



Solution Combustion Method (SCM) for the Formulation of $\text{BiFe}_{1-x+y}\text{Ba}_x\text{Ca}_y\text{O}_3$ ($x, y = 0.1, 0.15, 0.2, 0.25$) Nanopowder Samples

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Abstract: This paper presents the formulation of nanopowders of multiferroic samples like $\text{BiFe}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{O}_3$, $\text{BiFe}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{O}_3$, $\text{BiFe}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{O}_3$ and $\text{BiFe}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{O}_3$ by solution combustion method (SCM). The nanopowders of $\text{BiFe}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{O}_3$, $\text{BiFe}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{O}_3$, $\text{BiFe}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{O}_3$ and $\text{BiFe}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{O}_3$ samples were prepared using metal nitrates and glycine fuel as initial starting materials. These nanopowder samples were processed by grinding them in an acetone medium, calcining them at various temperatures, and finally producing pellets.

Keywords: Multiferroics, Nanopowder, BiFeO_3 , Ca and Ba dopants, SCM, applications.

I. INTRODUCTION

Ferroelectricity and ferromagnetism coexisting and interacting simultaneously in multiferroic materials [1]. At room temperature, BiFeO_3 manifests a ferroelectric Curie temperature $T_C = 1103 \text{ K}$ and G-type antiferromagnetic Neel temperature $T_N = 643 \text{ K}$ [2].

The multiferroic BiFeO_3 have number of key applications in spintronics [3], photovoltaics, optical filters [4], sensors [5], catalytic [6], piezoelectric devices, photosensitizers [7], multistate storage [8] and high density microactuators [9]. The number of formulation routes have been used for the synthesis of BiFeO_3 multiferroic ceramics like solid state reaction technique [10], combustion technique [11], auto-combustion method [12] and sol-gel auto combustion method [13].

In this article, we have synthesized the $\text{BiFe}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{O}_3$, $\text{BiFe}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{O}_3$, $\text{BiFe}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{O}_3$ and $\text{BiFe}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{O}_3$ multiferroic nanopowder samples through solution combustion method.

II. Experimental Procedure:

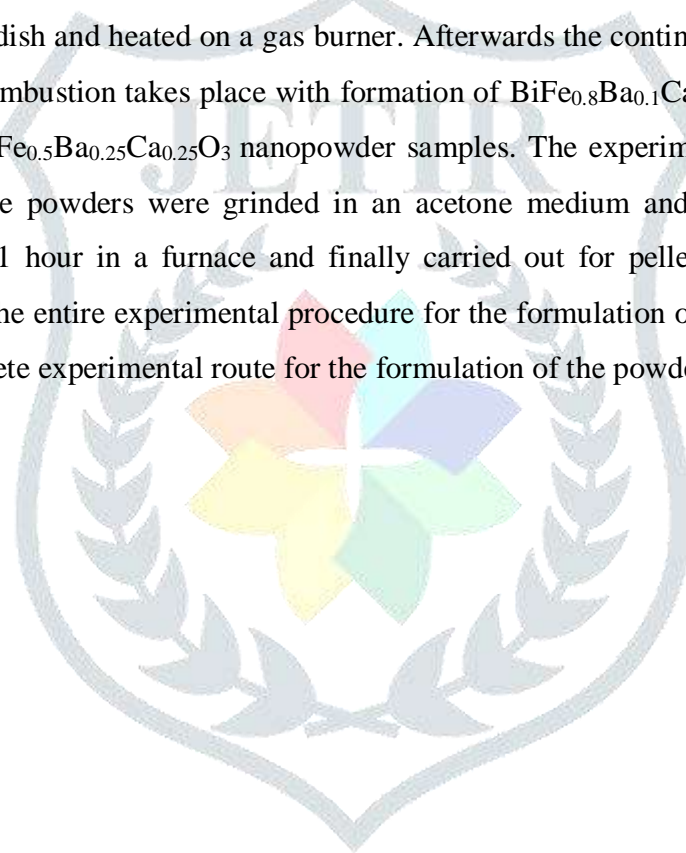
Initial Materials:

