Thermal studies of fly ashes expansion

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Abstract

The object of this study is to investigate the thermal properties of fly ashes from the last, farthest dedusting zone in terms of their use as ceramic masses additives. Siliceous fly ash is valuable additive to ceramic mass, which not only reduces its plasticity, but also actively affects sintering process and shapes the properties of the final material. The finest fly ash fractions are potentially useful flux materials in ceramics; however, a significant limitation in their use is due to thermal expansion/bloating occurring during high-temperature sintering. The bloating mechanism of fly ashes was investigated in relationship to their chemical composition with the use of DTA/TG/EGA analysis as well as high-temperature microscope. Chemical and phase compositions were studied by X-ray fluorescence and X-ray diffraction. Based on the results obtained, it can be concluded. The results indicate that bloating mechanism is caused by the co-occurrence of two phenomena accompanying sintering: appearance of high amount of liquid phase and simultaneous gas release from sintered material. The dominant mechanism is the simultaneous release of sulfur (IV) oxide and oxygen as a result of the redox reaction of removing SO3 from the vitreous phase.

Keywords Fly ash · Expansion · Bloating · Thermal studies

Introduction

Siliceous fly ash is a waste material created during combustion of coal in pulverized coal boilers, accumulated in the dust removal zone [1]. It is used in the building materials industry as a raw material for the production of Portland clinker [2], as an addition to cement [3, 4], mortar [5] and concrete [6–8], as well as sintered ceramic products [9, 10]. Fly ash was also researched as a starting material for the production of high-degree-of-consolidation and low-porosity ceramics [11, 12]. This type of material is sintered with a large proportion of the liquid phase at well-defined temperature intervals to avoid thermal softening and/or bloating.

Bloating phenomenon is the result of an intense gas emission from the material during sintering with a large proportion of the liquid phase [13, 14]. It is also a phenomenon that allows for the production of expanded clay aggregate [15, 16]. Part of the emitted gases (CO2) is the result of oxidation of unburned carbonaceous materials and thermal decomposition of carbonates. Others are the effect of redox reactions (reduction of iron oxides, but also sulfates and sulfides), and some are the result of reactions between clay components and admixtures [17–20]. The gases formed during the reaction cannot escape from the sintered material, which causes bubble creation; however, sufficiently slow heating of the material allows to complete these reactions before sintering and closing the pores [21].

In the case of sintered materials, fly ash is a partial substitute for clay minerals. In addition, due to the increased presence of alkali oxides, they act as a flux that accelerates the sintering process. The presence of alkali generates the formation of a liquid phase during firing, which accelerates mass transport and intensifies sintering. The addition of fly ash to the ceramic mixture allows to lower the temperature of the sintering process [22, 23]. The fine-grained fly ashes seem to be particularly interesting, because the content of valuable alkali is higher in the smaller fractions and at the same time their surface development is higher, which should be treated as the driving force of the sintering process. Unfortunately, the addition of fine-grained fly ash causes intense bloating of the sintered material, which reduces the safe temperature interval of the process.
In the case of sintered ceramics, there is also a phenomenon of “black core” associated with expansion, which is caused by the presence of organic compounds as well as reducing oxides [24]. This phenomenon is also found in sintered fly ashes and ceramic materials with the addition of fly ash [22, 23].

The aim of this work is to examine the phenomenon of bloating/expansion of fine fly ashes during high-temperature treatment. The performed research focuses mainly on differential thermal analysis and thermogravimetry combined with gas analysis in various measurement atmospheres.

**Method of examination**

The subject of the study was two fly ashes (X and Z) taken from two separate power plants based on conventional coal dust furnaces. Dust extraction was carried out with the help of several electrostatic precipitator zones arranged in series behind the combustion chamber. For the purpose of the experiment, only fly ashes from the last, farthest dedusting zone were taken, because these ashes are the most thermally swellable. It is worth noticing that they are the finest and the most interesting fly ashes for ceramic mixtures, in relation to ashes taken from other dust removal zones. Chemical and phase composition of fly ashes were studied with the use of X-ray fluorescence and XRD methods. The chemical analysis of fly ashes was carried out with WD XRF fluorescence spectrometer (Axios mAX Company). Powder diffraction measurements were carried out with the use of PANalytical X’Pert Pro MD diffractometer equipped with Cu K$_{\alpha1}$ X-ray line with the Bragg–Brentano standard setup comprising a Ge (111) Johansson monochromator at the incidence beam. The scanning range was 10°–70° with 0.008 step size and 20 s measurement time for each step. A quantitative analysis was carried out using the Rietveld method as described by Le Saout et al. [25]. In order to estimate the amorphous phase, XRD/Rietveld measurement was carried out with corundum as an internal standard. Knowing the percentage of corundum in the sample and the result of Rietveld analysis, the proportion of the glassy phase was calculated. The estimated amount of glassy phase contains crystalline components that are below their detection threshold.

