

Green Nanotechnology: Harnessing Rice Husk Ash for Nano-Silica and Characterization Insights

ABSTRACT

Aim: To synthesize and characterize amorphous silica nanoparticles from rice husk ash using an eco-friendly chemical treatment method.

Place and Year of Study: The study was conducted in the Department of Soil Science and Agricultural Chemistry, Tamil Nadu Agricultural University, Coimbatore- 641 003, in 2023.

Methodology: Silica nanoparticles were extracted from rice husk ash using acid-leaching and sol-gel methods and characterized using X-ray diffraction analysis (XRD), transmission electron microscopy (TEM), Fourier-transform infrared Spectroscopy (FTIR) and Scanning Electron Microscopy (SEM).

Results: Based on the XRD analysis and TEM, it was observed that both samples of silica had amorphous behavior, evident from the electron diffraction patterns. The FTIR spectra mainly consisted of siloxane and silanol groups. The SEM images showed the presence of primary nanoparticles with secondary microparticles, possibly due to their agglomeration.

Conclusion: Silica nanoparticles were successfully extracted from rice husk ash, making them applicable in fields such as microelectronics, sensors, and nano-additives.

Keywords: *Nanosilica . Rice husk ash . Amorphous silica . Silanol . Siloxane*

1. INTRODUCTION

Due to its numerous uses in solar panels, concrete, insulators, refractory ceramics, and other products, silica-based nanomaterials have recently attracted attention [1]. Previous publications have described the preparation of silica materials utilizing a variety of extraction procedures, including solution precipitation, sol-gel processing, thermochemical redox reaction, and vapour-phase reaction [2, 3].

Tetraethyl orthosilicate (TEOS), sodium silicate, and silicon alkoxide are typically utilized as precursors of silica for mass manufacture. At high temperature (1300 °C), silicon alkoxide is traditionally derived from raw silica sand via a multi-step carbo-thermal reaction method [4], while sodium silicate is extracted by heating quartz, an earth substance, with sodium carbonate [5]. Both of the aforementioned methods use a lot of energy and have environmental risks.

Additionally, these technologies produce greenhouse gases like CO₂, making them unsuitable for silica production in large quantities [6]. Furthermore, the primary raw material may become scarce if natural resources like quartz are used to produce silica. Therefore, in recent years, there has been an increase in interest among researchers in finding alternative sources of silica and ways to extract it. As a byproduct of agricultural practices, rice husk (RH) is widely accessible in the top rice-producing nations like India.

RH produces a large amount of rice husk ash (RHA) when burned as a fuel and offers extremely efficient thermal energy. It is uncommon for businesses to use RH as fuel, and when it is, a significant amount of RHA (around 20% of RH) is produced. Furthermore, disposal is the key issue for both RH and RHA, particularly for the rice milling sectors. In order to use RH/RHA for the manufacture of nanosilica, which typically resides in the amorphous phase in RH/RHA, numerous efforts have

been made. Fernandes et al. [7] and Shen [8] provided an overview of numerous techniques for extracting silica from RH/RHA and their long-term use in various industries.

Instead of sodium carbonate (Na_2CO_3), sodium hydroxide (NaOH) is preferable for the alkali- SiO_2 extraction method for producing silica from RHA because Na_2CO_3 releases an enormous amount of CO_2 during the extraction process, making the procedure unsustainable for the environment [13].

The recovered silica was then examined using XRD, FTIR, SEM-EDS and TEM to determine its Structural Composition. RHA Silica is the name given to the silica obtained from Rice husk ash.

2. MATERIALS AND METHODS

2.1. Materials

Rice Husk was collected. All chemicals like HCl, NaOH, H_2SO_4 etc. were obtained from Dept. of Soil Science and Agricultural chemistry, TAMILNADU AGRICULTURAL UNIVERSITY.

2.2. Methods

To obtain white color rice husk ash, dried black color rice husk was calcined at 550°C for five hours. Majority of the inorganic content, silica, is left over when the RHA is calcined, whereas the organic components, lignin and hemicellulose, are destroyed by heat [14]. The husk was acid leached for three hours in a magnetic stirrer with a 1 M HCl (1:10 ratio) solution. After being washed with water to remove extra acid from the surface, the acid-leached husk was dried in an oven for an entire night at 80°C .

