

ELIMINATION OF TOXIC GAS EMISSIONS BY MgO SOAKED EXPANDED CLAY IN WASTE COMBUSTION CHAMBERS

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Scope

- This paper serves to describe the elimination of toxic gas emissions in coal combustion chambers and activity of expanded clay soaked MgO and other alkali materials such as Lime, Hydrated lime, NaCl, MgCl₂ and KCl used in desulfurization. A chemical analysis of ash materials used to characterize the elimination of toxic substances was carried out. As well, the ability of the expanded clay to remove inorganic and organic sulphur and other pollutant substances were determined with their respective process control challenges.
- In this study, combustion tests of Şırnak asphaltite and different types of Turkish lignites; Kütahya Gediz, Tunçbilek, Soma Kısırakdere were carried out. Lime, Hydrated lime, Magnesia, NaCl, MgCl₂ and KCl were used as desulfurizing solid sorbent. The different type of solid sorbents such as Şırnak limestone, marly limestone and claystone, MgO soaked expanded clay use in coal desulfurization were virtually investigated and discussed. Combustion of solid fuels in the presence of expanded clay was managed at low particle sized such as 1-2mm. Expanded clay was examined as an absorbent for a conversion of toxic gas to friendly emissions. It can be a promising waste incineration for the production of electricity from nylon and plastic contaminated municipal wastes because of high activity in the collection and leaching in the toxic gas in the combustion reaction.

Introduction

- Coal, biomass or waste as a solid fuel, waste liquid fuels from natural sources and biodiesel have gained more market due to its use in electricity. Combustion of coal, biomass or waste creates toxic gas emissions for environmental concern. The electricity production of Turkey from the primary resources are natural gas imported and coal as high as 22% (Figure 1) (TTK, 2009, TKİ, 2009, IEA 2014). Reasons for growing interest in toxic gas emission control include its potential for reducing noxious emissions. Uses of fuels as potential contributions to rural economic development reduce reliance on high quality fuels, as an additional demand centre for electricity commodities and as a way to urbanization (Demirbaş and Balat, 2004).
- Desulfurization of coal has firstly applied for the flue gas in coke production and in fluidized bed combustion systems with limestone addition into the coal combustion chamber. The pre-combustion desulfurization methods have significantly been developed by wet desulfurization units in thermal power stations. However, small scale operation of wet desulfurization plants may not be economic (Wheelock, 1979). Especially in waste incineration pre combustion emission control may cost higher prices such as 60-90\$/ton. The expanded clay pellets soaked magnesia slurry may cost lower such as 3-5\$/ton.

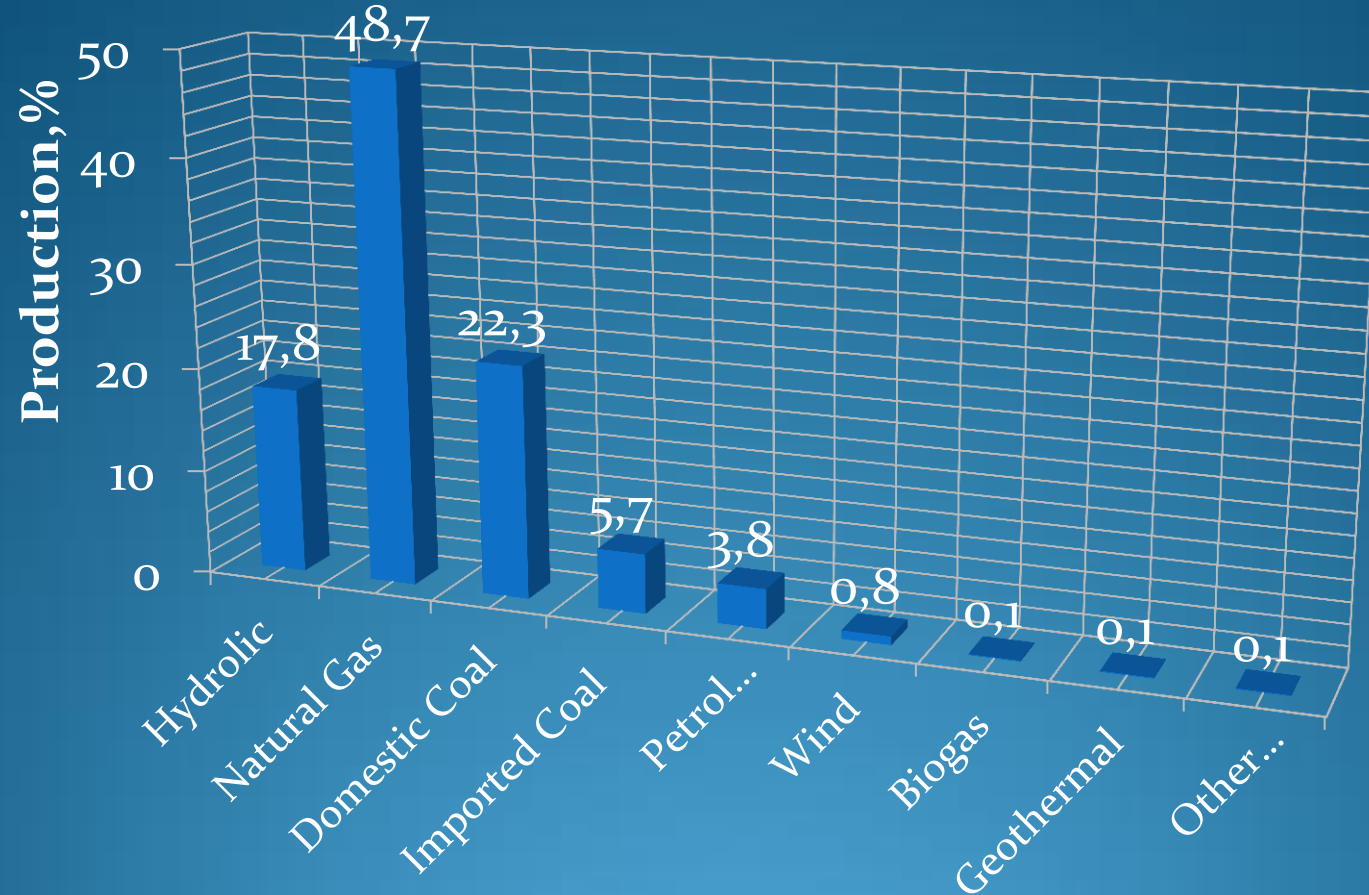


Fig. 1. The Revised Electricity Production from Turkish Primary Resources and Renewable Comparison.(IEA 2014)