The phenomenon of bloating of fired fly ashes was tested with heating microscope (Hesse Instruments Company). Fly ashes powders were shaped in small cylinders with a height of 3 mm and a diameter of 3 mm, put inside a small furnace and tested separately. The measurement was carried out in air atmosphere at a constant heating rate of 10 °C min$^{-1}$ up to temperature of 1303 °C.

The characteristic temperatures were determined by the samples size and shape change.

The software supplied with the instrument provides sintering curves from which four temperatures can be derived, namely:

1. The first shrinkage or sintering (BS)
2. The maximum shrinkage before softening starts (MS)
3. The softening point, at which the first signal of melting appears (SP)
4. The maximum of bloating—temperature at which the greatest bloating is observed (MB).

Temperatures correspond to a characteristic viscosity point, conventionally determined on the basis of geometrical considerations.

The bloating mechanism was studied with the use of DTA/TG method (thermal differential analysis coupled with thermogravimetry) and the evolved gas analysis EGA.

DTA/TG was performed with the use of STA 449F3 Jupiter Netzsch apparatus in the temperature range between 20 and 1120 °C at a heating rate of 15 °C min$^{-1}$. Average sample masses were between 85 and 95 mg, and the Al$_2$O$_3$-based open crucibles were used. Gas analysis was performed for m/z equal to 18, which corresponds to water secretion (Netzsch STA 449 with mass spectrometer Netzsch QMS 403). Air and helium were used as a carrier gas at a flow rate of 40 mL min$^{-1}$. The apparatus was calibrated in accordance with methodology presented in [26, 27]. The TG/QMS data were analyzed using Proteus software (version 5.2.0) from Netzsch. All the presented curves were corrected against empty runs.

**Results**

**Chemical characteristic**

The chemical composition of both fly ashes is presented in Table 1.

Fly ashes from both power plants have similar chemical composition, which consist mainly of silicon and aluminum oxides (a total of about 70%). The remaining part consists of iron oxide, calcium oxide, magnesium oxide and alkaline oxides (potassium and sodium). Fly ash X contains larger amount of two main oxides in comparison with ash Z and a smaller amount of other oxides. It is worth noticing that fly ash Z contains more than 45% iron oxide, more than 70% sodium oxide and greater amount of sulfur oxide (Table 1).

In Table 2 and Fig. 1, phase composition of fly ashes is presented.

Both fly ashes consist mainly of glassy phase, which may constitute more than 80% of the powder mass, according to the used method. The crystalline components of both ashes consist mainly of mullite and quartz but also hematite.
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Thermal behavior: heating microscope

In Fig. 2, the area changes of fly ash samples as a function of temperature are shown. The inset shows the heating microscope images registered for both fly ashes tested X and with their characteristic temperatures, which are: BS—the beginning of sintering; MS—the maximum shrinking before softening starts; SP—the softening point; MB—the maximum of bloating; and END—the end of measurement (Table 3).

Both analyzed fly ashes undergo thermal bloating. This process begins at a temperature about 1150 °C, but it is effective only after exceeding the 1200 °C. A greater degree of thermal bloating was recorded for fly ash X (230%) than for ash Z (180%).

TG/DTA/QMS thermal analysis of fly ashes with gas analysis in synthetic air

Figures 3–6 present the results of TG, DTG, DTA and the composition of the released gas products (CO₂—m/z 44, H₂O—m/z 17 and SO₃ m/z 64) in synthetic air atmosphere, respectively. All the results are presented on the basis of a temperature increase function up to 1100°C. QMID (quasi-multi-ion detection) stands for ion current for various gaseous (volatile) products expressed in amperes.