To extract the sodium silicate from the rice ash, dried rice husk ash was combined with 2N NaOH solution (1:8) and agitated in a magnetic stirrer for 120 minutes. To test for gel formation, the mixture was titrated against 0.6 M H_2SO_4 . After gathering the solid material in the centrifuge tube, it was centrifuged at 6000 rpm for 3 minutes while being washed four times with alcohol and twice with water. The solid was calcined at 400°C to 440°C for four hours after being dried in an oven at 80°C overnight.

2.3. SILICA RECOVERY

The recovery of silica from RHA was performed in triplicate and calculated using

$$\text{Silica Yield \%} = (\text{Mass of Silica obtained} / \text{Mass of RHA}) * 100$$

2.4. CHARACTERIZATION STUDY

RHA-Silica's microstructural, morphological, and compositional analysis were carried out using XRD, FTIR, SEM as material characterization techniques.

3. RESULTS AND DISCUSSION

3.1. XRD Characterization

RHA-Silica's XRD pattern is shown in Fig 1. For the silica sample, a large hump at a 2θ angle of 27° was seen, demonstrating the sample's amorphous composition. Ma et al. [6] and Yalc and Sevinc [9] also got similar results when they isolated silica nanoparticles from rice husk and found a large peak centered at a 2θ angle of 27° . Other scanning angles from 5° to 80° showed no strong peaks, indicating the absence of any organized crystalline structure. According to An et al.'s [15] analysis on silica's very high disordered particle shape, this also showed that. Additionally, the Bragg diffraction peaks' Gaussian peak fitting was carried out. Broader de-convoluted peaks were produced after the entire Bragg diffraction peak set was fitted with Gaussian functions (Fig. 1). This finding demonstrates that the sample has the normal amorphous components but no organized crystalline phases. Sampath et al.'s [10] use of the Gaussian function for an X-ray diffraction analysis of amorphous and nanocrystalline structure revealed that a larger Gaussian peak denotes the presence of an amorphous component.