- Conventional coal combustion systems using Stokers or grate chambers are not designed to treat potentially contaminated municipal organic waste in order to prevent by post combustion the potential spread of toxic emissions in coal and wastes (Tosun, 2013) and potential problems related to organic matter, phenol, such as undesirable colour, odour formation (Hartikainen et al. 2001). Solid adsorbents are needed in the combustion chamber systems typically consist of alkali salts intake, coagulation ash processes (Çulfaz et al. 1997, Tosun, 2007, Tosun, 2012).
- Specifically, combustion temperature and secondary air may improve to destroy or impair unwanted emissions through chemical adsorption (Sharma et al. 2008).
- The different type of chemical alkaline react with coal samples in combustion chamber at atmospheric pressure by the equations as given below (Kumar et al, 2000);
- $\text{SiO}_2 + 2\text{NaOH} / \text{Na}_2\text{CO}_3 \text{®} \text{Na}_2\text{SiO}_3 + \text{H}_2\text{O}$ (1)
- $\text{Al}_2\text{O}_3 + 2\text{NaOH} / \text{Na}_2\text{CO}_3 \text{®} \text{Na}_2\text{AlO}_2 + \text{H}_2\text{O}$ (2)
- $8\text{FeS}_2 / \text{S}_2 + 30\text{NaOH} / \text{Na}_2\text{CO}_3 \text{®} 14\text{Na}_2\text{S} + \text{Na}_2\text{S}_2\text{O}_3 / \text{Na}_2\text{SO}_3 + 15\text{H}_2\text{O} + 4\text{Fe}_2\text{O}_3$ (3)

Factors affecting toxic gas sorption in combustion

- Effective sorption in combustion processes depend on numerous factors including coal rank in carbonization, the volatile gaseous matter of coal such as presence of hydrogen, carbonyl gas and oxidation rate so stabilizing the desorbance, the settings of optimal diffusion conditions including structure defects (nitrogen, phosphorus, sulfur, etc.), temperature, oxygen content of coal, etc. and optimization of carbondioksit concentration ratios added the adsorption–desorption balance, the residence time and the spatial distribution of molecules in coal pores among other factors determining the efficiency of carbonization. as factors affecting the rate and extent of carbonization much dependent on the site activation, its desorption properties and its porosity. As discussed in the previous section, carbonization is a prerequisite step for oil generation from biomass wastes and coal (Wheelock, 1979).

METHOD AND MATERIALS

- In this research, representative specimens of the Şırnak asphaltite and different types of Turkish lignites; Kütahya Gediz, Tunçbilek, Soma Kısırakdere were crushed and comminuted to minus 1mm size by controlled screening. Air dried samples of 40-50 gr from each different coal types were prepared and sealed in nylon bags. The different type of solid sorbents such as Şırnak limestone, marly limestone and claystone were used in the combustion. The chemical analysis of the solid sorbents are given in Table 1. The results of proximate and ultimate analyses of various Turkish coals used in the experiments are given in Table 2 and Table 2. The qualities of processed coal products are ascertained by chemical and standard coal analysis of ASTM 3173-3177.

Table 1. The chemical analysis values of limestone, marl and claystone of Şırnak province

% Sorbent	Şırnak Limestone	Şırnak Marly limestone	Şırnak Marl	Şırnak Claystone	Expanded Clay soaked MgO
SiO ₂	3,53	9,42	24,14	48,53	44,3
Al ₂ O ₃	2,23	6,53	12,61	24,61	22,1
Fe ₂ O ₃	0,59	4,48	7,34	7,59	6,5
CaO	49,48	39,23	29,18	9,48	7,4
MgO	2,20	2,28	4,68	3,28	9,28
K ₂ O	0,41	0,53	3,32	2,51	2,51
Na ₂ O	0,35	0,24	1,11	0,35	0,35
Ignition Loss	46,19	26,11	21,43	6,09	0,09
SO ₃	0,32	0,21	0,20	0,32	

Table 2. Proximate Analysis of Turkish Lignite and Asphaltite. (ADB:Air dried base. DB:Dried base, DAB:Dried ashless base).

Coal Type	Ash,% ADB	Moisture ,% ADB	TotalS, % DB	Volatile Matter,% DAB
Şırnak Asphaltite	46.3	0.1	7.1	62.6
Tunçbilek Lignite	29.3	18.1	3.1	52.6
Kütahya Gediz	22.0	1.7	3.6	42.7
Soma Kısırakdere	13.8	14.0	2.2	40.4

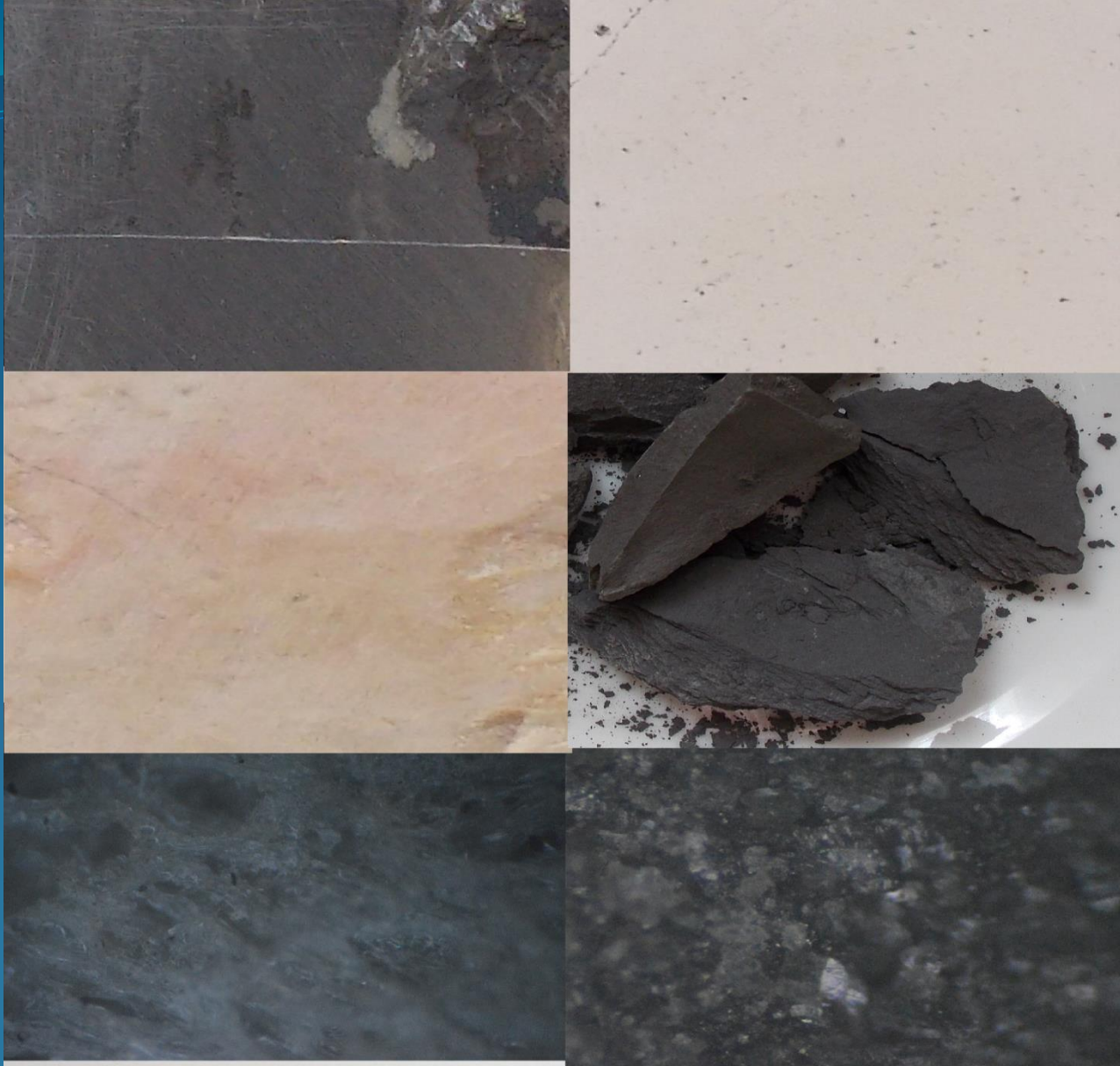


Fig. 2. Photos and Bright Sections and Parts of Şirnak limestone, marl, claystone



Film. Coal, Wood, waste pellet combustion.



