Flame Spray Pyrolysis Processing to Produce Metastable Phases of Metal Oxides

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Abstract

Functional metal oxides can exist as different polymorphs. Meta stable phases of these oxides may be produced by flame spray pyrolysis (FSP). This is a scalable, rapid solidification process that may be used for nano powder synthesis. This review describes the key features of the FSP process and equipment. It also provides examples of the use of FSP process to obtain a meta stable oxide phases at RT which have unique properties.

Keywords: Flame spray pyrolysis; Ceramics; Polymorphism; Metastability

Introduction

When bulk metal oxides are subject to differing conditions of pressure and temperature, they crystallize in Metastable forms that are structurally different [1]. However, the same Metastable components can be formed in nanoscale at normal conditions of temperature and pressure. The formation of metastable compounds has been related to Gibb’s free energy; a material, during nucleation, will exhibit a polymorph that has lowest free energy at a particular temperature and pressure. At a given thermodynamic condition, both Metastable and stable phases may coexist, but only one of the two phases is stable, with minimal free energy, and the other must be metastable and may transform into a stable state [1]. The free energy of a polymorph is governed by size, temperature and pressure. A slight change in either one of the condition may affect the formation of a polymorph.

To achieve a metastable form at room temperature, rapid solidification/quenching of these materials [2,3]. A number of studies focused on the formation of metastable complexes at room temperature by obtaining nanocrystalline solids by quenching from high temperatures. Among the various methods used, flame spray pyrolysis is of prime interest as the process offers high temperature gradient and low residence time (start of nucleation-quenching) [4]. Thus, generating favorable conditions towards formation of metastable phases. The process and its uses are described below.

The FSP process

Table 1: FSP parameters that affect the morphology and structure of resulting powders [7].

<table>
<thead>
<tr>
<th>Feature</th>
<th>Controlling Parameters</th>
</tr>
</thead>
<tbody>
<tr>
<td>Crystal size</td>
<td>Precursor composition, fuel and oxidant flow rate, burner configuration, size of droplet and flame</td>
</tr>
<tr>
<td>Crystal structure</td>
<td>Precursor flow rate and fuel/oxidant flow rate</td>
</tr>
<tr>
<td>Particle Morphology</td>
<td>Precursor concentration and fuel/oxidant flow rate</td>
</tr>
</tbody>
</table>

The earliest definition of the flame spray pyrolysis process has been: “pyrolyzing a suitable vapor of precursors that when decomposed give oxides with interesting catalytic properties”[5]. Flame spray pyrolysis is described as a break down process and it involves delivering a droplet of the precursor into the reactor by using an atomizer/ nebulizer. Since the reactor is at high temperature, the solvent evaporates and the remaining solute reacts with the surrounding gases. The composition of the final product is determined by the stoichiometric ratio of the solute dissolved in the precursor. The size and velocity of the droplets as well as the concentration of the precursor determine the particle size distribution of the powders deposited. Finally, the morphology of the nano particles is also affected by the surrounding gas flow rate, the temperature, and the type of
Flame-spray pyrolysis, while enabling kinetic control of metastable polymorphs, is also a very effective and scalable industrial process to synthesize oxide nano particles in large quantities while maintaining an excellent product quality. It has been used for the dry, one-step synthesis of catalysts, sensors, biomaterials, phosphors, and even nutritional supplements, each of which would require high quality metastable materials to ensure the desired functionality and provides precedent for FSP synthesized material quality. Flame spray pyrolysis has high processing temperature (as high as 2800K) and the highest gas flow rate of most oxide processing techniques. Based on these two factors, flame spray pyrolysis is the only processing method that can provide very short residence times (~ms time scale) in the highest temperature gradient region and is thus a powerful synthesis method for production of metastable species through kinetic control [8]. Various flame configurations are used for the manufacturing of nano particles, such as premixed and diffusion flames run in co-flow, or counter-flow conditions. In the diffusion flame configuration, the fuel and the oxidizer diffuse into each other inducing flame reaction and particle formation, while in premixed flames the precursor and the combustible gases are mixed before they enter the reaction zone (flame).

Types of FSP

Flame spray pyrolysis has been categorized into three different types:

Vapor assisted flame spray pyrolysis (VAFS): A commonly used industrial process for the synthesis of ceramics, such as, fumed silica, alumina and pigmentary titania [9]. The process uses volatile metal precursors, which are evaporated and then fed into a flame. The metal particles are converted into oxides and particle formation occurs by nucleation from gas phase. The major drawback of the process is its need for volatile precursors, which limits the variety of products that can be generated through the process.

