SYNTHESIS OF BARIUM TITANATE NANOMATERIAL BY CO-PRECIPITATION METHOD AND CHARACTERIZATION

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Abstract: Barium titanate $(BaTiO_3)$ nanoparticles have been synthesized by co-precipitation method. XRD analysis was used to study the structure and estimate the size of the particles. Scanning electron microscopy was used to study the morphology of the material. Thermogravimetric analysis (TGA) of BaTiO₃ particles.

Index Terms: Barium titanate, co-precipitation, nanomaterial, TGA.

I. INTRODUCTION

A material is a chemical substance or mixture of substances may be natural or man-made constitute foundation of technology [1]. Materials Science deals with the study of relationship between, processing structure, and properties of the materials. Nanotechnology is science, engineering, and technology conducted at the nanoscale, which is about 1 to 100nm combines knowledge from the fields of Physics, Chemistry, Biology, Medicine, Informatics, and Engineering. Barium titanate is the inorganic compound having oxides of barium and titanate with the chemical formula BaTiO₃ exist in the form of white powder and transparent as larger crystals. It is an dielectric materials due to its high dielectric constant, positive temperature coefficient, and nonlinear optical properties [2-3]. It has five phases as a solid, listing from high temperature to low temperature: hexagonal, cubic, tetragonal, orthorhombic, and rhombohedral crystal structure. All of the structures exhibit the ferroelectric effect except cubic. It is insoluble in water and soluble in concentrated sulfuric acid [4-6].

II. MATERIALS AND METHODS

The analytical grade materials were used to synthesize the required samples.

Synthesis of BaTiO3: The co-precipitation method was adopted to prepare BT nanoparticles.0.1M Titanium tetrachloride (TiCl₄) solution and 0.1M Bacl₂.2H₂O solutions were prepared in distilled water and both of these solutions were mixed and stirred for 5 minutes.0.1M solution of oxalic acid taken in 1000ml beaker and stirred for 2minutes and a mixture solution of TiCl₄ and Bacl₂ was added slowly drop wise from the burette with continuous vigorous stirring for 2 hours. The precipitation was allowed to settle down and dried at room temperature and grinded with mortar and pestle for 10minutes and calcinated for 2hours at 450°C [7].

III. RESULTS AND DISCUSSION

The XRD pattern of BaTiO₃ is as shown in Figure-1. The XRD pattern of BaTiO₃ reveals crystalline structure of the sample



formation of tetragonal phase of BaTiO₃, which is approved by the appearance of X-ray reflections at 2θ values 22.152°, 31.528°, 38.889°, 45.273°, 50.855°, 56.235°, 65.797°, 70.586°, 74.96° and 79.153° (JCPDS 05-0626). The average crystallite size of the BaTiO₃ crystals is estimated to about 20.766 nm by Scherrer's formula.

The SEM image of BaTiO₃ particles was obtained to analyze their surface morphology. The SEM photograph BaTiO₃ of magnification 20kx ,in Figure-2, indicates the existence of its crystalline structure. Both large and small size particles are appearing corresponding to crystal growth and well dispersion respectively.

The thermogravimetric analysis (TGA) graph of BaTiO₃ under heating rate of 10° C min⁻¹ is shown in Figure-3. The water weight loss is 2.71% up to the temperature 309°C. The weight loss at the temperature range 310°C to 660°C is 2.84% is due to of entrapped, carboxylates, nitrates, and unburnt carbon. And at the temperature range 660°C to 1000°C weight loss is



Figure-2: SEM image of BaTiO₃ nano-

1.94% due to decomposition of residual BaCO₃.

IV. CONCLUSION

The above discussion outlines the synthesis and characteriation BaTiO₃ nanomaterial. In this experimental work we have



Figure-3: Tthermogravimetric curve of BaTiO₃ nano-material

adopted co-precipitation method to synthsize BaTiO₃ powder. The particle size estimated as 20.766nm.SEM was used study the surface morphology of BaTiO₃ nanoparticles.TGA provides the knowledge of thermal properties of BaTiO₃ nanomaterial.

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