

CORROSION BEHAVIOUR OF ALUMINIUM 6061 NICKEL COATED CENOSPHERE COMPOSITE

Arvind R S¹, Prasanna Ram M¹, *Prashanth T¹, Jaimon Dennis Quadros²

¹Department of Mechanical Engineering, Global Academy of Technology, Bangalore, Karnataka-560098, India ² Department of Mechanical Engineering, Birla Institute of Technology, RAK- 41222, United Arab Emirates https://doi.org/10.37255/jme.v15i3pp54-59

ABSTRACT

Cenosphere fly ash is one of the most inexpensive and low-density material which is abundantly available as a solid waste by-product of coal combustion in thermal power plants. Aluminium metal matrix composites with Nickel coated cenospheres as the reinforcement is prepared by stir casting route. The composites are prepared with varying percentages of cenospheres in the percentage of 2-10% by weight of the composite. Immersion corrosion tests are conducted on the composites in three different medium and for three different time durations. It is evident from the test results as well as the microstructure images that the weight loss of samples with 8% Nickel coated cenospheres has shown least corrosion or the highest corrosion resistance when compared to the counterparts.

Keywords: Aluminium 6061 composites, fly ash cenospheres, Nickel coated cenospheres, corrosion resistance, immersion corrosion.

1. Introduction

Aluminium and its alloys are of high specific strength and low density. Therefore, they have a wide range of application in the automotive and aerospace industries [1]. Metal matrix composites (MMCs)are gaining widespread popularity in several technological fields owing to its improved mechanical properties when compared with conventional metals/alloys [2]. The reinforcements are generally used to improve the base metal properties such as strength, stiffness, conductivity, etc. [3]

India is the second-largest producer of coal in the world, generates copious amounts of fly ash. Since fly ash is a very light particle, it needs to be stored under an artificial storage pond; therefore, the demand to determine the usage of fly ash in metal matrix composites is increasing.

Though considered as a waste, cenosphere is a useful by-product to be used to produce newer materials otherwise cheaply which poses significant environmental and disposable problems. Fly-ash cenospheres have unique properties such as low density, non-toxic, non-metallic hollow microparticles and are light in weight. The particle size of cenospheres varies from 1 to 20 µm and their mean particle diameter in the order of 8µm. Cenosphere has a size range from 1 to 500 microns with an average compressive strength up to 7000 psi. Their colour ranges from white to dark grey. [4]

Cenosphere reinforced aluminium enhances the chemical stability of aluminium-cenosphere composites during synthesis and reheating [5]. The weight loss of the MMC material followed descending order as percent filler increased till 6% cenosphere, further increase of cenosphere resulted with an increase in weight loss for both heat-treated and untreated composites [6]. Cenospheres as filler in aluminium casting reduce cost, decreases density and increases hardness, stiffness, wear and abrasion resistance [7]. They are considered as potential materials for components like pulleys, oil pans, intake manifolds and valve cover [8]. The use of Aluminium-fly ash composites reduces emissions like CO2, which otherwise come from aluminium production. In transporting the components, low weight materials consume less energy benefiting the environment [9] Zhu and Hihara [10] have reported on the corrosion performance of a continuous alumina-fibre reinforced metal-matrix composite (MMC) and its monolithic matrix alloy (Al-2%Cu-T6) in 3.15wt% sodium chloride solution.

Nunes et al [11] have studied the corrosion behaviour solution of alumina-aluminium and SiC-Al in sodium chloride. Immersion and anodic polarisation corrosion tests have been carried out. It is reported that composites have exhibited lower corrosion resistance when compared. Nickel and copper coated reinforcements have produced beneficial effects in particular to reduction in porosity level and reduce

*Corresponding Author - E- mail: prashanth.t@gat.ac.in

interfacial reactions during the processing of metal matrix composites [12-14].