Film. Coal, Wood, waste pellet combustion.

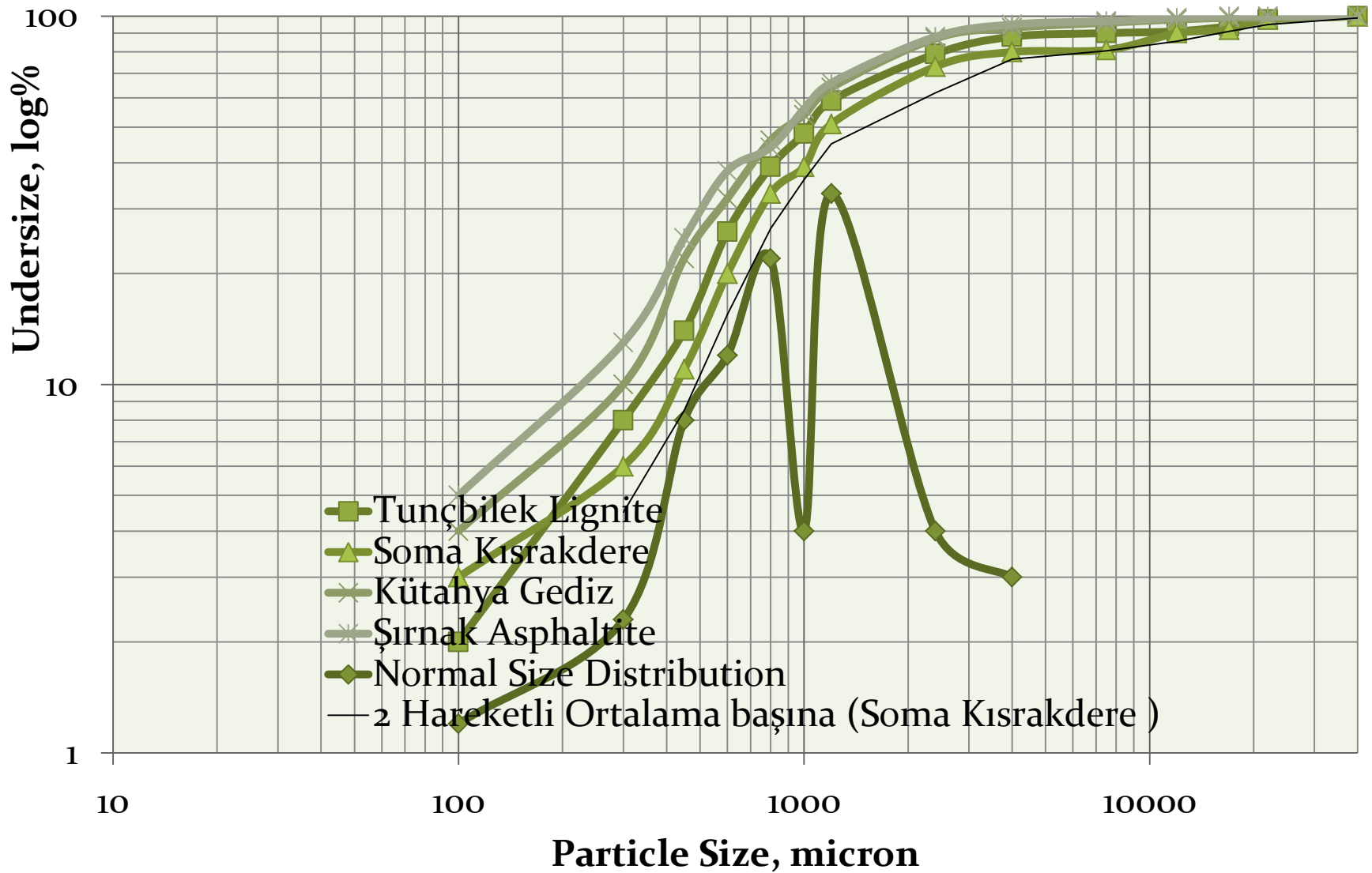


Fig. 3. Particle Size Distribution of Turkish High Sulfur Coals.

