A STUDY ON THE CHARACTERIZATION OF RED MUD

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Abstract

Red Mud (RM) is a solid waste product obtained from aluminum industries via Bayer’s process. India is one of the largest producers of RM producing around 4 million tons annually. Its disposal is one of the major concerns as it is highly alkaline in nature and is a potential threat to air, water and land. Currently RM is being dumped on land or in RM ponds near the alumina refineries and with the cost of the land going high it is becoming a big problem for the industries. Due to the differences in the ore type in its production process, there is a variation in the mineralogical composition between the residues from India and other countries. Its properties such as iron content in the form of ferric oxide (Fe₂O₃), high surface area etc. makes it attractive potential reactants for many reactions. In this paper the structural and phase compositions of RM is being studied with the help of PSA, XRD, FESEM, EDX, BET and FT-IR techniques.

Key words: Red Mud, PSA, XRD, FESEM, EDX, FT-IR

I. INTRODUCTION

Red Mud (RM) is a solid waste product obtained from aluminum industries which is reddish brown in color. RM is mainly made up of oxides of iron, aluminum, silicon and titanium. The primary composition of red mud is Al₂O₃ (17-20%), Fe₂O₃ (48-54%), SiO₂ (4-6%), TiO₂ (3-4%), Na₂O (3-5%), CaO (1-2%) [1]. RM is usually considered as an inert solid waste but it is strongly alkaline and highly corrosive in nature [2]. It is usually discharged as highly alkaline slurry (pH 10-13.5) with 15-40% solids from the industries and because of this highly alkaline character, its disposal is a big problem for the industries as it may lead to soil, water and air pollution. The chemical and mineralogical composition of RM may however slightly change, depending on the source of bauxite and on the technological processing conditions. It is reported that 0.8–1.5 tons of red mud is generated per ton of alumina produced [3]. Globally 60-120 million tons of RM is produced annually and about 2 million tons of RM is produced in India [4].

RM can be utilized in various ways like recovery of Fe, Al, Na [5], use in cement production [6], brick production [7], glass production [8], aerated concrete blocks [9], absorb heavy metallic ions like Cu²⁺, Zn²⁺, Ni²⁺, Cd²⁺ [10] and also for the removal of H₂S [11] and Fluoride [12]. During recent years various studies have been carried out on the physical and chemical properties but the characterization results obtained in each case varies from place to place. In the present paper RM sample is analyzed by PSA, XRD, FESEM, EDX, BET and FT-IR techniques. This research gives emphasize on the comprehensive utilization of RM and provides an important base for further studies of it.

II. MATERIAL AND EXPERIMENTAL METHODS

A. Materials

The Red Mud used in this study is collected from NALCO, Damanjodi, Odisha, India. The samples are obtained in slurry form. On drying this slurry, RM lumps are produced. The collected RM lumps are sieved after ball milling it for 45 minutes. The average particle size of RM is found to be 63 microns by sieve analysis. The samples are then stored in plastic sealed packets for further analysis.

B. Experimental Methods

The particle size of the RM samples is measured using Malvern Mastersizer (Range: 0.02µ to 2000µ). The XRD patterns are analyzed using Philips X’Pert X-Ray diffractometer with a Cu Kα radiation source in a
2θ range of 10° to 70° at spanning range of 3° min⁻¹. FESEM and Energy Dispersive X-Ray (EDX) are used to study the surface morphology and elemental composition of the samples respectively using Nova Nano SEM 450. The FR-IR spectra of the samples are obtained by using Perkin Elmer FT-IR Spectrometer (Spectrum RX-I). The range is taken between 4000-400 cm⁻¹. The BET surface area of the samples is also measured by Quantachrome Autosorb-1 using liquid N₂.

II. RESULTS AND DISCUSSIONS

A. PSA Analysis:

The particle size distribution graph of the RM particle is shown in Fig. 1. It can be observed that the particle size of the sieved RM particle varies from 0.0136µ to 990.5362µ with an average particle size of 12.106µ. Such wide range of particle size is observed because simple sieving using a sieve shaker is just a raw method to obtain a specific particle size.

B. XRD Analysis:

The elemental analysis and phase characterization of RM have been reported many times before but the analysis data of the composition of RM is not uniform. From Fig. 2 it can be seen that the main components are Hematite (Fe₂O₃), Gibbsite (Al(OH)₃), Rutile (TiO₂), Calcite (CaCO₃), Sodium aluminum silicate (Na(AlSiO₄)), Dicalcium silicate (Ca₂SiO₄), and Quartz (SiO₂). The XRD pattern of the elements are referred form JCPDS file of X’Pert HighScore software. Using peak broadening technique of X’Pert HighScore software for XRD analysis, one can clearly differentiate the peaks.

C. FE-SEM and EDX Analysis:

To analyze the morphological structure of the RM sample, the sample is dispersed in methanol and ultrasonicated for 45 minutes. Then the samples are observed under Field Emission Scanning Electron Microscopy. The FESEM images of RM sample is shown in Fig. 3. It is observed that the arrangement of the particles is relatively loose with high porosity and small particle size.

The EDX analysis of the RM sample ensures the presence of particular elements in it and is shown in Fig. 6. The main elements observed are Fe, O, Ti, Al and Si.
D. FT-IR Analysis:

The relative FT-IR spectrum of RM is shown in Fig. 7. It is observed from the figure that broad bands occurred around 2919.50 cm⁻¹. This may be due to the stretching vibrations of O-H bonds and H-O-H bending vibrations of interlayer adsorbed H₂O molecule [13]. Here the change in the dipole moment of the water hydroxyl-stretching plays an important role and so the intensity of the infrared spectrum is more. Stretching vibrations of C=O is found at 1449.40 cm⁻¹ confirms the presence of carbonate groups [14]. The main reason being the presence of chemisorbed CO₂ in RM. Characteristic bands of Si-O and O-Si-O group are also observed at 989.30 cm⁻¹ and 534.50 cm⁻¹ confirms the presence of silicate groups. Presence of Al³⁺-O²⁻ bonds are also observed near 805.5 cm⁻¹. Minor stretching vibrations of Fe-O is also observed in the region around 440 cm⁻¹.

E. BET Analysis:

The sample is outgassed at a temperature of 150°C for about 1.5 hours. The surface area observed is 32.20 m²/g and the pore volume being 0.175 cm³/g. Thus the analysis suggests that the surface area of the RM sample is quite descent to make it use as a reactant or a catalyst till some extent.

III. CONCLUSIONS

In this paper using the PSA analysis, the average particle of the RM sample is found to be 12.106µ. The XRD results shows the presence of certain compounds which are again confirmed by the EDX and FT-IR analysis. The FESEM image shows the morphological structure of RM where the particles are relatively loose or particles are poorly crystallized and with high porosity. The BET analysis shows that the RM sample has a descent surface area. All the obtained results will provide an important base for the further studies of comprehensive utilization of RM.

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