Introduction

The project work was carried out with an objective to extract silica nanoparticles from rice husk ash (RHA), the silica content in rice husk ash and to prepare composite using extracted silica nanoparticles to find out whether nanoparticles affect in mechanical properties of composite or not. Nanoparticles are nanomaterials having all three dimensions in the range of 1-100 nm [3]. All nanoparticles are nanomaterials but not all nanomaterials are nanoparticles. Nanomaterials are materials having at least one of their dimensions in the range of 1-100 nm. This range is called nanoscale.

Rice is the second most widely consumed food item globally, with rice paddy production registering about 758 million metric tons in 2070, a number that will increase gradually due to the projected demand of the world population [4]. The RHA is an agricultural waste, which is generally used as ameliorants to break up clay soils and improved soil structure but is also used for the production of silica. The properties of RHA depend on the ecological circumstances of its origin as well as the process applied for burning the husk [5]. Silica which is the constituents of silicon and oxygen has the molecular formula SiO$_2$. Silica has been widely used in vegetable oil refining, pharmaceutical products, detergents, adhesive, chromatograph column planning and ceramics [6-8]. Not only this, silica is also found to be used in water purification, vulcanizing rubber, control of insect pests, in stored food stuffs, manufacturing refractory bricks, and as admixture in low cost concrete block manufacturing [9].

Abundantly available waste RHA has been widely used as raw materials for the preparation of silica gel...
and powder following (Kamath and Protocor, 1998) [1].

Nanoparticles exhibit attractive properties such as low thermal expansion, high mp, high thermal stability, high thermal resistance and chemical stability [10,11]. The interfacial area that creates a significant volume fraction of the interfacial polymer with the properties different from bulk polymer even at low filter loadings [12,13]. Several researchers have studied the effect of particle size and volume fraction on the mechanical response of composite [14-18]. However, among the numerous polymer composites; silica polymer nanocomposites are most commonly reported in the literature and are also employed in variety of applications, such as electronics, automotive and aerospace industries as well as used in many industrial products due to their good mechanical characteristics [19].

Ceramics is defined as the solid materials that are not metal, plastic, or derived from plants or animals [20]. Ceramics have some distinctive properties including good chemical inertness, high temperature stability brittleness, high mp and an electrical insulation capability [21]. Furthermore, Chemical bonding (both covalent and non covalent) between filler and a polymer improves materials compatibility, and thus enhances certain properties of matrix above and beyond what is accomplished by single doping with filler [22-24]. Using similarly sized dopant and matrix elementary building blocks may have certain distinctive advantages [25].

Materials and Methods

Chemicals used: Distilled water, 1N NaOH, 1N HCl

Apparatus used: Beaker, conical flask, burner, wire gauze, magnetic stirrer, filter paper (Whatmann 41), hot air oven

Preparation of chemical reagents

Preparation of 1N NaOH

Mass of NaOH required (w) = NEV/1000

Where, N= Normality, E= Equivalent wt.,

V=Volume required in mL

For 600 mL, mass of 1N NaOH required was 24 g. It was taken in a volumetric flask and little amount of distilled water was added to dissolve it. After it got fully dissolved, distilled water was added up to the mark of 600 mL.

Preparation of 1N HCl

As concentrated HCl has approximately strength of 36%, its normality calculated to be 11.6 N by using the following formula:

\[
\text{Normality} = \frac{\text{(% by mass x specific gravity x 10)}}{\text{Equivalent weight}}
\]

Where, % by mass = 36

Specific gravity = 1.1789

Equivalent weight = 36.5

For 200 mL of 1 N dilute HCl, 17.24 mL of concentrated HCl was added up to 200 mL of distilled water. It was calculated by the following way:

\[
(\text{Conc. HCl})_1 V_1 N_1 = V_2 N_2 (\text{1 N Dil. HCl})
\]

Where, \(V_1\) = Volume of concentrated HCl to be calculated

\(N_1\) = Normality of Concentrated HCl (11.6 N)

\(V_2\) = Volume of dilute HCl to be prepared (200 mL)

\(N_2\) = Normality of dilute HCl (1 N)

Preparation of rice husk ash from rice husk

Rice Husk was collected from the nearest Rice milling industry, which was then burnt and converted into ash. The ash converted was then made into fine powder.

Extraction of silica from rice husk ash

This was done by Sol-gel method. In this method, 50 g RHA was weighed and dispersed in 300 mL of distilled water in a beaker. The pH of solution maintained at 7 by testing with litmus paper. The solution was then stirred by magnetic stirrer for 2 hours. It was then filtered by Whatmann 41 filter paper and the residue was washed with 500 mL of water. The residue was then dispersed in 300 mL 1N NaOH and boiled with constant stirring for 1 hour. The filtrate was collected and allowed to cool. Then, the filtrate was titrated with 1N HCl till the pH of filtrate became neutral.

When pH of filtrate became 7, gel obtained. The gel was allowed to age for 18 hours. After 18 hours, 500 mL of distilled water was added into volumetric flask containing gel and crushed the gel. After crushing the
gel, the solution was left for some time, later which two layers were seen. Upper layer comprised of water called supernatant and lower layers consisted of gel.

The supernatant was discarded by the help of dropper and the washing process was repeated for 3 more times. The gel obtained after removing supernatant was kept in an oven for 18 hours at 80 degree Celsius. The gel produced called xerogels, which was then kept in air tight plastic bottle.