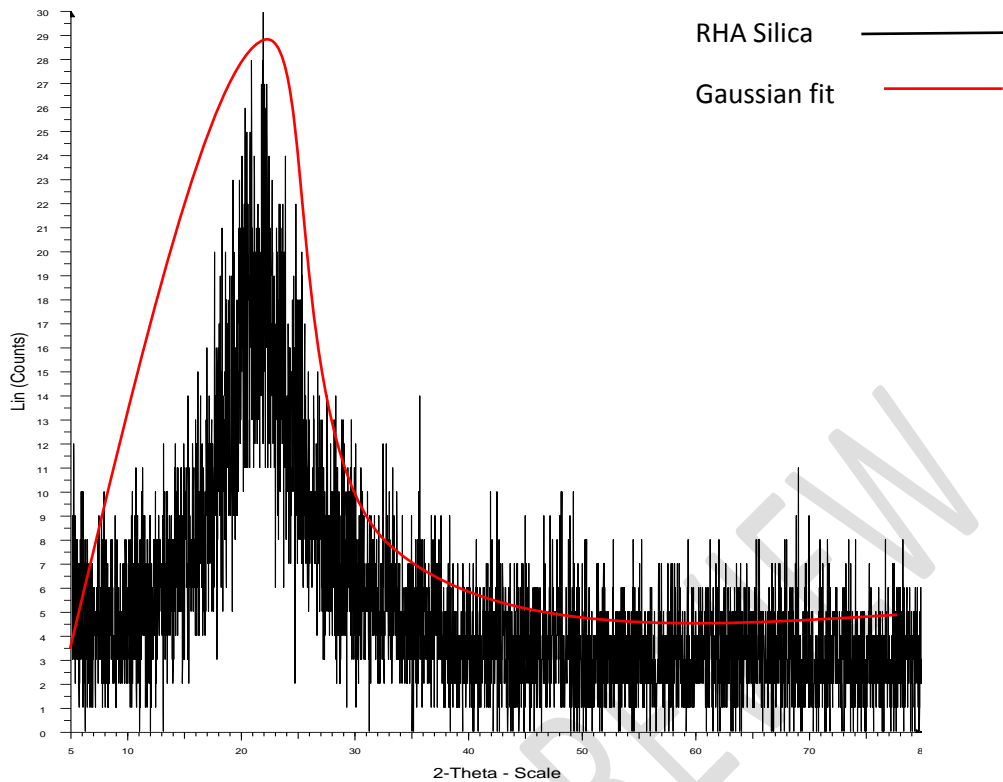


Fig 1. XRD of RHA Silica

3.2. FTIR Characterization

While bands at No.6 with position at 1640.16 for RHA-Silica sample is due to bending vibration of H₂O molecule in the Si-OH group, bands at No.2 with position 3416.28 of RHA-Silica are attributed to stretching vibration of O-H group, respectively [16]. The Si-O-Si asymmetric stretching vibration was shown by the bands at No.7 with Position 1100.19. The Si-O bond's symmetric stretching vibration caused the wavenumber of No.10 to have band position 616.145. In contrast, the Si-O bending of the siloxane group for RHA-Silica caused the wavenumber of No.11 to have band position 468.617.

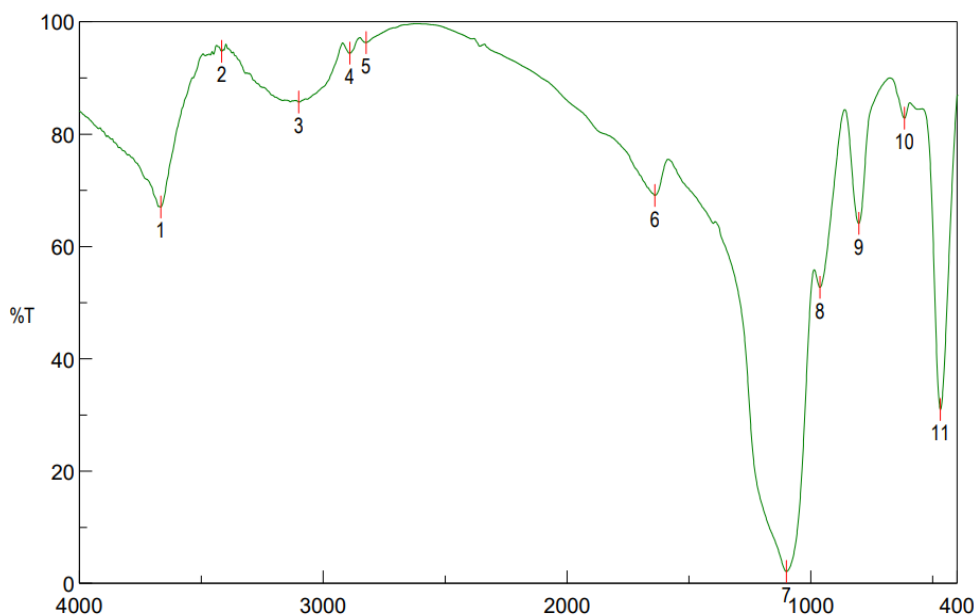


Fig 2. FTIR Spectra of RHA Silica

Chart 1. The position and intensity

No	Position	Intensity
1	3666.02	66.985
2	3416.28	94.7105
3	3100.01	85.6836
4	2890.77	94.3866
5	2824.24	96.281
6	1640.16	69.0455
7	1100.19	2.0944
8	962.305	52.7116
9	803.206	64.0635
10	616.145	82.8119
11	468.617	30.9392

Table 1. Absorption bands of silica sample

Band origin	Position/cm ⁻¹	RHA Silica
Bending vibration of Si-O	Siloxane group	468.617
Symmetric stretching vibration of Si-O	(Si-O-Si)	616.145
Asymmetric stretching vibration of Si-O-Si		1100.19
Absorption of single broad band	Silanol group	-
Bending vibration of H-O-H	(Si-OH)	1640.16
Stretching vibration of O-H		3416.28

