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## Original Research Paper

## Cleaning of high pyritic sulfur fine coal via flotation

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#### ABSTRACT

Large quantities of fine coal have been produced during mining and processing activities of coals. Use of these fine coals after cleaning is important in terms of recovering of economically valuable energy source and prevention of environmental pollution. In the present study, amenability of a high ash-sulfur coal sample to flotation process was investigated. Coal particle size and collector dosage were selected as variables. Increasing coal particle size affected sulfur and ash rejections adversely. On the other hand, more combustible matter was recovered at coarser particle sizes. Maximum combustible recovery, ash rejection, pyritic sulfur rejection and sulfate sulfur rejection were achieved to be 82.56%, 82.84%, 90.04% and 81.45%, respectively. Polished section examinations and scanning electron microscopy-energy dispersive spectroscopy (SEM-EDS) analysis confirmed the rejection of significant level of ash and sulfur from the coal.

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#### 1. Introduction

Coal is an important resource for humanity due to its huge reserves and role as fuel. However it is also a candidate for environmental pollution because of its ash and sulfur content [1]. Sulfur restricts coal usage due to its negative effect on environmental pollution [2]. Environmental awareness on sulfur emissions have been increased in all over the world. Therefore, all coal producers will have to be able to meet coal standards in the near future [3].

Sulfur levels in coals are highly variable in the world. It ranges from 0.5% to 11% [3]. Coal with sulfur percentage of over 3% is defined as high sulfur coal [4]. Sulfur in coal can be present in organic and inorganic forms [1,5,6]. Organic sulfur is chemically bonded with the carbon in coal [6]. The inorganic sulfur is mostly found in disulfides and sulfates [5]. Pyrite is major sulfur containing inorganic matter in coal [7].

Combustion of low quality coal causes several environment 58 problems such as acid rains, air pollution [6,8,9], food pollution 59 [6], loss in energy recovery, waste accumulation [8]. Coal cleaning 60 61 before combustion is required to increase the quality of coal and to prevent its negative effects on environment. The methods applied 62 63 for coal cleaning may be physical, physico-chemical, chemical. In addition, several methods assisted by microwave, ultrasonic and 64 65 bacteria have been improved recently [10].

Among cleaning methods, froth flotation, oil agglomeration,flocculation, and enhanced gravity separation methods have been

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used for cleaning of fine coal [11]. However, flotation is the most common method among them [12] for removal of ash sources [13] and inorganic sulfur [6,14,15] from fine coal. Effective sulfur reduction by physical coal cleaning processes cannot be achieved unless sulfur is reasonably well liberated from coal. Hence, it is important to emphasize that the sulfur removal potential of a coal depends on both the quantity and degree of liberation of pyrite [16]. Separation by flotation is based on differences in hydrophobicity of organic coal and mineral matters. Mineral matters are generally hydrophilic unlike hydrophobic coal matter [17]. In flotation, hydrophilic mineral matter remained in the pulp while coal is taken into the froth [2].

Many parameters, such as feed rate, mineralogy, particle size, pulp density, agitation, pH, types-dosages of collectors and frothers influence the combustible recovery, ash rejection, sulfur rejection in coal flotation [3]. Particle size is one of the most important parameters among them [18]. The effect of size fractions on coal flotation has been previously investigated.

Li et al. [19] investigated the effect of particle size on selectivity of the flotation process. The best selectivity was obtained for the middle size fraction of 0.25–0.075 mm, while the flotation selectivity of larger (0.5–0.25 mm) and smaller (–0.075 mm) particles was diminished. Jorjani et al. [20] examined the different size fractions of coal in term of their maceral composition and flotation response. Among the different size fractions of -500 + 150, -150 + 75, -75+ 40, -40 + 25, and  $-25 \,\mu$ m, -40 + 25, and  $-25 \,\mu$ m size fractions were found to have greater amount of liptinite. It was stated that liptinite rich samples had better flotation response and separation

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E. Sahinoglu/Advanced Powder Technology xxx (2018) xxx-xxx