For both fly ashes in the temperature range between 20 and 200 °C, endothermic effect occurs (Figs. 3, 5) associated with the first mass loss, amounting to approx. 0.26 wt%. The mass loss is caused by the thermodesorption of physically adsorbed water confirmed by the increase in the intensity...
of the released gas products (m/z: 17) (Figs. 4, 6). The process of residual carbon combustion starts at the temperature range of 200 °C–230 °C. According to DTA curves of fly ash, X and Z reflect the occurrence of exothermic effects at 400°C, which is due to the combustion of residual carbon present in ash. The process is accompanied by a mass loss (TG curve from Figs. 3, 5). This is also proved by the significant increase in signals of m/z 44 on the recorded gas product curves (Figs. 4, 6).

Moreover, probably the combustion of hydrocarbons takes place, followed by water release. Then, the amorphous carbon (soot) is burned, as well as coke breeze. In the high temperature range of 700–800 °C, endothermic effect took place. It resulted from the thermal dissociation of calcium carbonate CaCO3 and was responsible for mass loss, amounting to 0.41% (fly ash X) and 0.42% (fly ash Z) (see DTA/TG curve in Fig. 3 and DTA/TG curve in Fig. 5). Thermal dissociation of CaCO3 is proved by the signal enhancement for gas products m/z 44 (CO2) (Figs. 4, 6). The most important several subsequent thermal processes take place at temperatures above 800 °C (Figs. 3, 5). DTA curve of fly ash X reveals exothermic effect followed by endothermic one. From other studies [28], it is known that among the exothermic effects, the sintering process and the crystallization of fly ash glassy phase are observed, while endothermic effects are associated with the appearance of the liquid

![Fig. 3 DTA and TG/DTG curves of fly ash X](image)

![Fig. 4 Analysis of H2O, CO2 and SO2 emission during thermal treatment of fly ash X](image)
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Phase and probably sublimation process. DTG curves above 800 °C demonstrated constant mass loss (over 1.5%). The maximum rate of mass loss is observed at 1016 °C (fly ash X) and 1039 °C (fly ash Z). The simultaneous increase in the intensity of the released gas products (CO₂—m/z 44 and SO₂—m/z 64) is observed (Figs. 4, 6). The registered released gas products of H₂O—m/z 17 prove that this loss is connected with moisture evaporation.

The sources of released gases are diverse. One of them is the release of gases closed in fly ash microspheres. This loss of H₂O and CO₂ and SO₂ is connected with the process of oxidation of the combustion of coal. The second source may be gas phase reactions. The authors preliminarily assumed that this is the main source of SO₂ release from ashes, because the release of this gas occurs only after exceeding the temperature of about 1000 °C.

Thermal analysis of fly ashes with gas analysis in helium atmosphere

The mass loss (TG curve in Figs. 3, 5) above the temperature of 800 °C is higher than the mass loss of combustion and mass loss of dehydration effects separately; however, during these two processes, the largest release of gases was recorded (Figs. 4, 6). Probably above 800 °C, gases released cannot be detected in air atmosphere. To address this problem, we performed DTA/TG analysis in which mass loss was monitored while heating fly ashes in a nonoxidizing...
helium atmosphere, followed by heating both samples in an air up to 800 °C. The results of DTG-MS tests are presented in Figs. 7–10 for the X and Z fly ashes heated in helium. The released gas products (O$_2$—m/z 32 and SO$_3$ m/z 64) are observed.

By running DTG-MS in different conditions, we found that before the temperature of 800 °C, there was an oxygen desorption from the measuring system and the sample itself (Figs. 7, 9). Oxygen desorption, along with the retention time of helium permeation and temperature was gradually extinguished. After exceeding the temperature of 800 °C, there is a significant increase of oxygen release from the sample. Moreover, after analysis of the MS results, we determined a completely different characteristic of SO$_2$ evolution. In the air, SO$_2$ release starts at about 1000 °C (Figs. 4, 6), in helium at around 850 °C (Figs. 8, 10).

