



# Solution Combustion Method (SCM) For The Preparation Of $\text{Bi}_{1-x+y}\text{Ba}_x\text{Ca}_y\text{FeO}_3$ (X, Y = 0.1, 0.15, 0.2, 0.25) Nanopowder Samples

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**Abstract:** This paper describes the synthesis of nanopowder multiferroic ceramics samples such as  $\text{Bi}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{FeO}_3$ ,  $\text{Bi}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{FeO}_3$ ,  $\text{Bi}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{FeO}_3$  and  $\text{Bi}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{FeO}_3$  samples using solution combustion method. These nanopowder samples of  $\text{Bi}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{FeO}_3$ ,  $\text{Bi}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{FeO}_3$ ,  $\text{Bi}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{FeO}_3$  and  $\text{Bi}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{FeO}_3$  were prepared using metal nitrates and glycine fuel as preliminary precursors. These nanopowder ceramics were grinded finally in an acetone medium, calcined at different optimized temperatures and finally pelletized in the form of pellets.

**Keywords:** Multiferroic, Nanopowders,  $\text{BiFeO}_3$ , Ba and Ca dopants, SCM, applications.

## I. INTRODUCTION

Several ferroic properties are displayed by multiferroic materials like ferroelectricity, ferroelasticity, and ferromagnetism in a single phase [1]. At room temperature,  $\text{BiFeO}_3$  has ferroelectric Curie temperature ( $T_C = 1103$  K) and G-type antiferromagnetic Neel temperature ( $T_N = 643$  K) [2]. The multiferroic bismuth ferrite ( $\text{BiFeO}_3$ ) have number of important applications such as nanoelectronics [3], memory devices [4], optical filters [5], piezoelectric devices [6] and microelectronic devices [7].

The number of synthesis routes have been used for the preparation of  $\text{BiFeO}_3$  ceramics such as solid state reaction route [8], mechanochemical synthesis technique [9], Sol gel method [10], hydrothermal method [11], citrate gel technique [12], Co-precipitation technique [13] and liquid phase precipitation route [14]. In this article, we have synthesized the  $\text{Bi}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{FeO}_3$ ,  $\text{Bi}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{FeO}_3$ ,  $\text{Bi}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{FeO}_3$  and  $\text{Bi}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{FeO}_3$  multiferroic ceramics via solution combustion method.

## II. EXPERIMENTAL PROCEDURE:

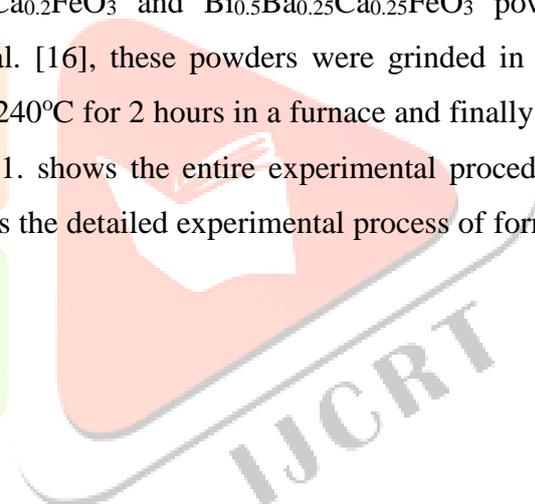
### Initial Materials:

The bismuth nitrate, barium nitrate, calcium nitrate, ferric nitrate and glycine were used as primary starting materials for the formulation of Bi site codoping of Ba and Ca in BiFeO<sub>3</sub>.

### Synthesis Process:

The preparation of Bi<sub>0.8</sub>Ba<sub>0.1</sub>Ca<sub>0.1</sub>FeO<sub>3</sub>, Bi<sub>0.7</sub>Ba<sub>0.15</sub>Ca<sub>0.15</sub>FeO<sub>3</sub>, Bi<sub>0.6</sub>Ba<sub>0.2</sub>Ca<sub>0.2</sub>FeO<sub>3</sub> and Bi<sub>0.5</sub>Ba<sub>0.25</sub>Ca<sub>0.25</sub>FeO<sub>3</sub> ceramic samples were carried out using the precursors such as bismuth nitrate, barium nitrate, calcium nitrate, ferric nitrate as an oxidizers while glycine was used as a fuel. In order to prepare the mixture of samples, the oxidizer (O) to fuel (F) ratio was properly considered using the oxidizing and reducing valences of the metal nitrates and fuel [15].

