# Unlocking the Potential of *Phaseolus Vulgaris* Particulates on AA6063 Alloy for Enhanced Mechanical, Corrosion and Opto-electrical Performance

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#### Abstract

AMMC are notable for their extensive research and unique property profiles, surpassing those of traditional solid metals. Their remarkable strength sets them apart, making them highly sought-after and versatile across diverse industries. The integration of reinforcements into AMMC boosts their properties, including increased tensile and compressive strength, improved wear resistance, reduced thermal expansion and enhanced hardness, characteristics unattainable with conventional materials. However, the use of CMC, such as SiC and Al<sub>2</sub>O<sub>3</sub>, escalates the production cost of AMMC, thereby limiting their application in various industries. In recent times, researchers have increasingly turned to natural reinforcements derived from waste sources, such as bamboo leaves, rice and palm kernel shells. However, investigations into PV potential as reinforcements for AMMC have been limited, prompting the initiation of this study. This research aimed to examine mechanical, microstructural, electrochemical and optoelectronic properties of AMMC reinforced with PVP. The study demonstrated an increase in hardness values, from 28.1 to 38.5 HRB, with 25% PVP reinforcement, and a significant reduction in CR, from 12.06 to 6.13 mm/yr. Electrical resistivity and conductivity measurements showed variations across composite samples, while microstructural analysis confirmed uniform PVP dispersion and integration within the metal matrix, alongside additional crystalline formations.

Keywords: Al; corrosion; HRB; optoelectrical; PV; PVP; reinforcement.

#### Introduction<sup>•</sup>

Many engineering materials in today's world require properties such as increased strength, weight reduction and cost-effectiveness, particularly in terms of energy usage. The search for such materials arises from the aim to produce lightweight structural elements and engine parts with high strength-to-weight ratios, hence reducing excessive fuel consumption and CO<sub>2</sub> emissions [1-4]. Due to rigorous demands of engineering applications, materials must exhibit a range of complex

<sup>•</sup>The abbreviations list is in page 361.

properties, including high strength, exceptional wear resistance, improved corrosion resistance and other desirable mechanical, electrical and chemical characteristics. However, it is rare to find all these properties in a single metallic material. Researchers have found that achieving such desirable features is only possible through merging different materials, to create a combination with enhanced properties [5-7].

The combined materials are commonly known as composites, which have the unique potential to improve selected characteristics of monolithic materials, while remaining superior to their counterparts. They are generally classified according to the composition of their base matrix phase: CMC, MMC and PMC [8-12]. MMC are highly engineered materials renowned for their ability to blend diverse properties by combining matrices with hard particulate reinforcements, thereby attaining substantial engineering and industrial importance compared to traditional metallic alloys. Within the spectrum of MMC, AMMC are notable for their extensive unique property profiles, surpassing those of traditional solid metals. Their remarkable strength sets them apart, making AMMC highly sought-after and versatile across diverse industries [13-15].

AMMC efficiency relies on various factors impacting their properties, such as type, volume and weight fraction, as well as particle size, distribution and shape of reinforcements. Additionally, production method employed also significantly influences overall effectiveness of composites [16-19]. Research of those factors reveals that reinforcements play a key role in sustainable AMMC. In manufacturing, AMMC, such as Al-Si, Al-Cu and Al-Si-Mg, serve as MMC, while reinforcements like SiC, Al<sub>2</sub>O<sub>3</sub>, C, B, B<sub>4</sub>C, AlN, SiO<sub>2</sub> and BN are incorporated into them. Specifically, the integration of reinforcements into Al matrixes enhances their properties, including increased tensile and compressive strength, improved wear resistance, reduced thermal expansion and greater hardness, characteristics unattainable with conventional materials [20-22]. However, the use of CMC such as SiC and Al<sub>2</sub>O<sub>3</sub> escalates the production cost of AMMC, thereby limiting their application in various industries [23].