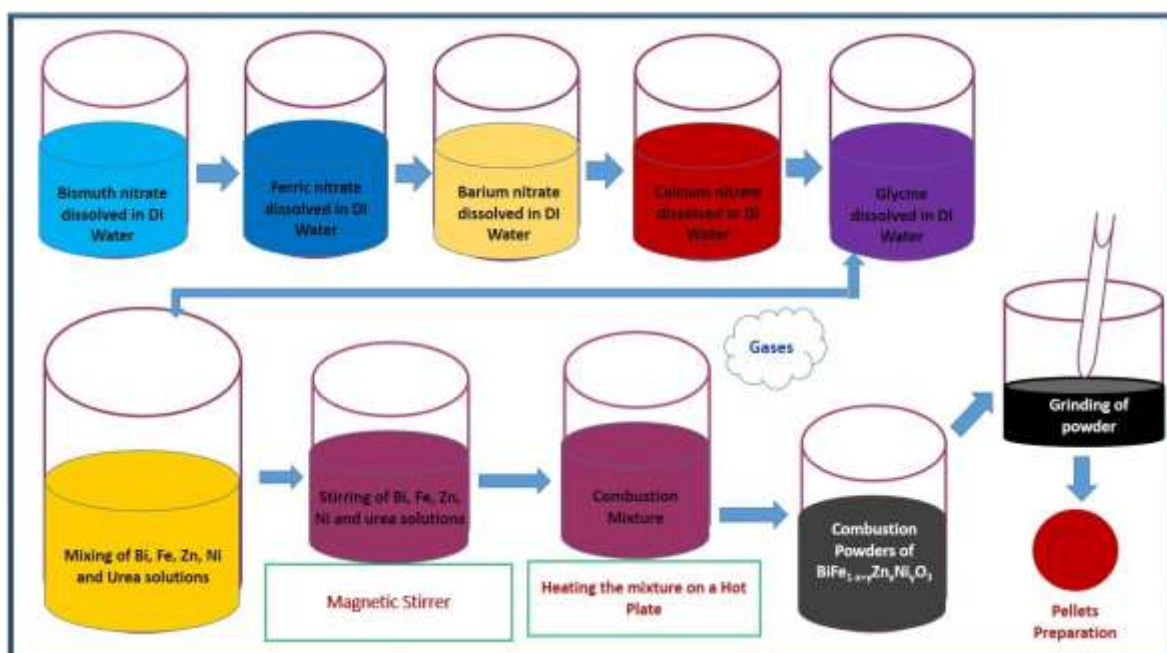
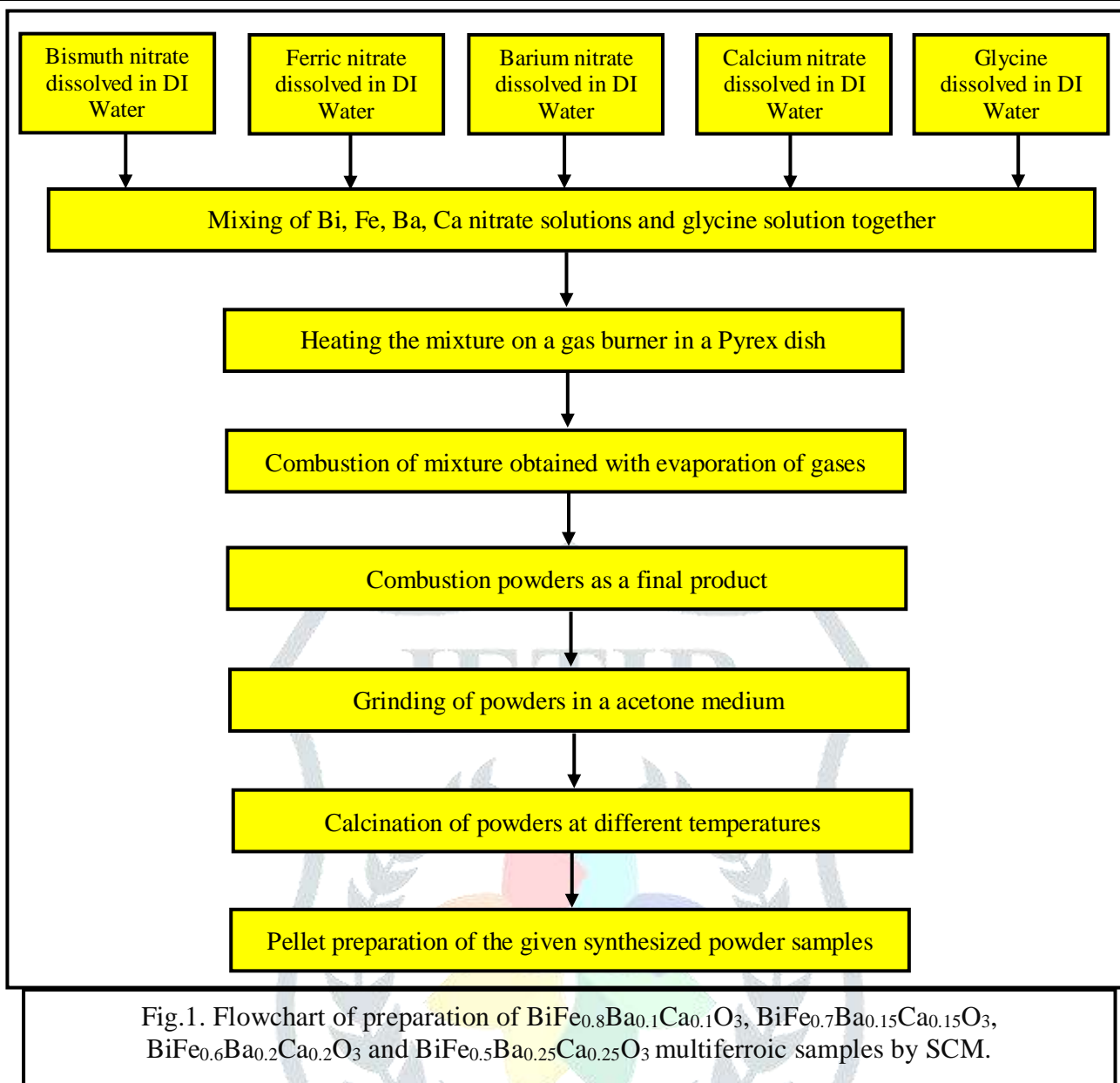
The bismuth nitrate, ferric nitrate, barium nitrate, calcium nitrate and glycine were used as initial starting materials.