The bismuth nitrate, barium nitrate, calcium nitrate, ferric nitrate and glycine taken in a stoichiometric quantity and were dissolved in a distilled water in a separate beakers after that, these solution were mixed together and conveyed in a pyrex dish for heating on a gas burner. Afterwards the continuous heating, the water gets evaporated and finally a combustion takes place with formation of Bi<sub>0.8</sub>Ba<sub>0.1</sub>Ca<sub>0.1</sub>FeO<sub>3</sub>, Bi<sub>0.7</sub>Ba<sub>0.15</sub>Ca<sub>0.15</sub>FeO<sub>3</sub>, Bi<sub>0.6</sub>Ba<sub>0.2</sub>Ca<sub>0.2</sub>FeO<sub>3</sub> and Bi<sub>0.5</sub>Ba<sub>0.25</sub>Ca<sub>0.25</sub>FeO<sub>3</sub> powders. The experimental procedure was given by Chaudhari et.al. [16], these powders were grinded in an acetone medium and finally calcined at 210°C, 220°C, 230°C, 240°C for 2 hours in a furnace and finally carried out for pellet formation. The following flowchart in Fig.1. shows the entire experimental procedure for the formulation of these powder samples and Fig 2. depicts the detailed experimental process of formulation of the powder samples.



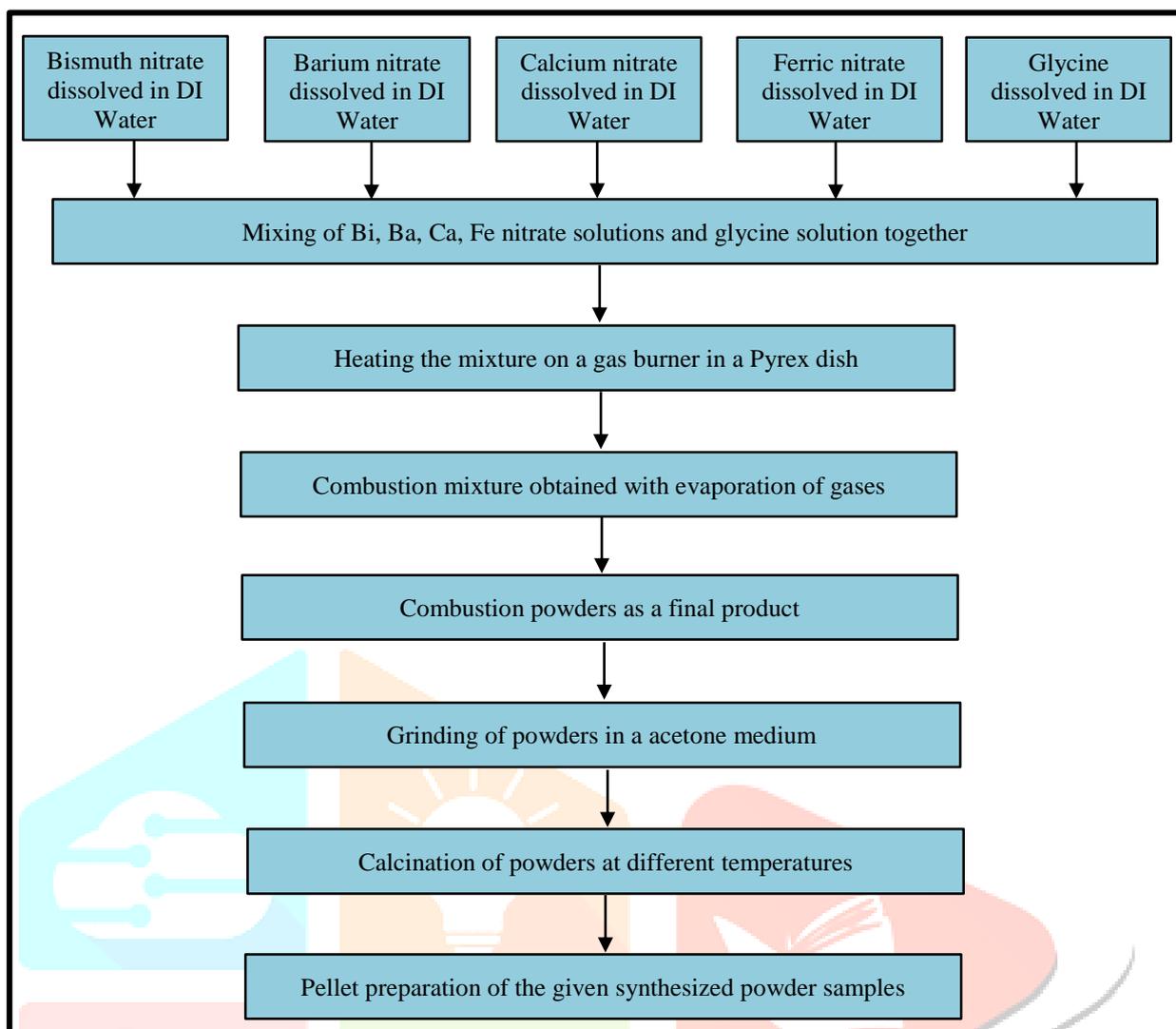


Fig.1. Flowchart of preparation of  $\text{Bi}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{FeO}_3$ ,  $\text{Bi}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{FeO}_3$ ,  $\text{Bi}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{FeO}_3$  and  $\text{Bi}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{FeO}_3$  multiferroic samples by SCM.

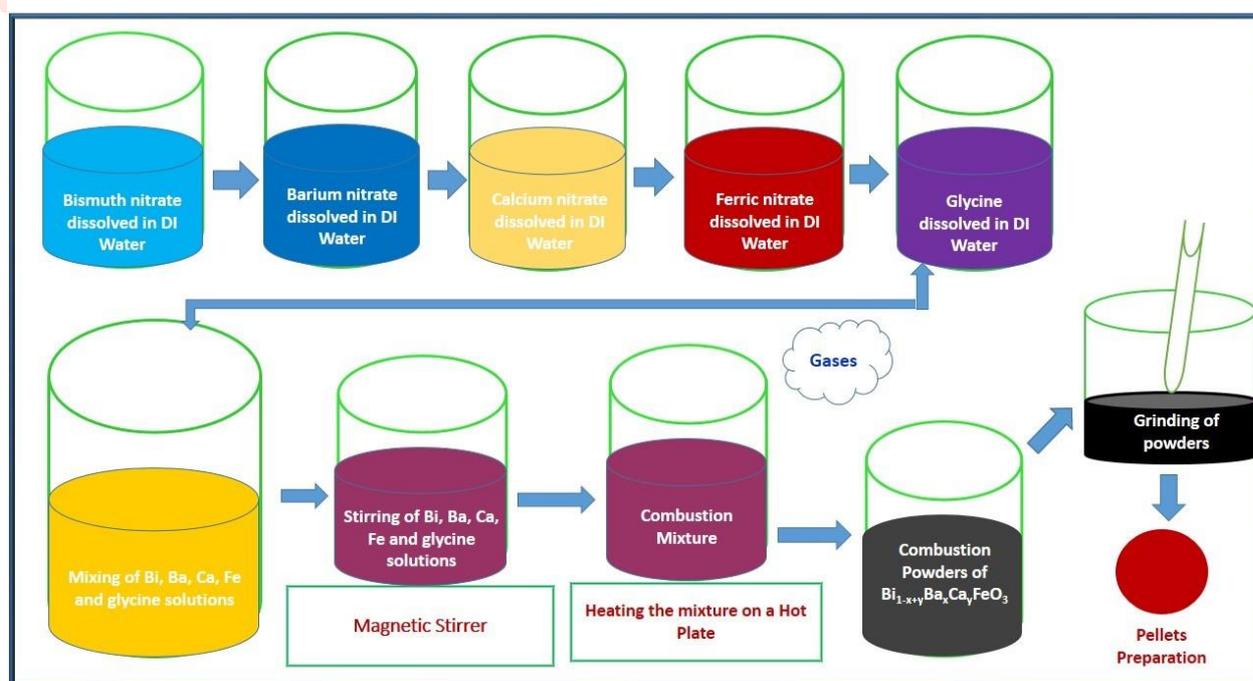


Fig.2. Experimental process of formulation of  $\text{Bi}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{FeO}_3$ ,  $\text{Bi}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{FeO}_3$ ,  $\text{Bi}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{FeO}_3$  and  $\text{Bi}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{FeO}_3$  powder samples.