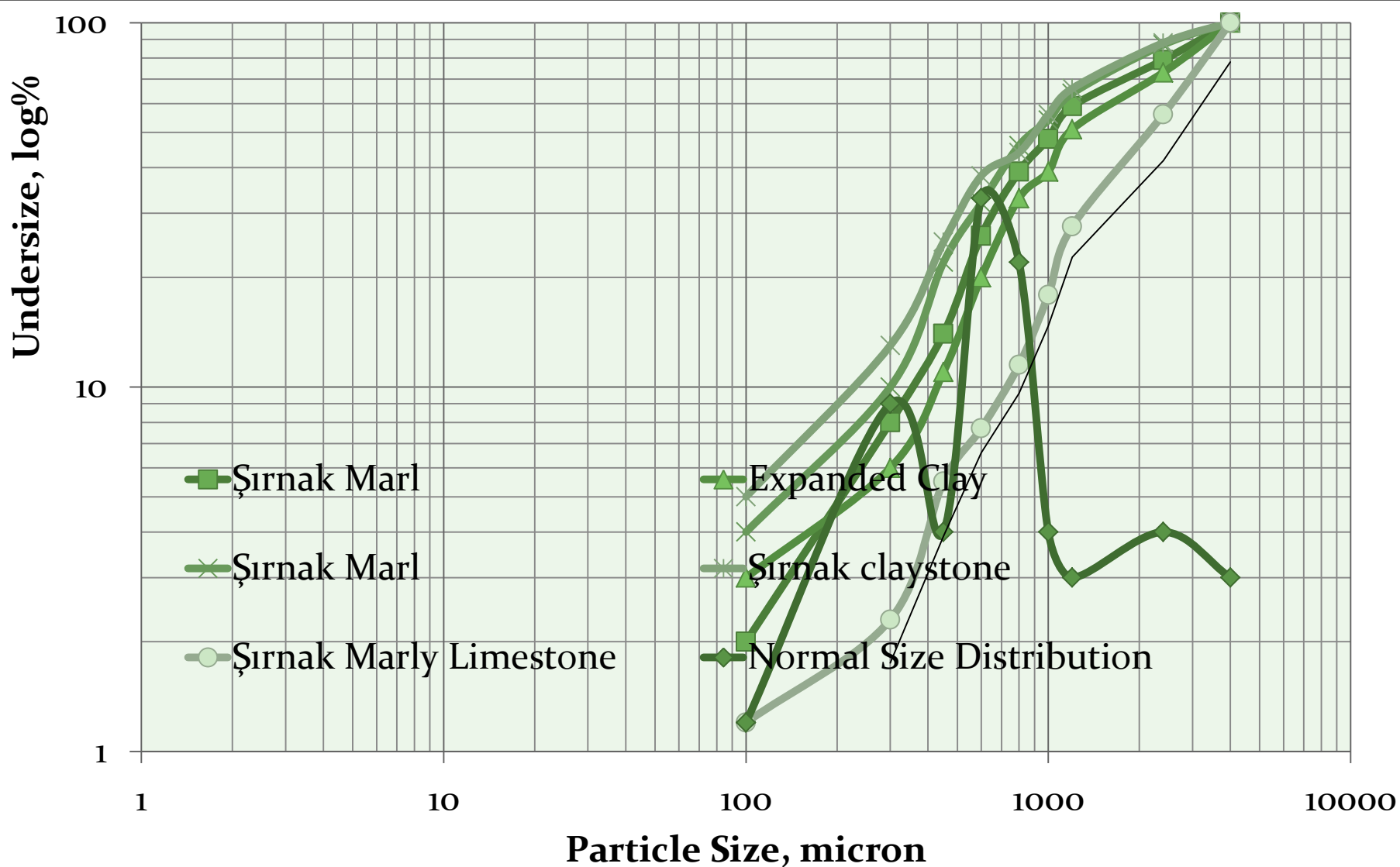


Fig. 4. Particle Size Distribution of of Solid Sorbents of Şırnak limestone, marly limestone, marl, claystone, expanded clay

As seen from Figure 3, 80% of weights of samples were under 3 mm. The lignite samples were mainly distributed between 1mm and 3 mm size fractions. As seen from Figure 4, 80% of weights of solid sorbents were under 2 mm. Two normal distributions are seen from Figure 4 due to different mechanical breakage manners of solid sorbents. Especially, hardly crushed particle size fraction of sorbents was ranging between 2 and 3 mm.

Combustion experiments were carried out in a benchmarked laboratory type 1m kiln reactor put in the furnace at atmospheric pressure at a temperature precision of $\pm 5^{\circ}\text{C}$ as seen from Figure 5.

Coal Combustion

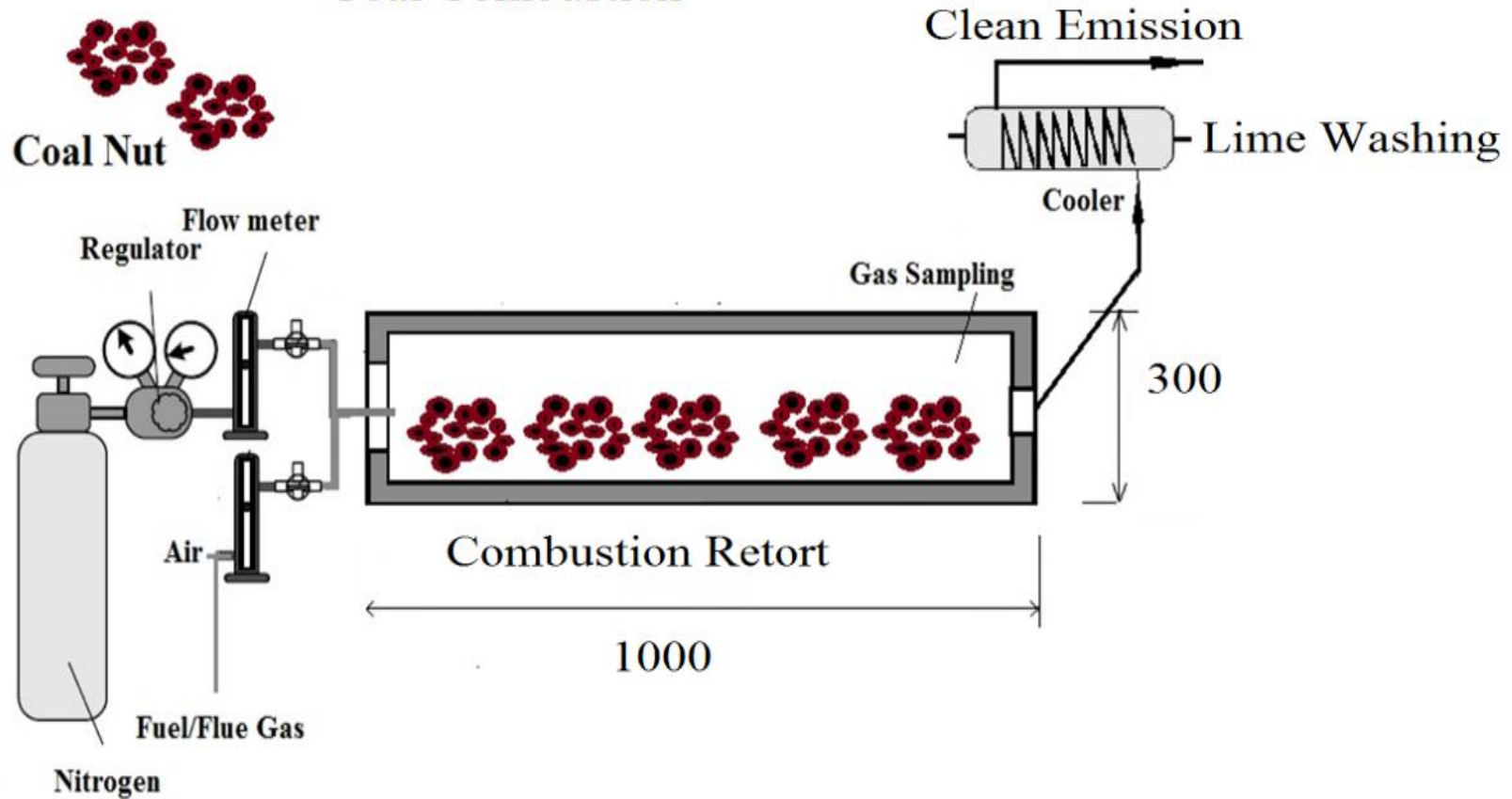


Fig. 5. Use Coal Combustion Retort subjected to solid sorbent

RESULTS AND DISCUSSION

Turkish lignites may not be destroyed by controlled crushing and screening till reducing particle size of specimens to minus 10 mm. As seen from Figure 3, normal distribution of coal size was determined as two different fractions and highly sufficient in order to combustion and react with solid sorbents.



- Fig. 9. Expanded Clay MgO Soaked photo 500X, a.prior to combustion at 800°C



Fig. 9. Expanded Clay MgO Soaked photo 500X,
b. Combustion at 800°C

Combustion tests were carried out under atmospheric pressure at a constant time period of 3 hours previously determined over 1-2 kg lignite samples. MgO soaked expanded clay were made by calcination of Şırnak claystone at 800 °C and soaking with MgO and SEM picture was illustrated in Figure 6. Total solid sorbent weight was hold constant at a quarter of coal weight. Various sorbents were used at 1/4 weight rate into to coal samples. The effect of particle size of solid sorbents were investigated over the combustion of Şırnak Asphaltite and carried out well on emitted gas substance subjected to reaction with expanded clay in combustion, as shown in Fig. 7.