Synthesis Process:

The preparation of $\text{BiFe}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{O}_3$, $\text{BiFe}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{O}_3$, $\text{BiFe}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{O}_3$ and $\text{BiFe}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{O}_3$ ceramic samples were carried out using the precursors such as bismuth nitrate, ferric nitrate, barium nitrate, calcium nitrate as an oxidizers while glycine was used as a fuel. The oxidizer (O) to fuel (F) ratio was measured accurately to form the sample mixture by measuring the oxidizing and reducing valences of the fuel [14].

The bismuth nitrate, ferric nitrate, barium nitrate, calcium nitrate and glycine taken in a stoichiometric extent and were dissolved in a distilled water in a separate beakers thereafter, The mixture of these solutions was transferred to a Pyrex dish and heated on a gas burner. Afterwards the continuous heating, the water gets evaporated and finally a combustion takes place with formation of $\text{BiFe}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{O}_3$, $\text{BiFe}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{O}_3$, $\text{BiFe}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{O}_3$ and $\text{BiFe}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{O}_3$ nanopowder samples. The experimental process was given by Chaudhari et.al. [15], these powders were grinded in an acetone medium and finally calcined at 200°C , 210°C , 220°C , 230°C for 1 hour 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. describes the complete experimental route for the formulation of the powder samples.