Flame aerosol spray pyrolysis (FASP): It uses non-combustible liquid metal precursors (such as aqueous solutions of metallic salts). The precursor liquid is dispersed into tiny droplets and pyrolyzed by an external flame [9]. The flame is usually produced by hydrocarbons or \( \text{H}_2/\text{O}_2 \). Upon evaporation, the particle formation process is similar to VAFS. Liquid feed flame spray pyrolysis (LF-FSP):

This was developed by R.M Laine at University of Michigan, Ann Arbor [10]. The liquid fed FSP was able to overcome the drawback of FASP by fabricating more complex metal oxide particles. The process uses a metal oxide precursor in the form of liquid; its composition is generally in the form of metal chlorides, metal nitrates or metal acetates dissolved in aqueous or alcohol based solutions [11] (Figure 1). The process features short residence times (a few milliseconds) and results in formation of homogenous, nano sized particles [12].

Components of the FSP

Atomizer: The purpose of Atomizer is to create droplets. Three different kinds of atomizers [13] exist and are described below:

Ultrasound nebulizer: Used to form sprays using high-frequency vibrations and converting liquid into mist. The mist enters the flame and follows the VAFS route, giving small particles.

Two-fluid nozzle: A device that disperses liquid into spray, the breakup of liquid into spray occurs with the help of an atomizing gas.

Electro-sprayer: It uses high voltage to convert liquid into aerosol. It is useful for producing ions from macromolecules.

A two-fluid nozzle is the most commonly used atomizer for FSP system.

Burner: The purpose of burner is to generate a flame. Two different types of burner exist, which are a premixed burner and a diffusion burner: The premixed burner mixes the fuel and oxidant [13]. FSP generates a self-sustaining flame so that the burner system for it is an ignition source. The advantage for it is that it could directly control the particle size by controlling the combustion enthalpy and the metal concentration of the precursor [4].

Collector: The purpose of collector is to collect the final product from FSP process. Usually a substrate or a filter is used for collection of particles.

Liquid precursor selection

The selection of liquid precursor is an important step in FSP particle synthesis. The selection of metallic precursor and solvent should be performed with keeping various factors in mind, such as, melting/decomposition temperatures, miscibility and chemical stability. These factors govern the overall particle formation in flame. Nitrates, acetates and acetyl acetonates are the natural choices for FSP as they were economical and readily available. However, they do not always yield homogeneous morphology [4].
The low combustion enthalpies of precursors coupled with high melting points results in formation of in-homogenous particles [14]. However, this drawback can sometimes be remediated by processing parameters. Following is how the particle formation is affected by liquid precursor formulation.

Hollow particles are formed when the boiling point of solvent is less than the melting point of solute or if the metal precursor precipitates on the surface of solvent. These parameters can be overcome by providing sufficient heat to the precursor. For formation of homogenous particles the melting point of precursor solute should be lower than the boiling point of solvent.

**Mechanism for particle formation in FSP**

**One droplet to one particle (ODOP):** As the name suggests, the formation of particles through this mechanism is by conversion of a single droplet into a single particle. The first step involves solvent evaporation and allows solute to disperse in the reactor. The solute then reacts with carrier gas or one-another to form particles. In final step, the particles undergo sintering and densification. The resultant of ODOP mechanism, yields particles in the micron size range. A process usually follows ODOP when the melting point of solute is higher than solvent [4].

The particle size for ODOP [13] can be predicted and is given by:

$$dp=\left(\frac{C\cdot Md^3}{N\cdot \rho}\right)^{1/3}$$

Where,

- **Dp** is the average particle size
- **C** is the concentration of the droplet
- **D** is the diameter of the droplet
- **M** is the molecular weight of particles
- **N** is the stoichiometric ratio
- **ρ** is the particle density

**One droplet to multiple particles (ODMP):** The steps involved in this mechanism are similar to that of ODOP. The difference between the two mechanisms is based on the precursor. Precursors tend to follow ODMP if there is presence of an organic additive. The additive leads to crack down of particles during the sintering process, which results in formation of multiple particles. The particles obtained exhibit a wide size range and are usually nano-crystalline [13].