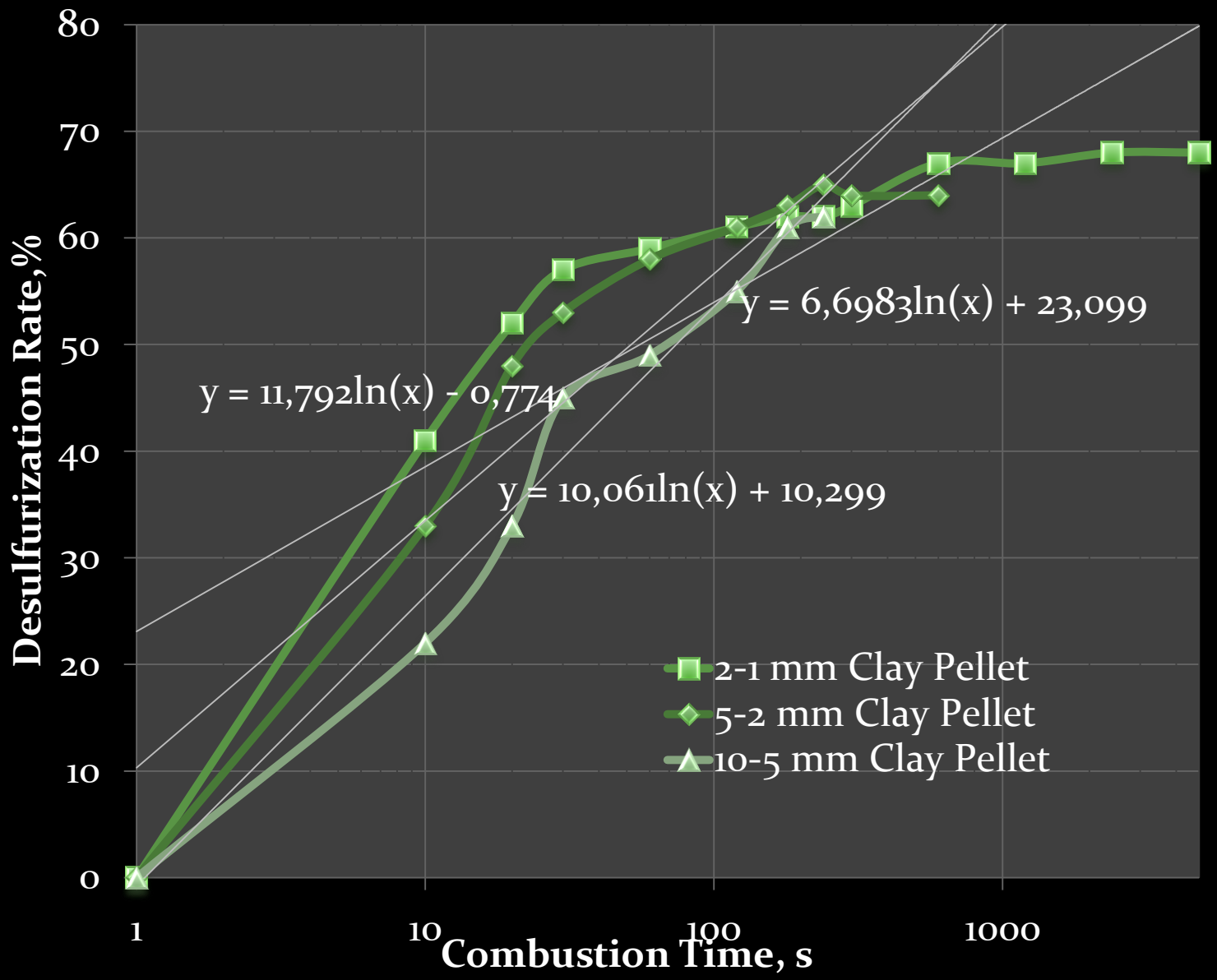


Fig. 7. The Effect of Particle Size of Expanded Clay on Desulfurization

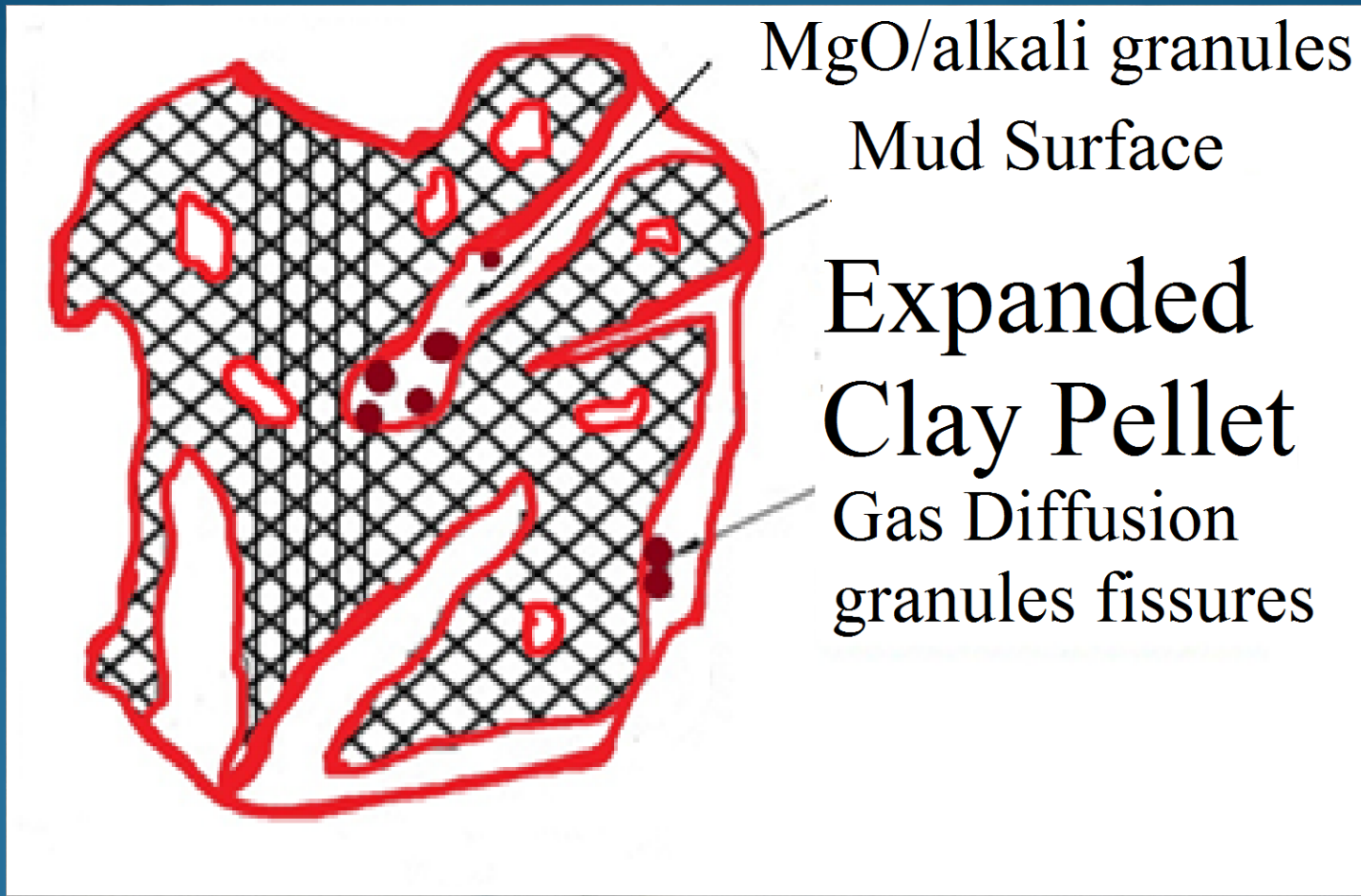


Fig. 8 Expanded Clay surface adsorption and pore entrapment.