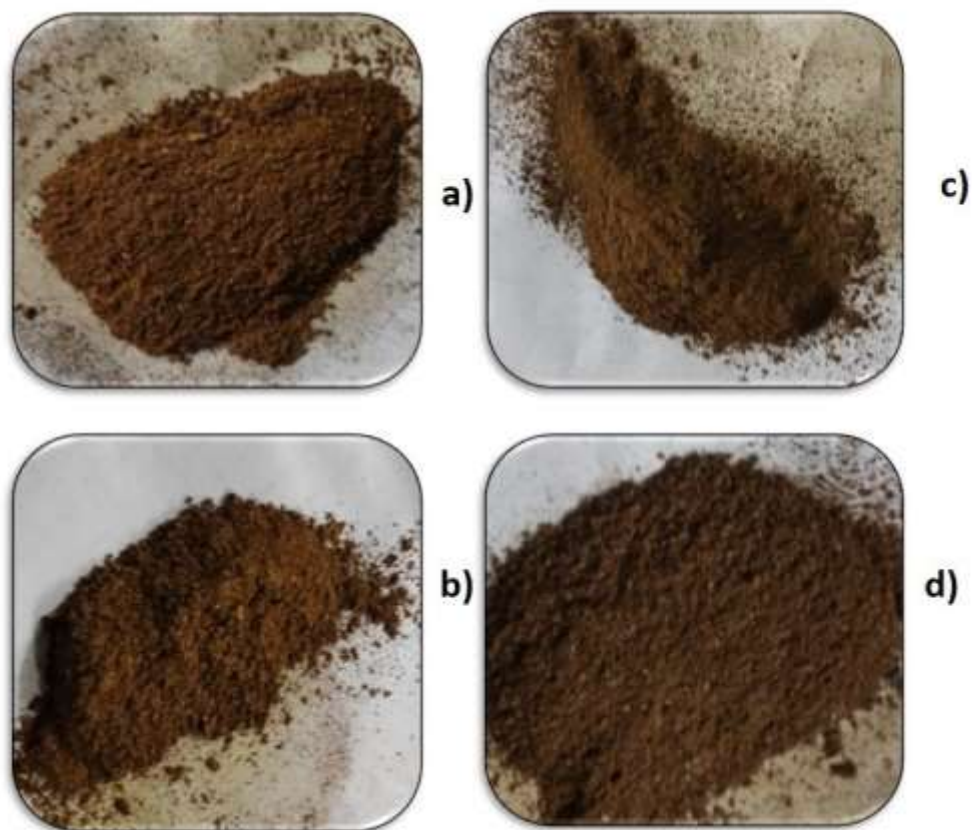


Fig.3 (a), (b), (c), (d) shows the synthesized powder samples of $\text{BiFe}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{O}_3$, $\text{BiFe}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{O}_3$, $\text{BiFe}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{O}_3$ and $\text{BiFe}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{O}_3$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

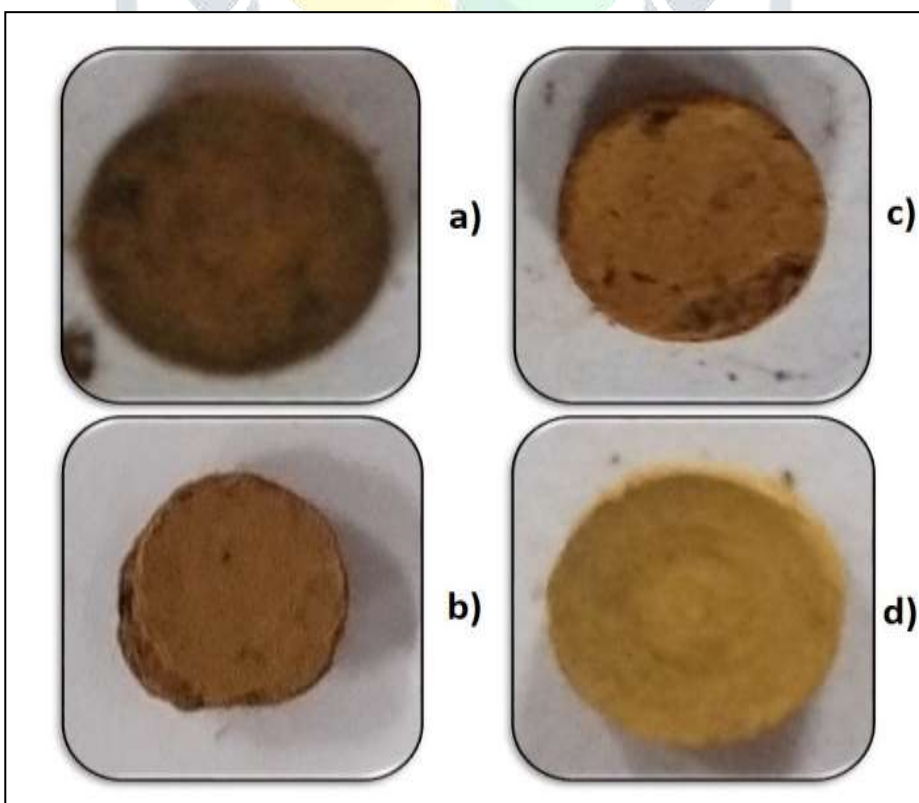


Fig. 4 (a), (b), (c), (d) shows the pellets of the formulated samples of $\text{BiFe}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{O}_3$, $\text{BiFe}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{O}_3$, $\text{BiFe}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{O}_3$ and $\text{BiFe}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{O}_3$ ceramics by SCM.