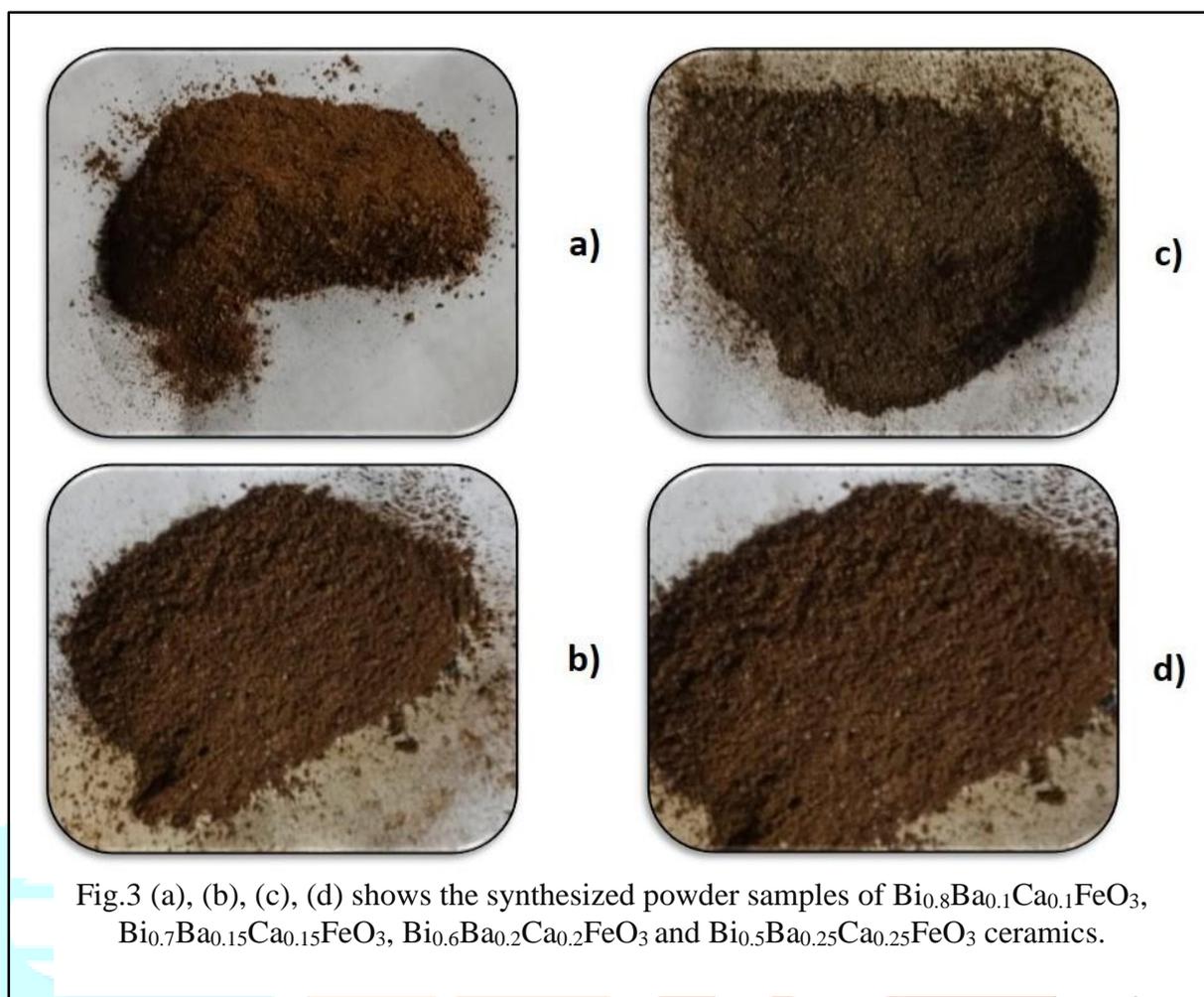


Fig.3 (a), (b), (c), (d) shows the synthesized powder samples of  $\text{Bi}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{FeO}_3$ ,  $\text{Bi}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{FeO}_3$ ,  $\text{Bi}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{FeO}_3$  and  $\text{Bi}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{FeO}_3$  ceramics.

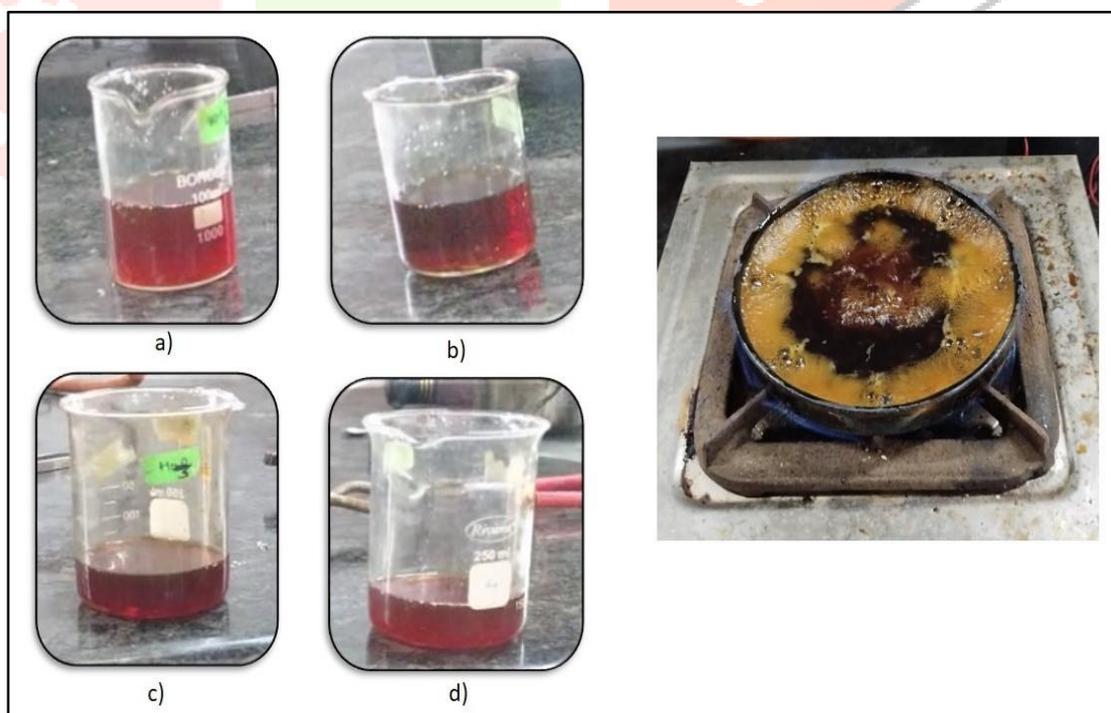
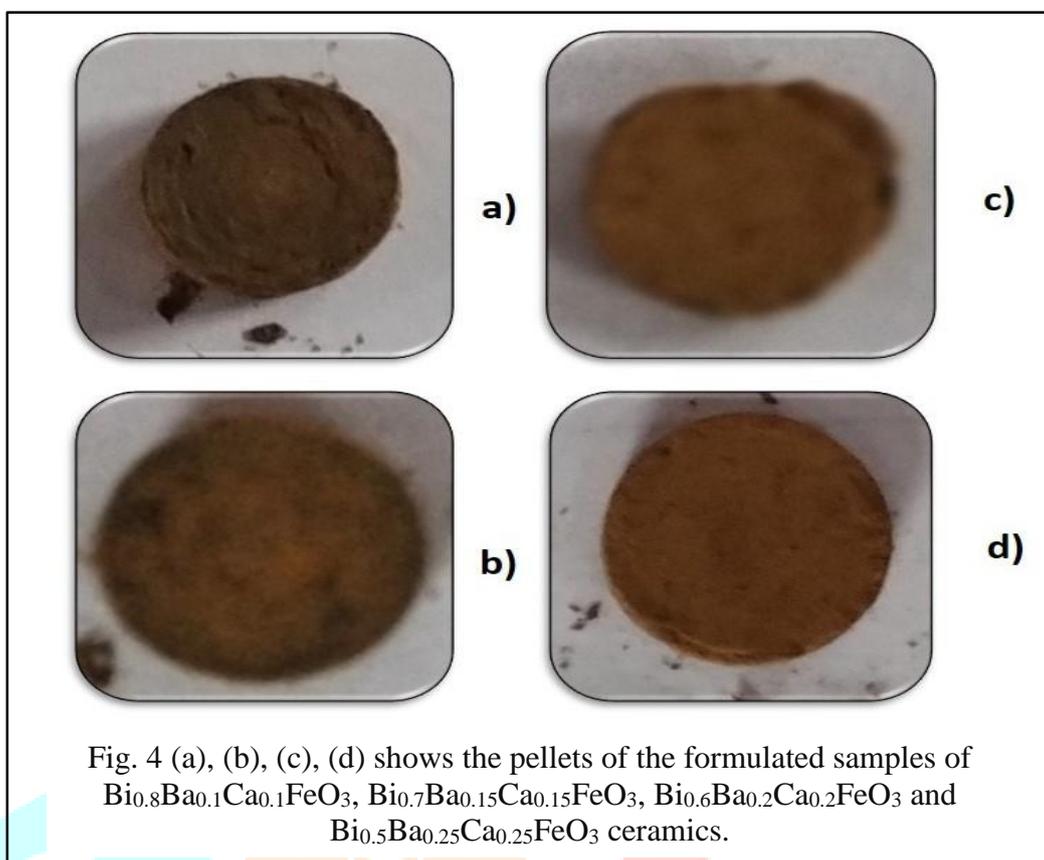