Currently, electroless deposition of metallic coatings on the ceramic reinforcement is gaining popularity owing to its advantages such as uniformity in coatings over the surfaces regardless of size and shape. Further, this technique is autocatalytic, which means no conduction surfaces are required. The published literature on advanced materials, such as Aluminium nickel-coated composites, is somewhat limited and is primarily concerned with applications of fly ash particles for the synthesis of these materials. Therefore, it was thought worthwhile to study the corrosion behaviour of this composite as well as present the pitting morphologies of the corroded surface. The present work is dedicated to such an investigation.

2. Experimental Procedure

A batch of 3.5kgs of Aluminium 6061 alloy was melted using a 6KW electric furnace. The melt was degassed using commercially available chlorine-based tablets (Hexachloroethane). The molten metal was agitated by use of mechanical stirrer rotating at a speed of 300 rpm to create a fine vortex. Preheated cenospheres (preheated to 2000C for 2 hrs.) were added slowly into the vortex while continuing the stirring process. The stirring duration was 10 min. The composites melt maintained at a temperature of 710°C was then poured into preheated metallic moulds. The stirrer blades used were made of stainless steel and were coated with a ceramic material to minimise the iron pickup by the molten metal. The amount of cenospheres was varied from 2 to 8 wt. % in steps of 2%. [15].

2.1 Immersion corrosion test

Immersion test were carried out as per ASTM G31 test procedure. Polished samples of Al 6061 alloy and all the composites were immersed in 3.5% NaCl solution for a total duration of 25 days. Weight loss measurements of the samples were done at the end of every 5 days. Corroded surfaces of the samples were cleaned with acetone before weighing using an electronic balance of accuracy of 0.001 grams. The corresponding changes in the weights were noted. The corrosion rate was calculated using the following relationship.

Corrosion rate (mpy) \rightarrow 534*weight loss / (density*area of exposure*time of exposure) (1)

A single step activation and sensitisation step was carried out. We also used Sodium Hypophosphite to reduce Nickel in the presence of PdCl2. The composition of the chemicals used for the coating process is as shown in Table 1.

NiSO4 +2NaH2PO2+2H2O \rightarrow Ni+2NaH2PO3- +H2+ H2SO4 (2)

The nickel reduction only takes place on specific catalytic surfaces, including Nickel itself, which makes the reduction process autocatalytic as shown in Equation 2. The reducing agent (PdCl2) used for electroless plating not only supplies the electrons for the reduction, but some elements in the reducing agent can be incorporated into the nickel deposit improving its properties e.g. when sodium hypophosphite is used as a reducing agent the resultant deposit is a nickelphosphorus alloy.

During the plating process, the nickel sulfate and sodium hypophosphite raw materials are continuously depleted and must be replenished in order to maintain the chemical balance of the bath and addition of either ammonia or sodium hydroxide are necessary to keep the pH in the preferred range. This results in an accumulation of sodium and sulfate ions, along with orthophosphate, as the electroless nickel bath ages and consequently, the plating rate can decrease from 18 mm hr-1 to less than 10 mm hr-1

 Table 1 Bath composition for electroless nickel

 deposition

Bath constituents	Quantity (g/litre)
Nickel Chloride	30
Sodium acetate	10
Sodium Hypophosphite, monohydrate (NaH2PO2, H2O)	20
Glycin (H2NCH2COOH)	10
Sodium hydroxide	0.02

2.2. SEM studies

Scanning electron microscope studies were carried out using JSM 840a Joel scanning electron microscope on Aluminium 6061, Aluminium cenosphere composite and aluminium nickel-coated cenosphere composite at Indian Institute of Science (IISc), Bangalore. The sizes of the specimens were machined to 20mm (diameter)

3. Results and Discussion

Fig.1 to Fig. 5. shows the SEM micrographs of Nickel coated cenosphere Al 6061 composites. From the micrographs, it is observed that the Nickel coated cenosphere particles are dispersed uniformly throughout the matrix. Further, SEM images clearly indicate that

reinforcement particulates are well bonded to the aluminium matrix and that there exist very minimum micro porosities in all the composites. This fact supports the good bond between the matrix and reinforcement. [16-17].