Although molecular gas diffusion is believed to be the primary mass transport process in the combustion chamber, complex convective gas emissions proliferated the alkali clusters below 1-2mm size and exothermic combustion reactions increased toxic substances in the gas form, a relatively porous structure of expanded clay interstitial spaces and cracks reduced over 5mm size. The combustion gases substances towards the expanded clay surface through this surface alkali is primarily accomplished by molecular diffusion across the micro cracks and alkali clusters. A reduction in the size of combustion coal also permitted more clay substrate to diffuse through the surface towards the expanded clay sites subsequently increase adsorption as illustrated in Figure 8.

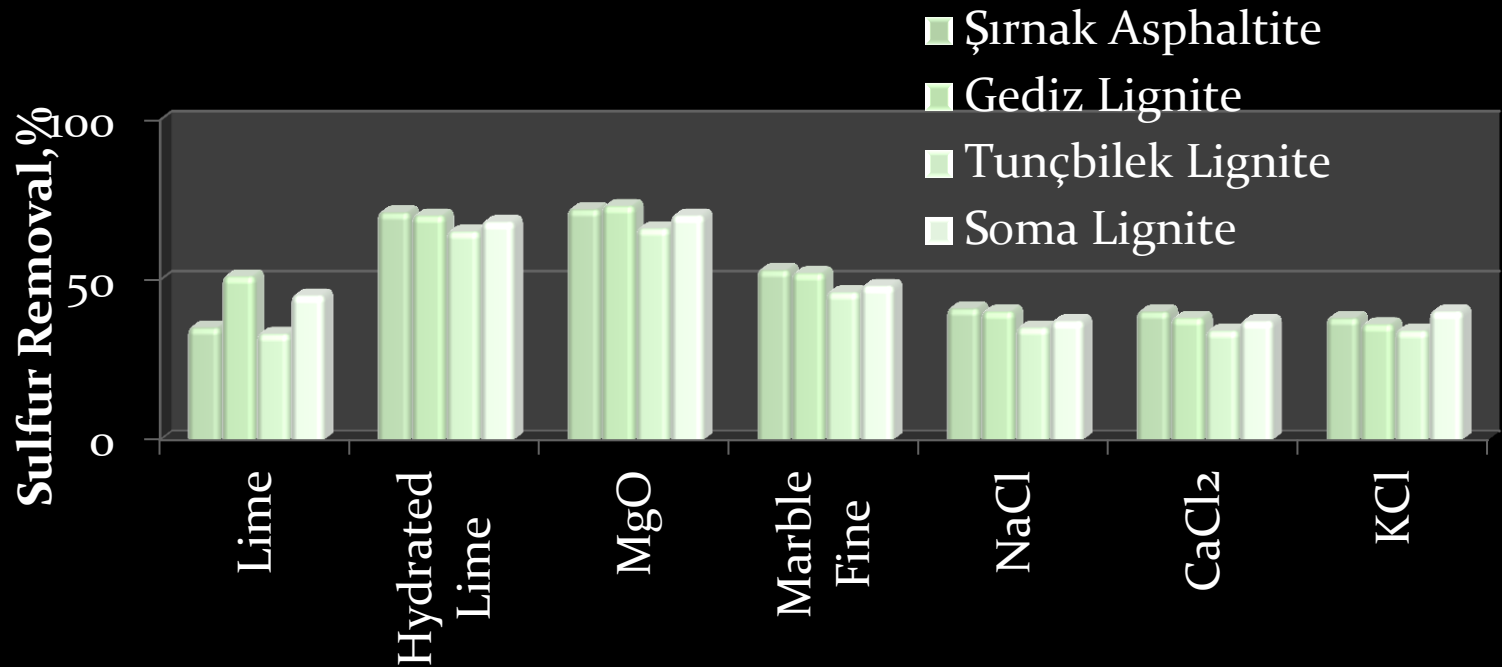


Fig. 10 The effect of alkali type on Desulfurization in Kiln Combustion of Turkish Lignites at 800 °C

Fig. 11.
 The
 Combusti
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 Turkish
 Coals with
 Expanded
 Clay
 %10MgO
 Soaked,
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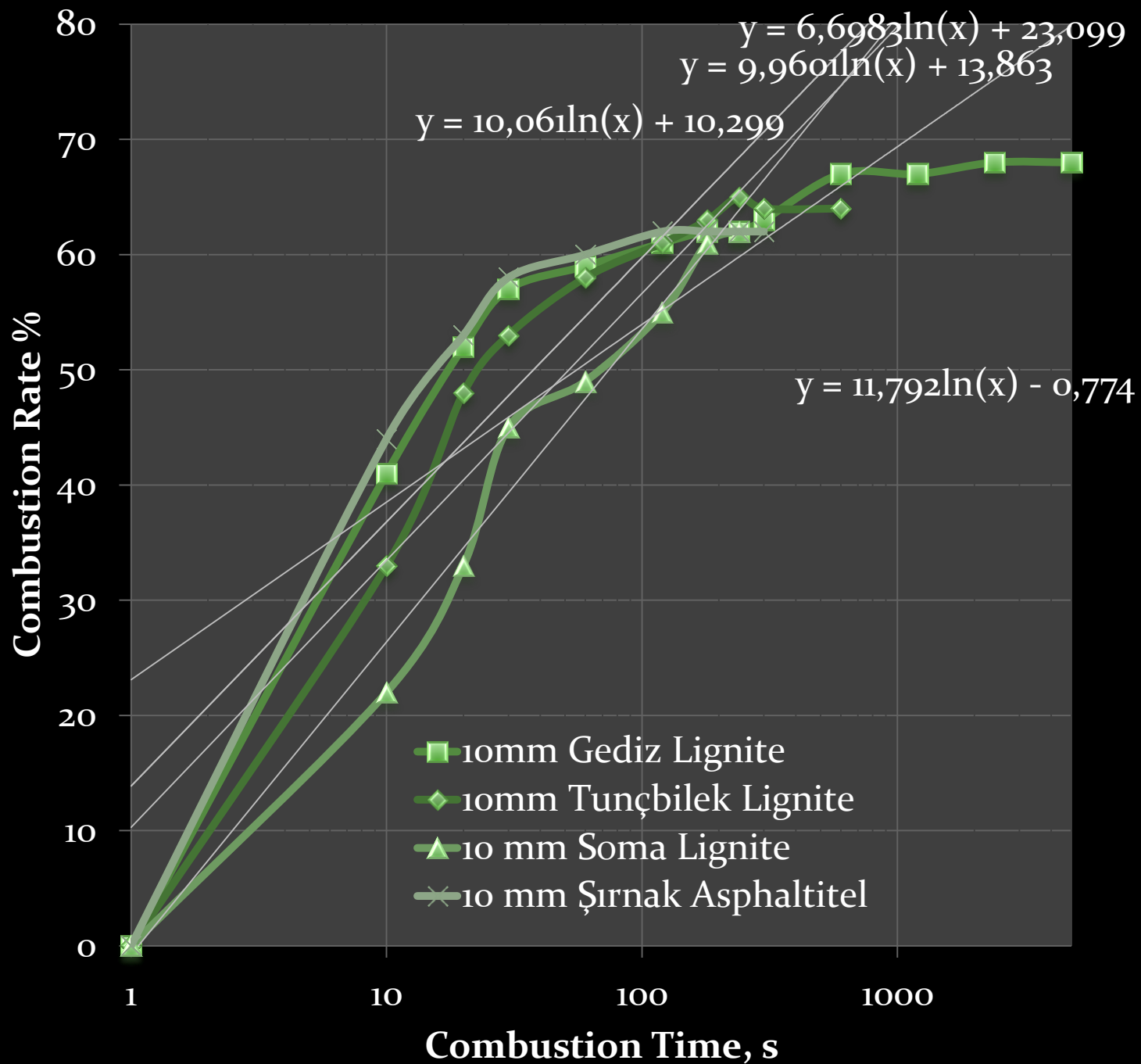


