laser irradiation and no signs of decomposition; crack or fracture were observed even for continued irradiation with laser power of 800 mW.

RESULTS AND DISCUSSION

The growth kinetics and the quality of the crystals grown from solutions are considerably influenced by pH of the solution. The amino acids contain both acidic

(-NH₃) and basic (-COO) groups, they are amphoteric. The predominant form of the amino acid depends on the pH of the solution. In an acidic solution, the COOH group and the molecule has an overall positive charge. As the pH is lowered, the - COOH loses its proton at about pH 2. This point is called pK_{a1}), the first acid-dissociation constant. As the pH is raised, the $-NH_3^+$ group loses its proton at about pH 9.6. This point is called pK_{a2} , the second acid - dissociation constant. Above this pH, the molecule has an over all negative charge.⁸So, we have grown crystals of GSC at: 1. pH of 1.1 which is less than pH 2 and 2.pH of 10.8 which is greater than pH 9.6. At pH 1.1 the GSC crystals show high intensity of SHG, the crystal structure is Hexagonal. At pH 6.0 the GSC crystals show very low intensity of SHG, the crystal structure is monoclinic. At pH 10.8 the GSC crystal shows still lower intensity of SHG and the crystal structure is Hexagonal.

From the FTIR studies we find for pH = 1.1, the absorption peak at 3444 cm⁻¹ rather than OH may indicate the presence of NH bond. NH stretching (amide) may be indicated by the absorption band at 1632 cm and second peak of absorption at 1495 cm⁻¹ may confirm the NH stretching. The C=O presence

may be indicated by the absorption bands at 1331 cm⁻¹ and 1254 cm⁻¹. The CH₂ vibrations are indicated by the absorption bands at 887 cm⁻¹, 672 cm⁻¹ and 502 cm⁻¹.⁹

The FTIR for pH = 10.8 shows, the presence of NH bond is indicated by the absorption peak at 3436 cm⁻¹ and NH stretching (amide) is indicated by the absorption at 1672 cm⁻¹ and the second peak of absorption at 1481 cm⁻¹, confirms the NH stretching. The C=O presence is indicated by the absorption band at 1333 cm⁻¹. The CH₂ vibrations are indicated by the absorption bands at 893 cm⁻¹ 686 cm⁻¹ and 501 cm⁻¹.

From the XRF studies we find the existence of Cl in the GSC pH = 1.1 and this enhances the NLO property. Further the existence of Na is more in the in the GSC, pH = 10.8 and this lowers the NLO property.

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Compd.	а	b	с	α	β	γ	Cell vol.
GSC(pH = 1.1)	7.0335	7.0449	5.4761	90	90	120	235.2621
GSC(pH = 6.0)	5.1061	11.9765	5.4652	90	111.78°	90	310.3383
GSC(pH=10.8)	7.0323	7.0338	5.4916	90	90	120	235.3147

Table-1

Table-2

GSC at pH=10.8					
Frequency cm ⁻¹	Assignment of vibration				
3436 (s)	Cisoid secondary amide				
1627 (s)	C = O stretching				
1489 (w)	NH in plane bending				
1439 (m)	CH ₃ bending				
1410 (s)	CH ₃ bending				
1333 (m)	C – N Stretching				
1125 (w)	CH ₃ symmetric bending				
1040 (w)	$= CH_2 rocking$				

929 (m)	- CH out of plane bending					
893 (w)	- COO stretching					
686 (m)	- CH out of plane bending					
GSC at pH = 6						
2923 (s)	CH ₂ symmetry stretching					
2854 (s)	CH ₂ symmetry stretching					
2603 (s)	NH ₂ ⁺ asymmetric stretching					
2171 (m)	S-C = N stretching					
1600 (s)	C=O stretching					
1500 (s)	NO ₂ symmetry stretching					
1436 (s)	N=O symmetry stretching					
1394 (s)	N=O stretching					
1323 (s)	C-N stretching					
1126 (s)	C-O-C symmetry stretching					
1043 (m)	C-O stretchingand O-H plane bending					
929 (s)	CH ₂ out of plane bending					
889 (s)	CH ₂ out of plane bending					
684 (s)	O-N=O bending					
	GSC at pH = 1.1					
3444 (s)	Cisoid secondary amide					
3238 (m)	N-H stretching					
2609 (s)	NH ₃ stretching					
1632 (s)	NO ₂ asymmetry stretching					
1495 (m)	N-H in plane bending					
1410 (m)	C – O Stretching					
1331 (m)	C-N stretching					
1254 (m)	C-N stretching					
1125 (m)	N-H stretching					
1027 (w)	C-O stretching					
887 (m)	C-O-O stretching					
672 (w)	O-C-N stretching					

Table-3: Power output of SHG signal developed in the KDP and the GSC crystals (grown from different pH)

KDP mW	GSC pH = 1.1 mW	GSC pH = 6.0 mW	GSC pH = 10.8 mW
150	200	125	100
290	310	240	200
330	290	210	150



Fig-1: GSC-pH=1.1



Fig 2: GSC-pH=10.6



Fig 3:GSC=pH-6.0

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