Fig.5 (a), (b), (c), (d) shows the prepared solutions of  $\text{Bi}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{FeO}_3$ ,  $\text{Bi}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{FeO}_3$ ,  $\text{Bi}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{FeO}_3$  and  $\text{Bi}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{FeO}_3$  samples and a combustion reaction on a gas burner.



### III. Results and discussion:

Fig.1. shows the flowchart of sample preparation of  $\text{Bi}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{FeO}_3$ ,  $\text{Bi}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{FeO}_3$ ,  $\text{Bi}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{FeO}_3$  and  $\text{Bi}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{FeO}_3$  materials in powder form and Fig 2. illustrates the full experimental process of formulation of the powder samples.

Fig.3 (a), (b), (c), (d) shows the synthesized  $\text{Bi}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{FeO}_3$ ,  $\text{Bi}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{FeO}_3$ ,  $\text{Bi}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{FeO}_3$  and  $\text{Bi}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{FeO}_3$  nanopowder samples. Fig. 4 (a), (b), (c), (d) shows the pellets of the prepared  $\text{Bi}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{FeO}_3$ ,  $\text{Bi}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{FeO}_3$ ,  $\text{Bi}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{FeO}_3$  and  $\text{Bi}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{FeO}_3$  samples. These formulated powder samples were grinded in an acetone medium, calcined carried out at temperatures of 210°C, 220°C, 230°C, 240°C for 2 hours in a furnace. Fig. 5 (a), (b), (c), (d) shows the prepared solutions of  $\text{Bi}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{FeO}_3$ ,  $\text{Bi}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{FeO}_3$ ,  $\text{Bi}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{FeO}_3$  and  $\text{Bi}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{FeO}_3$  samples and a combustion reaction on a gas burner.

### IV. Conclusion:

In the present article, we have effectively produced the multiferroic  $\text{Bi}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{FeO}_3$ ,  $\text{Bi}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{FeO}_3$ ,  $\text{Bi}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{FeO}_3$  and  $\text{Bi}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{FeO}_3$  nanopowder samples using solution combustion method (SCM).

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