Fig. 12.
The sulphur removal effect of Expanded Clay %10MgO Soaked, Combustion on Şırnak Asphaltite at 800°C

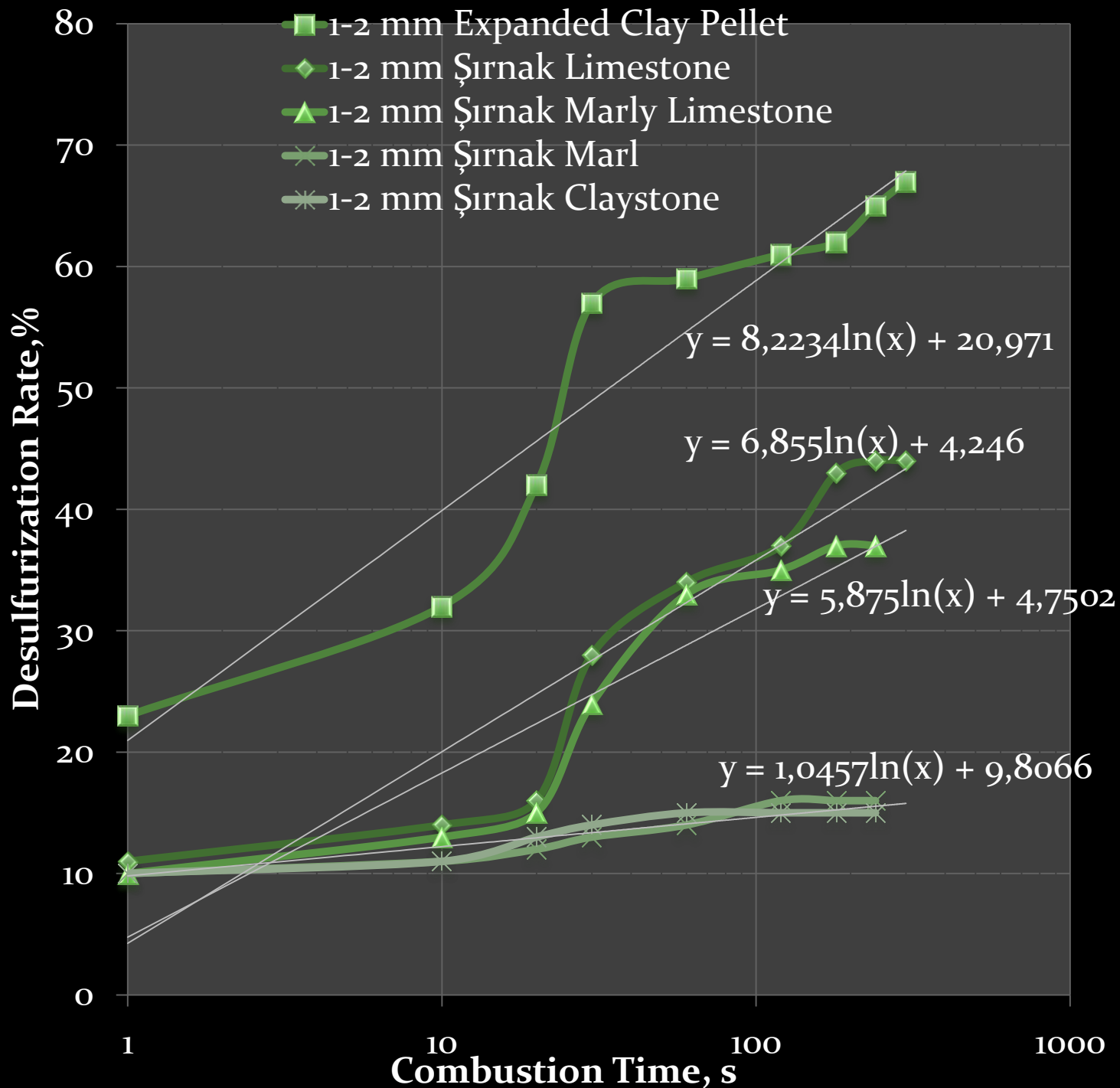
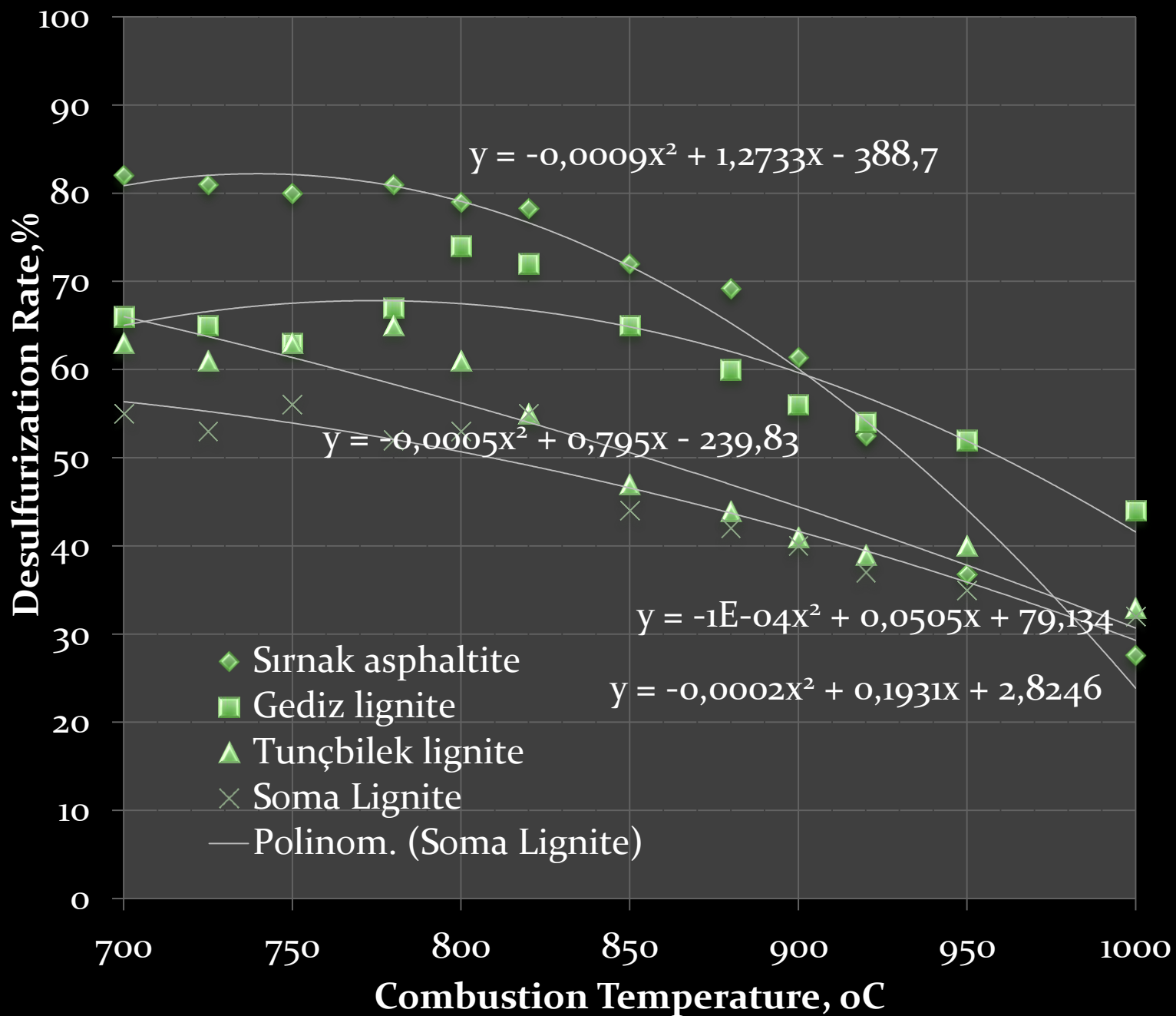