**Gas phase mechanism:** For this mechanism, the precursor is an organo metallic compound. The precursor evaporates quickly once it enters the flame, producing metal vapors. The formations of particles occur in various steps [4] and are given as a flow chart in (Figure 2) below: In the gas phase mechanism, the metal vapor may collide with the carrier gas to nucleate thereby forming clusters. Alternatively, the vapor may also nucleate on the surface of formed clusters resulting in new particles growth. The higher the number of clusters is being formed, the higher is the probability for them to collide with each other and form strong adhesive (physical) bonds or chemical bonds. The formations of bonds result in coagulation to produce nano particles. Finally, these nano particles produce single particles during coalescence process. The coalescence process results in aggregation of particles in the high temperature zone. To control the particle size by not allowing them to aggregate into larger particles, the lower residence time of precursor is needed [13]. The lower residence time can be achieved by a higher flow-rate of the precursor solution or high organic content of the organic precursor solvent, which results in bigger flame height.

**Figure 2:** Formation of particles through gas phase mechanism.

**Metastable phase formation**

Flame spray pyrolysis has resulted in the formation of many metastable phases for ceramic oxides. The high temperature of operation coupled with rapid solidification and small particle sizes offer perfect conditions for formation of metastable polymorphs [4]. The size of particles formed through FSP are affected by various parameters such as composition of precursor, fuel and dispersion/oxidant flow rate, size of droplet and temperature of flame [15]. These parameters are interrelated as a high oxidant flow gives rise to low flame temperature. A non-combustible solute lowers the flame temperature, and lower oxidant flow rate decreases the droplet atomization and yields larger droplet. An optimum particle size and phase distribution, as well as tailored polymorph selection can be obtained from the same precursor by properly adjusting the processing parameters. For example, a metastable tetragonal-BaTiO$_3$ was produced using FSP [16], from organic precursors of Ba and Ti dissolved in HNO$_3$ and citric acid. Pure phases with low particle sizes were achieved for the precursor with the highest concentration of citric acid. This finding was associated with lower residence time of the vapors. In another study by a different group, the same recipe was used except citric acid was replaced by water [17]. The addition of water to the precursor mixture resulted in lower flame temperatures, bigger particle sizes, and the formation of a cubic-BaTiO$_3$ phase. The lower flame temperature due to the addition of water increased the residence time for the particles which in turn led to formation of
larger particles and stabilization of the cubic polymorph. Organic precursors (organic solute in organic solvent) are the best choice for the formation of metastable polymorphs as they offer a high temperature gradient and low residence time.

Tungsten trioxide is a well-known metal oxide that finds widespread use in gas sensing, electrochromic, and catalytic applications [18-20]. Polymorphic transformations of the pseudo-cubic lattice result in a wide variation in its electronic properties. WO$_3$ usually crystallizes in one of the following lattice configurations: triclinic, monoclinic, orthorhombic, and tetragonal (Figure 3). All of these polymorphs are distorted forms of a cubic ReO$_3$ lattice, with increasing order of crystallographic symmetry from the triclinic to tetragonal lattice. The crystal lattice is composed of a framework of metal-oxygen octahedral units, where the metal atoms are located at the center of the oxygen octahedron with varying amounts of metal-oxygen bond lengths and thus varying amounts of octahedral distortion. This distortion in turn serves to stabilize the different polymorphs. (Figure 4) demonstrates that high quality ε-WO$_3$ nano powders have been synthesized using FSP by our group. The ε-phase for WO$_3$ is very important for gas sensing applications as it acts as a selective sensor for acetone detection [19].

![Figure 3: Phase stability fields for WO$_3$ polymorphs in the bulk material (from ref. [17]).](image_url)

![Figure 4(a & b): XRD patterns of WO$_3$ nanocrystals prepared by the FSP process that show a "rare" polymorphic structure. The main peaks can be assigned to monoclinic ε-WO$_3$ phase (JCPDS No.: 87-2386). (b) HRTEM micrograph of the as-synthesized WO$_3$ nano crystals.](image_url)

**Conclusion**

The operating principle and key features of the flame spray pyrolysis process have been reviewed in this work. It has been shown that this atomization process may be fine-tuned to guide the selection of particular polymorphs of metal oxides as the final product. Material precursor parameters, such as the solvent used, play a key role to determining the final particle size and the phase of the powders produced.

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