Fig. 1 Al 6061 (200x)

The few pores observed in the as-cast composite were mostly associated with the reinforcement particles and particle clusters. The latter feature was especially pronounced in the composite with a higher weight fraction of the reinforcement in the composites. The extent of distribution of reinforcement particle in the composites is more homogeneous when compared with cast composites, suggesting that rearrangement and recrystallisation had taken place.



Fig. 2 Al 6061- 2% Nickel coated cenosphere



Fig. 3 Al 6061-4% Nickel coated cenosphere



Fig. 4 Al 6061- 6% Nickel coated cenosphere



Fig. 5 Al 6061-8% Nickel coated cenosphere

3.1 Effects of Reinforcement

Fig. 6. shows the effect of reinforcement on corrosion rate in mills per year of Al 6061 alloy and Al 6061-nickel coated composites after 60 days of immersion in 3.5%NaCl solution. It is observed that the corrosion rate of Al 6061 alloy decreases with increase in the percentage of Nickel coated cenospheres as reinforcement. This can be attributed to the fact that the coating of Nickel on the cenosphere provides effective barriers for the corrosion to occur. It is therefore observed that the composite containing 8% by weight of Nickel coated cenospheres has shown better corrosion resistance when compared to its counterparts.



Fig. 6 Variation of corrosion rate of Al 6061 – nickelcoated composite cenospheres after 10 days of immersion in 3.5%NaCl solution

3.2 Effect of immersion duration



Fig. 7 Variation of corrosion rate in mpy of Al 6061 matrix alloy and Al 6061-nickelcoated cenosphere composites

Fig. 7 shows the corrosion rate in mpy as a function of immersion time in the number of days in 3.5% NaCl for Al 6061 matrix alloy and Nickel coated cenosphere composites. It is observed that the corrosion rate drops drastically in the composites studied. It is further noted that the corrosion rate of Al 6061 matrix alloy and its composite systems varies in a narrow band. This can be attributed to the facts as discussed in the earlier section

3.3 Pitting morphology

Fig. 8 and Fig. 9 shows the SEM photographs of the corroded surface of Al 6061 matrix alloy and Al 6061-nickel coated cenosphere composites, respectively. It is observed that the developed composites possess a lesser number of pits when compared with the Al 6061 matrix alloy. Bhat et al. [18] have observed a similar trend in his study on aluminium matrix composites



Fig. 8 Al 6061 alloy (200x)

Fig. 10 shows the SEM photographs of the corroded surface of Al 6061-8 wt% Nickel coated cenosphere composites for varying time durations. It is observed that the damage on the composite decreases with an increase in the percentage of reinforcement. However, increase in time duration has led to the formation of a passive layer on to the surface of the composite material leading to decrease in the corrosion rate days which is well supports the obtained results discussed in earlier section.



Fig. 9(a) Al 6061-4wt% nickel coated cenosphere(200x)



Fig. 9(b) Al 6061-8 wt% nickel coated cenosphere(200x)



Fig. 10(a) 5 days (200x)



Fig. 10(b) 15 days (200x)



Fig. 10(c) 20 days (200x)

Fig. 10 SEM photographs of corroded surfaces of Al 6061-8 2wt%nickel coated cenosphere composites for different time durations

4. Conclusions

- i. Al 6061-nickel coated cenosphere composites have been successfully produced by liquid metallurgy route.
- ii. Al 6061-nickel cenosphere composites (8%) possess inferior corrosion resistance in 3.5%NaCl medium when compared with Al 6061 alloy.
- iii. It is observed from the pitting morphology studies that the pits are deeper in the reinforced composites as compared to the unreinforced counterparts.
- iv. The studies clearly show that the reinforcement has, in particular, a higher role to play in the improvement of corrosion resistance of the composite.

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