Probably, during the thermal treatment, the dominant reaction process that causes emission of oxygen and sulfur dioxide is the decomposition of sulfur trioxide (which is redox reaction, where sulfur is reduced):

$$\text{SO}_3 \leftrightarrow \text{SO}_2 \uparrow + \frac{1}{2} \text{O}_2 \uparrow$$ (1)

This is a conventional reaction, due to the fact that SO$_3$ in the ashes exists in the form of SO$_4^{2-}$ sulfate ion incorporated into the glass. When helium environment was applied during DTG-MS test, the reaction striving for chemical equilibrium is shifted to the right, due to the negligible partial pressure of oxygen. This is due to a lack of oxygen in the measurement atmosphere.

As a consequence, the reaction begins at a lower temperature when the measurement was performed in helium atmosphere, in contrast to that one performed in air, and larger total amount of emitted SO$_2$ was also observed. The SO$_2$ release starts at a temperature of about 1000 °C (Figs. 4, 6), so it does not coincide with the beginning of temperature
where the mass loss of about 800–850 °C was observed (TG curves in Figs. 4, 6).

There may be other sources of gas emissions than those mentioned in reaction 1. If during the thermal treatment only the above reaction would occur, then the shapes and temperature ranges of the SO$_2$ and O$_2$ emission curves should be very similar. There are, however, some deviations which indicate that there may be other oxygen-emitting reaction mechanisms. For example, during the crystallization of the glassy phase of ashes, iron silicates are formed (fayalite) [22]. The presence of this phase is associated with the reduction of iron from +3 to +2 oxidation state, which is accompanied by the process of oxygen emission.

Change in chemical composition during thermal treatment

Chemical reactions involving the gas phase should cause differences in the chemical composition before and after the reaction. Therefore, fly ash and ash sinters from 1000 and 1100 °C were subjected to XRF analysis. The results of these analyses are presented in Table 4 in elemental form.

Seemingly, the results do not provide helpful information in explaining the issue of thermal bloating of fly ashes. First, it can be noted that there is less sulfur concentration in the sinters than in the raw ashes. This is particularly visible in the fly ash Z and the analog sinter from a temperature of 1100 °C. The second important information is the oxygen content in ashes and sinters from a temperature of 1100 °C. Oxygen is the main elemental component of the studied
materials and accounts for almost 50% of all elements. Taking into account the loss of ignition observed on the TG curves, an increase in the concentration of oxygen should be expected, assuming that it has neither been absorbed nor released. In brackets in Table 4, an estimation of the content of this element is given with the assumptions given above. As can be seen, the real values are more than 1% lower, which probably confirms that the oxygen was emitted during sintering. In the case of other elements, there were no unambiguous decreases in content, but only a slight increase related to the losses of ignition occurring during sintering of fly ashes.

### Conclusions

Rietveld quantitative X-ray diffraction analysis confirmed that both fly ashes studied consist mainly of glassy phase, which may constitute more than 80% of the powder mass. The presence of this vitreous phase indicates that a liquid phase is formed during sintering, which enhances the process and causes the shrinkage shown by the HMTA microscope studies. Moreover, fine-grained fly ashes have a high susceptibility to bloating. This phenomenon is favored by the high content of alkali oxides, fine granulation and the large amount of gases released from the liquid phase formed. The onset of the bloating phenomenon recorded by HMTA microscope occurred at much higher temperatures above 1130 °C. This discrepancy is a result of liquid phase occurrence, but on the other hand, it might be connected with a small sample size. The main gases that cause direct thermal bloating are SO₂ and O₂. By running DTG-MS in different conditions, we found that before the temperature of 800 °C, there was an oxygen desorption from the measuring system and the sample itself. We determined a completely different characteristic of SO₂ evolution. When helium environment was applied during DTG-MS test, SO₂ release starts at around 850 °C in comparison with DTG-MS test performed in air where the release of SO₂ was observed at about 1000 °C. Moreover, the release of oxygen registered may be the result of redox reaction of iron compounds, e.g., fayalite crystallization, hematite reduction to magnetite. Thermal analysis carried out in a different atmosphere proved that the firing atmosphere itself has a significant effect on the rate and temperature of gas emission. Fine-grained, alkaline-rich fly ashes can then be valuable flux additives in a sintered ceramic. Probably, the addition of suitable oxidants can effectively inhibit the redox reactions and eliminate the phenomenon of bloating.

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