96 the maceral groups by particle size is as important as other estab-97 lished factors in flotation. Wei et al. [21] examined the role of dif-98 ferent parameters in float stability that is an important 99 determinant of performance of the flotation and revealed that froth 100 stability changed depending on coal particle size, process water quality, and chemical reagents dosage. The froth stability in flota-101 102 tion of fine coal particles  $(-106 \,\mu m)$  with process water was found to be very high due to the interaction between particle size and 103 salts. Mixing of coarse and fine coals, dilution of process water with 104 de-ionised water and reducing reagent dosages had adverse effects 105 on froth stability. Liang et al. [22] reported that more stable frothes 106 were created by finer hydrophilic particles having no considerable 107 effect in maintaining the froth in the froth decay process. Fine and 108 moderate hydrophobic particles enhanced the froth stability in the 109 110 froth formation process. Fine particles with high hydrophobicity 111 had the minimum effect in stabilizing the froth due to the lowest 112 water solid ratio. Sahoo et al. [23] classified the coal into five discrete size fractions  $(-500 + 150 \,\mu\text{m}, -150 + 75 \,\mu\text{m}, -75 + 36 \,\mu\text{m},$ 113  $-36 + 25 \,\mu$ m, and  $-25 \,\mu$ m) and determined the effect of size distri-114 bution on flotation kinetics. It was reported that combustible 115 116 recovery was not be affected significantly by feed size. Maximum 117 recovery was achieved at the greatest particle of  $-500 + 150 \,\mu\text{m}$ .

Most of the previous studies some of which was mentioned 118 119 above evaluated the performance of flotation process in terms of 120 only combustible recovery and ash rejection. However, current 121 study focuses also on pyritic sulfur rejection. Pyritic sulfur rejec-122 tion is comes first to determine the contribution of any coal cleaning process to environmentally friendly energy generation. In this 123 study, a coal sample with considerably high pyritic and sulfate sul-124 125 fur content was used. No previous study related to flotation of 126 Muzret coal has not been met before. Extremely high pyritic and

#### Table 1

Proximate, sulfur and calorific value analyses of the coal sample.

	Air dried	Dried
Proximate analysis		
Moisture (%)	2.25	_
Ash (%)	34.85	35.65
Volatile matter (%)	10.73	10.98
Fixed carbon (%)	52.17	53.37
Sulfur analysis		
Sulfate sulfur (%)	0.99	1.01
Pyritic sulfur (%)	5.44	5.57
Organic sulfur (%)	1.3	1.33
Total sulfur (%)	7.73	7.91
Calorific value analysis		
Calorific value (kcal/kg)	4970	5084

sulfate sulfur content of Muzret coal make its flotation worth 127 researching. In the present study, flotation performance the coal 128 sample of different particle sizes was investigated by using differ-129 ent dosage of kerosene as collector. Combustible recovery, ash 130 rejection, pyritic sulfur rejection, and sulfate sulfur rejection of 131 the process were determined. The findings obtained in the study 132 were supported by advanced characterization techniques (i.e. pol-133 ished section and SEM-EDS examinations), which provided an 134 invaluable insight into the flotation process. 135

#### 2. Materials and methods

Coal sample was obtained from a coal deposit in Muzret-Artvin 137 region of Turkey. A fine coal sample of  $\approx$ 150 kg was taken from 138 coal deposit. Large portion of the sample was below a particle size 139 of 0.5 mm. Remaining part whose size was over 0.5 mm was 140 reduced to -0.5 by controlled dry grinding. In controlled grinding, 141 time of the grinding was kept short to prevent overgrinding. After 142 screening, oversize material was subjected to another short-time 143 grinding. Representative part divided from main sample was sub-144 jected to size reduction process to obtain material with size frac-145 tion of -0.25 mm and -0.125 mm. Cone - quartering and chute 146 rifflers was used as repeated cycles successively for dividing the 147 sample into small quantities. Final samples of 100 g to be used in 148 the tests were prepared by using the method of chessboard 149 squares. 150



Fig. 2. Particle size distributions of coal samples.



Fig. 1. Pyrite grains of different size and shapes in coal structure.

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