3.3. SEM, EDX and Particle size characterization

SEM micrographs **Fig 3a**, particle size histograms **Fig 3b** and EDX peaks **Fig 3c** are showed. Spherical SiO_2 -NPs were observed with uniform surface morphology in RHA-Silica along with agglomeration. The agglomeration effect may be due to the dominance of strong cohesive intramolecular forces instead of gravitational force [11]. The elemental composition analysis was performed through EDX study and the results are presented in Table 2. It was observed that only two strong peaks of O K α and Si K α were present in both RHA-Silica. The results obtained are in accordance with the results of Mourhly et al. [12], who worked on mesoporous nano-silica. From particle size distribution histograms, it was noticed that average particle size of the SiO_2 -NPs extracted from RHA was 17.80 ± 10.10 nm. However, it is worth mentioning here that a few secondary micro-particles were also observed may be due to the agglomeration effect.

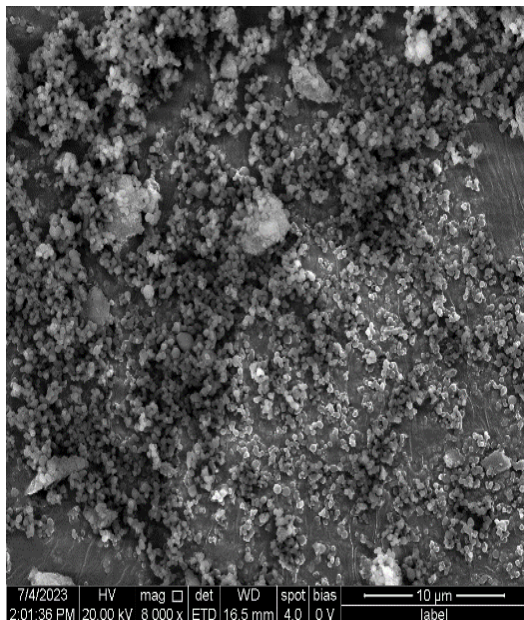


Fig 3a. SEM image

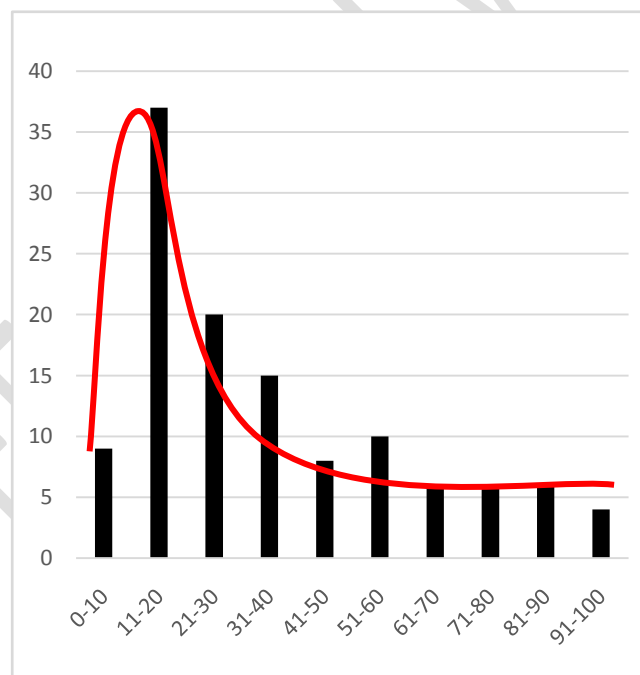