The method applied following (Kamath and Protocor, 1998) [1] can be explained step wise as:

1. Dispersed RHA (50 g) in Distilled water (500 mL) → Adjusted pH (7)
2. Filtrate (residue 40 g) → Stirred (1 hr)
3. Washed residue with water (500 mL) → Dispersed residue in 1 N NaOH (500 mL)
4. Filtrate (residue 40 g) → Boiled with stirring (1 hr)
5. Washed residue with boiling water (500 mL) → Filtrate collected and allowed to cool
6. Allowed the gel to age for 18 hrs → Titrated filtrate with 1N HCl (pH 7)
7. Added water (500 mL) and crushed the gel → Centrifuged
8. Discarded the supernatant and repeated the washing
9. Dried at 80 degree Celsius
10. Extracted silica (16 g)

Results and Discussion

FTIR analysis

The FTIR of silica nanoparticles shows various peaks as a result of the absorption of light of a wavelength in infra red region. From the graph of Transmittance% vs. wave number per cm, it is cleared that rocking vibration occurs at 800 cm\(^{-1}\). Similarly; Si-O asymmetrical stretching vibration occurs at 1068 cm\(^{-1}\).

<table>
<thead>
<tr>
<th>S. N.</th>
<th>Wave number cm(^{-1})</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.</td>
<td>800</td>
<td>Rocking vibration</td>
</tr>
<tr>
<td>2.</td>
<td>1068</td>
<td>Si-O-Si asymmetrical stretching vibration</td>
</tr>
</tbody>
</table>

Preparation of ceramics matrix nanocomposites (CMNs)

The CMNs, silica-sand cement block was prepared. For this purpose, the extracted silica from RHA was mixed to the sand cement mixture, which is believed to increase the mechanical property. Firstly extracted silica, sand and cement were mixed in the porcelain basin in the ratio of 1:4:1. Then, distilled water (generally 26-33% of the total weight of mixtures) was added on it till the required texture was obtained. After then, it was framed in a 2 cm cube mold and the curing process was done by immersing the block in distilled water. As 3 days strength test was followed, the blocks were removed from distilled water after low-cost days and allowed it to dry, free from any moisture. The compressive test of the block was then done.

Ceramics composites

The preparation method was the same as that of CMNs, silica-sand cement block. The only difference is that silica nanoparticles were not added to cement and sand.

In this process, sand and cement were taken in the
XRD analysis

Confirmation of the effectiveness and the size of silica nanoparticles were provided by X-ray studies (XRD). The center of peak and the value of FWHM required to calculate the size of nanoparticles was determined by using origin software.

On the basis of X-ray curves, Lorentzian fitting was done. It is intended to a one-mode system and believes to describe a decaying system over time. The data of XRD (two theta degree value vs. intensity) was copied into origin software. Then, two strong peaks were selected efficiently and analyzed in multi-peaks fitting. As mentioned above, Lorentzian fitting was done which then by double-clicking on the center of peaks gave the value of angle of the center of peak (2θ) and the FWHM value.

The center of peak and FWHM value on using origin software found to be 32.10910 and 0.37580. The value of θ calculated by dividing the center of the peak by 2 and FWHM value was then converted into radian which found to be 6.55×10^-3. The wavelength used was 1.5406Å and Scheerer's constant value (κ) used was 0.94. Thus, by using Scheerer's formula, the size of the particle was determined.

\[ D = \frac{K\lambda}{\beta \cos\theta} \]

Thus, putting all the values, the particle size was found to be 23 nm.

Compressive test

At first, the cross-sectional area and the breaking load of both the blocks, ceramics matrix nanocomposite (containing silica nanoparticles), and another ceramics composite (without silica nanoparticles) were measured and their value was found to be:

<table>
<thead>
<tr>
<th>Block</th>
<th>Cross-sectional Area(cm²)</th>
<th>Breaking load(N)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ceramics Matrix Nanocomposite</td>
<td>4</td>
<td>40</td>
</tr>
<tr>
<td>Ceramics Composite block</td>
<td>4</td>
<td>24</td>
</tr>
</tbody>
</table>

Thus, by putting the value in a formula,

\[ \text{Compressive strength} = \frac{\text{Breaking load}}{\text{Cross-sectional area}} \]

It was found that the compressive strength of block containing silica nanoparticles (ceramics matrix nanocomposite) was 10N/cm² and 6 N/cm² for the block having no silica (ceramics composite).
Conclusion

Silica nanoparticles were synthesized from Rice Husk Ash (RHA) by using the sol-gel method. The synthesized silica nanoparticles found to be 26 g (52%), which is almost equal to 60% and seems to follow the citation [1,2]. The silica nanoparticles were characterized by spectroscopic techniques like FTIR and XRD. As per the result of XRD, the particle size was found to be 23 nm. By analyzing the data of FTIR, rocking vibration, and Si-O-Si asymmetrical stretching vibration was found. Similarly, synthesized silica nanoparticles were then incorporated in sand cement mixture in 1:4:1 ratio to prepare ceramics matrix nanocomposite block. In the same way, ceramics composite block was also prepared where silica nanoparticles were not incorporated in the sand-cement mixture. The mechanical property of prepared blocks was characterized by compressive strength test using the Instron testing machine. As per the result of compressive test, the ceramics matrix nanocomposite block containing the silica nanoparticles found to have almost 1.67 times greater strength than the ceramics composite block containing no silica. This proves that silica nanoparticles enhances the mechanical strength and hence it can be used in several construction materials.

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References

2. A. Chakraverty and S. Kalemullah, Conversion of