Thus, researchers have been investigating alternate reinforcement materials with the aim of cutting AMMC production costs. These studies have revealed an array of naturally occurring particles, including agro-waste and industrial by-products, showing promise as feasible alternatives to typical reinforcements in AMMC. Given its abundance and economic viability, agrowaste has emerged as an especially favourable reinforcement material. Extensive research efforts on diverse agricultural by-products have revealed their intrinsic richness in Si and Mg, making them potential reinforcing possibilities [24]. The realm of agrowaste presents a promising avenue for exploring particulate reinforcements. It offers a dual benefit of addressing pollution challenges and enhancing the strength of metal matrices. In recent times, researchers have increasingly turned to natural reinforcements derived from waste sources such as bamboo leaves [25-28], rice husk [29-31] and palm kernel shells [32-36]. However, investigations into the potential of PV shells as reinforcements for AMMC have been limited, prompting the initiation of this study. Thus, the primary objective of this research was to examine mechanical, microstructural, electrochemical and optoelectronic properties of AMMC reinforced with PVP.

### **Research methodology**

Al alloy (AA6063), which was sourced locally from the metal market, served as base material. PVP were prepared by cleansing them with acetone and warm water, air-drying for 48 h, crushing, milling and sieving, to achieve the desired particle size of 45  $\mu$ m. Final particulate size can be observed in Fig. 1.



Figure 1: Photo of PVP sample.

Chemical compositions of AA6063 and PVP, after characterization, are detailed in Tables 1 and 2, respectively.

Table 1: Chemical composition of AA6063.

Table 1. Chemical composition of 74 (0005).										
Elements	Fe	Cu	Mg	Mn	Si	Ti	Zn	Cr	Al	
%Wt.	0.11	0.01	0.79	0.02	0.88	0.008	< 0.01	< 0.01	Bal.	

Table 2: Chemica	l composition	of PVP.
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Compound Ct	CaO	Al <sub>2</sub> O <sub>3</sub>	K <sub>2</sub> O	P <sub>2</sub> O5	SiO2	MgO	Fe <sub>2</sub> O <sub>3</sub>	MnO	Na <sub>2</sub> O
(wt%)	41.21	15.43	7.32	2.39	15.69	4.85	9.71	0.72	1.76

Stir casting technique was employed to fabricate Al metal matrix composite. This method involved uniformly mixing the reinforcement into the matrix material, using a mechanical stirrer and pouring the composite mix into the desired mould geometry. Five samples were prepared with varying wt.% of PVP (0, 10, 15, 20 and 25%), following ASTM B179 standard specifications. For mechanical assessment, HRB was measured employing an HR-150A HRB testing machine (ASTM E18-22). Samples were indented for 10 secs, at a force of 5 kg, with multiple measurements taken to ensure accuracy. Corrosion testing utilized an AUTO LAB potentiostat galvanostat (PGSTAT 101), in a 1 M HCl environment with a three-electrode polarization setup employing Ag/AgCl and graphite electrodes. Microstructural analysis was conducted using a JOEL JSM-7600F machine for SEM/EDXS, and Rigaku D/Max-IIC X-ray diffractometer for XRD experiments. Electrical conductivity and resistivity were assessed using a custom electrical setup (Fig. 2)

[37]. Fig. 3 shows the empirical electrical setup used to investigate magnetic permeability of the composite, in response to inductive change.



Figure 2: Experimental electrical setup [37].



Figure 3: Magnetic saturation test circuit.

## **Results and discussion**

## Hardness values of samples

HRB value of the developed composites is shown in Fig. 4. According to HRB readings, the samples with highest Ct of PVP showed the greatest reading and helped raise the substrate material's hardness from 28.1 to a final composite deposition of 38.5 HRB. Overall, there was an increase in hardness values for samples containing different PVP sizes, from 5.33 to 37.01%. The enhanced

strength observed in the fabricated composites can be largely attributed to robust interfacial bonding and adhesion between soft and ductile AA6063 matrix and PVP reinforcement. This bonding significantly augmented load-bearing capacity of the composite, thus minimizing matrix deformation. Additionally, uniform dispersion of reinforcements within matrix phase contributed to observed results [23, 38, 39]. However, it should not be assumed that increasing the Ct of particulate reinforcements will consistently lead to more hardness, as this is limited by the availability of intermetallic ions required for bonding with the lattice [31].



Figure 4: Hardness values of samples.