Fig 3b. Particle size

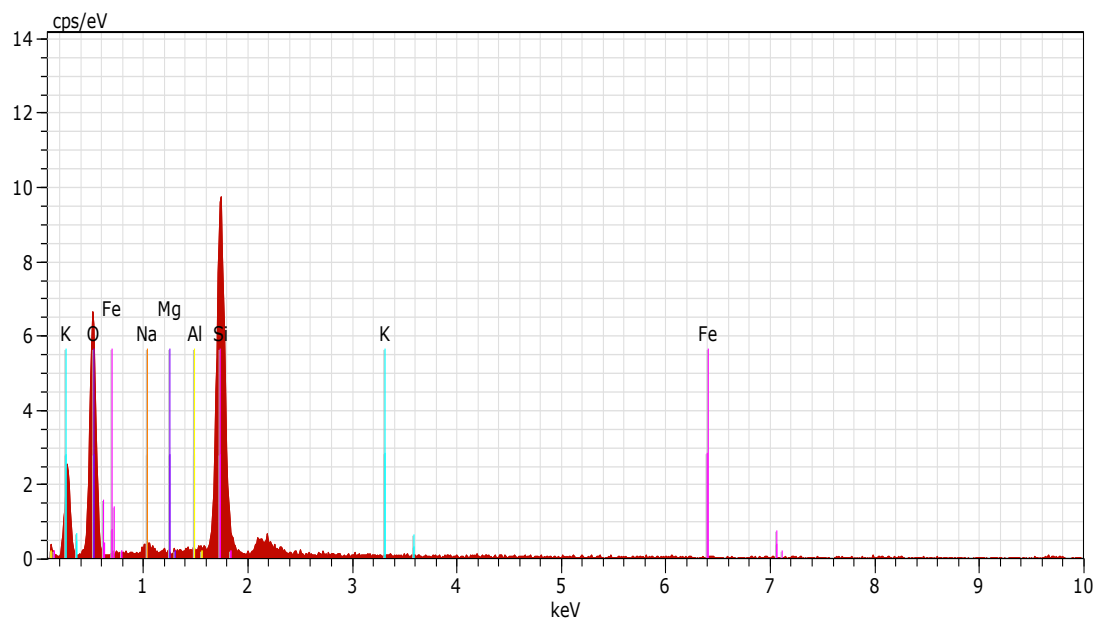


Fig 3c. EDX

Table 2. Elemental composition of silica obtained from EDX analysis

Elements	RHA Silica	
	Weight %	Atomic %
O k	63.7775.35	
Si k	32.77	22.05
Na k	2.04	2.04
Al k	0.46	0.32
Fe k	0.34	0.11
Mg k	0.12	0.09
K k	0.06	0.03
Total	100	100

3.4. TEM Characterization

In **Fig 4.** The microstructures of silica samples were also verified through FETEM and the TEM image clearly showed spherically shaped nanoparticles (size <200nm) with high agglomeration.

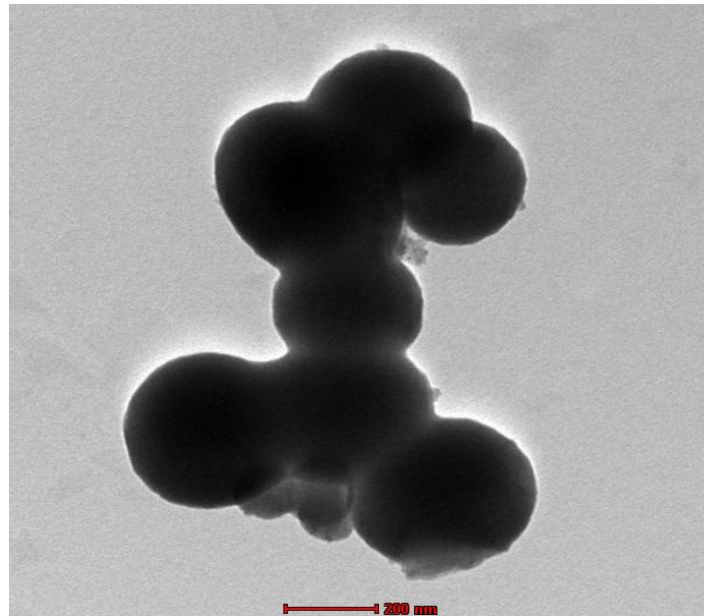


Fig 4. TEM Image of RHA Silica

4. Conclusion

According to the research, silica extraction (SiO_2) utilizing the acid-leaching and sol-gel methods with rice husk base material was accomplished satisfactorily. The crystal structure of extracted silica is amorphous. A possible green resource is rice husk. The suggested process for making nano silica from rice husk is straightforward, suitable for large production, and reduces waste disposal issues.

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