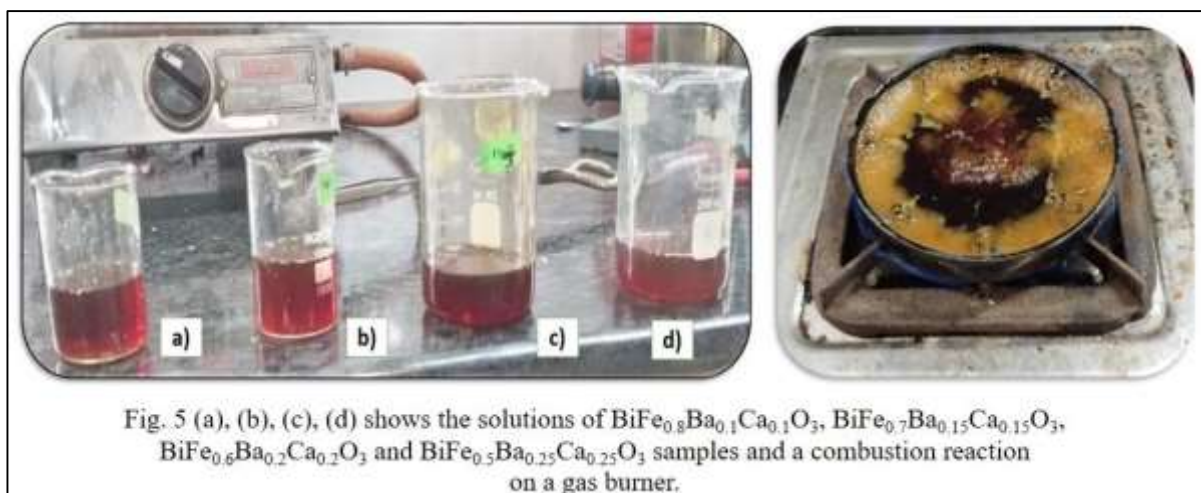


Fig. 5 (a), (b), (c), (d) shows the solutions of $\text{BiFe}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{O}_3$, $\text{BiFe}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{O}_3$, $\text{BiFe}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{O}_3$ and $\text{BiFe}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{O}_3$ samples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

III. RESULTS AND DISCUSSION:

Fig.1. shows the flowchart of sample preparation of $\text{BiFe}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{O}_3$, $\text{BiFe}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{O}_3$, $\text{BiFe}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{O}_3$ and $\text{BiFe}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{O}_3$ materials and Fig.2. presents the whole experimental scheme of synthesis of the nanopowder samples. Fig.3 (a), (b), (c), (d) shows the synthesized $\text{BiFe}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{O}_3$, $\text{BiFe}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{O}_3$, $\text{BiFe}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{O}_3$ and $\text{BiFe}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{O}_3$ powder samples. Fig. 4 (a), (b), (c), (d) shows the pellets of the prepared $\text{BiFe}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{O}_3$, $\text{BiFe}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{O}_3$, $\text{BiFe}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{O}_3$ and $\text{BiFe}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{O}_3$ samples. These formulated powder samples were grinded in an acetone medium, calcined carried out at temperatures of 200°C , 210°C , 220°C , 230°C for 1 hour in a furnace. Fig. 5 (a), (b), (c), (d) shows $\text{BiFe}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{O}_3$, $\text{BiFe}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{O}_3$, $\text{BiFe}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{O}_3$ and $\text{BiFe}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{O}_3$ samples solutions and the combustion reaction carried out on a gas burner.

IV. Conclusion:

In the present article, we have effectively synthesized the multiferroic $\text{BiFe}_{0.8}\text{Ba}_{0.1}\text{Ca}_{0.1}\text{O}_3$, $\text{BiFe}_{0.7}\text{Ba}_{0.15}\text{Ca}_{0.15}\text{O}_3$, $\text{BiFe}_{0.6}\text{Ba}_{0.2}\text{Ca}_{0.2}\text{O}_3$ and $\text{BiFe}_{0.5}\text{Ba}_{0.25}\text{Ca}_{0.25}\text{O}_3$ powder samples and finally pellets formation. At different temperatures, these powder samples were calcined.

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