Fig. 13. The combustion temperature on the Desulfurization removal of Expanded Clay %10MgO Soaked.



The gaseous reacted adsorbate then adsorbs to the sorbent in an certain amount that is equal to the amount of previous adsorbate that was partially degraded on the surface of the expanded clay and removing aliphatic hydrocarbons and phenols/chlorinated phenols, carbonyl toxins, along with organic matter related odour substances.

In the combustion experiments, the experimental condition is calculated on the basis of the ash composition in the ambient state. So neither the contained water vapour nor the condensing hydrocarbons are taken into account.

Among various cheap alkali fines such as Lime, Hydrated lime, Magnesia, NaCl, MgCl₂ and KCl, magnesia were found as efficient desulfurizing solid sorbent. 72-74% desulfurization rates with Şırnak asphaltite, Gediz and Soma lignite could be reached. This cheap alkali sorbent fines may be so feasible at the side of cost and sorbent production. The high amount of gaseous emissions might be reduced instead of massive alkali sorbent use. Advanced coal washing of Turkish lignite may not be feasible. However, the combustion with solid expanded clay soaked alkali fine of Turkish lignites can be managed. The heavy metal and toxic emissions of soot, nitrogen oxides and sulfur oxides with expanded clay soaked MgO managed at high elimination rates ranging 72-74%.

The approach of combustion kinetics assumed basically that the process exponentially was developed itself, as seen in Figure 11 with all specific features. The elimination of emissions with expanded clay sorbent was a decisive factor for the path of the kinetics of combustion reactions of coal. Therefore a static model of coal combustion was developed at 10mm of coal nut size. Instead of fluid bed combustion, packed bed combustion of coarse size coals is highly governed by slow combustion reactions, sufficiently sorption of toxic gas.

The different type of solid sorbents such as Şırnak limestone, marly limestone, marl and claystone, MgO soaked expanded clay were used at 1-2 mm size in coal combustion at 800°C and the effect of the massive solid sorbent type on elimination of toxic gas emissions were investigated and the results were illustrated in Figure 12. In comparison of use massive solid sorbents in combustion of solid fuels with the presence of expanded clay it was found that at low particle sized such as 1-2mm lower surface area of massive solid sorbents reduced desulfurization rate. As seen in Figure 12, the expanded clay soaked MgO examined was more efficient as an absorbent for a conversion of toxic gas to friendly emissions. It can be a promising waste incineration for the production of electricity from nylon and plastic contaminated municipal wastes because of high activity in the collection and leaching in the toxic gas in the combustion reaction. The desulfurization rates reached to 74% with Şırnak asphaltite.

In the combustion experiments with addition expanded clay soaked 10% MgO, reactor temperature changed between 700°C and 1000°C. Products received from combustion of lignite with solid sorbent at 1-2 mm size at a quarter weight rate to coal following 3 hours combustion were subjected to analysis for sulfur hold-up determination. Test results of combustion by expanded clay between 700 °C and 1000 °C were seen in Figure 13.

From the point of view of temperature effect experimentation, the resulted ash and sorbents analysis following combustion showed that increasing temperature reaching 1000 °C in combustion kiln for biomass, lignite and other coal samples were suddenly combusted the volatile substance without reacting sorbent matter in the kiln even in comparison with different sorbent evaluation and so we may reduce the effect of ash sorption.

CONCLUSIONS

The higher desulfurization yields were provided in combustion tests with using the expanded clays with MgO soaked with high sulfur coals in the kiln apparatus at a quarter weight rate to coal.

Combustion of different types of Turkish lignite was successfully processed in terms of desulfurization and depend on the volatile matter.

At higher rates of combustion of different types of Turkish lignite could be obtained from the tests using high combustion temperature of 1000°C. It has been clearly determined that CO₂ and steam much beneficial in elimination of toxic matter of different types of Turkish lignite. Şırnak asphaltites should be also desulfurized at high rate of 88 and 74% and high ash content reduced toxic gas emission in combustion.

Benefaction from high sulfur Turkish lignite in the various parametric combustion systems, in order to receive clean energy clean liquid and gaseous products must be managed in low temperature reactions with expanded clay soaked with MgO. It is also advised that the high amount of elimination of toxic gas will be managed at low combustion temperatures over 700 °C and more environmental friendly gaseous emissions were provided by 1-2 mm sized expanded clay in comparison same sized limestone.

Nut sized coal below 10mm sized in retort combustion carried out for Turkish lignite and Şırnak asphaltite showed sufficient desulphurization rates at 800 °C and even other lignites showed similar trend.

In order to optimize the desulfurization rates of the coal combustion and for the elimination high rate of detoxification in waste incineration process as given in Figure 13 the low 800 °C combustion was advised at nut size solid fuel combustion and with finer particle size fractions of coal specimens, the reactor temperature should be optimized to lower combustion kinetics mixed with expanded clay soaked MgO at 10% weight rate. However, as seen in Figure 12 it was showed that desulfurization rate reduced lower than 42 and 37% level at the addition a quarter weight of Şırnak limestone and marly limestone in to the combustion, respectively. That soot remained in ash was among 3,8-5,6%. Therefore it was supposed that porous expanded clay layers, improved finer fill of porous clay layers with alkali and ash metal catalysts of Turkish lignite and even exhibited sufficient gas permeability in small particle size fractions.

Thanks for attention.

Şirnak University, Turkey