## Corrosion evaluation of samples

Table 3 indicates results of polarization test and Tafel plot in terms of CR,  $J_{corr}$ ,  $E_{corr}$  and  $R_p$ .

Table 3: Tafel plots of samples.							
Samples	E <sub>corr</sub> (V)	J <sub>corr</sub> (A/cm <sup>2</sup> )	CR (mm/yr)	Rp (Ω)			
Control	-0.383	2.01E-02	12.06	3.58E+01			
10% PVP	-0.381	1.77E-02	10.34	4.02E+01			
15% PVP	-0.394	1.57E-02	9.31	3.88E+01			
20% PVP	-0.416	1.69E-02	8.63	3.44E+01			
25% PVP	-0.442	1.75E-02	6.13	5.42E+01			

In the presence of Cl<sup>-</sup> ions, the control sample corroded rapidly, with a CR of 12.06 mm/yr compared to Al with 25% PVP sample, which had the lowest CR of 6.13 mm/yr. Similar reduction trend in CR values from reinforced samples was observed as the Ct of PVP increased from 10 to 25%, gaining a 41% decrease in CR. This confirms the inherent inhibitive nature of PVP. Fig. 5 shows corrosion response stability throughout each composite sample during a 120 secs.

timeframe. The plot shows that the sample without PVP reinforcement consistently had the highest potential across the 120 secs. period. This specimen had the most noticeable shift towards the positive region, indicating a higher potential. In contrast, samples with different Ct of PVP showed lower values, shifting towards negative potential range. Notably, the specimen with 20% PVP reinforcement had the lowest potential, lower than -0.5 V. The curve's flow towards the negative region indicates that PVP had more impact on the cathodic than on the anodic reaction, implying that it was cathodic inhibitory [40-42].



Figure 5: OCP plots of PVP samples.

The influence of corrosion polarization's cathodic effect is further elucidated in Fig. 6, which showcases a decrease in both cathodic reactions of sample corrosion, as the reinforcement Ct increased from 10 to 25 wt.%.



## **Opto-electrical response of the samples**

Figs. 7 and 8 depict the results of electrical resistivity and conductivity for the produced coating samples.



Figure 7: Electrical resistivity of composite samples.

Interestingly, the trend in electrical resistivity did not follow a uniform pattern, as the results show a paired array, with the control and the most reinforced sample exhibiting the same resistivity value of 5.11  $\Omega$ m, while 10 and 20% reinforced samples had similar values of 5.09 and 5.07  $\Omega$ m, respectively (Fig. 8).



Figure 8: Electrical conductivity of composite samples.

Conversely, a reverse pattern was observed for conductivity plots, with the control and the most reinforced sample having the same conductivity value of 0.1957  $\Omega m^{-1}$ , while 10 and 20% reinforced samples had similar values of 0.1964 and 0.1971  $\Omega m^{-1}$ , respectively. However, 15% PVP sample exhibited highest conductivity of 0.1943  $\Omega m^{-1}$ . Contrary to expectations based on prior research studies, this study revealed a deviation attributed to non-uniform distribution of PVP across Al matrix [37, 43, 44]. Magnetic permeability response of the samples is shown in Fig. 9. Al, unlike Fe, is not ferromagnetic, which corresponds to its low magnetic permeability. Experimental investigation aimed at assessing the influence of PVP on specimens' enhancement revealed a

negligible alteration in metal ferromagnetism, evidenced by voltage variation across diverse samples with increasing inductance, which amounted to a total voltage change of merely 0.17 V in the most reinforced sample.



Figure 9: Magnetic permeability response of the samples.

## Microstructural analysis of samples

SEM micrographs and EDXS analysis of composites with lowest reinforcement Ct (10% PVP) are depicted in Fig. 10.



From SEM images, it is evident that PVP adequately dispersed within the matrix, allowing for visualization of grain boundaries and pores throughout Al matrix. EDXS spectra indicate peaks corresponding to Al, Ca, Si, Mg, C and O, which confirmed that CaO,  $SiO_2$  and  $Al_2O_3$  were major constituents of the reinforcements, as confirmed by PVP characterization. Additionally, oxygen

presence suggests its contribution for silica phase of PV. Similar observations have been reported by previous studies [29, 45].

Fig. 11 presents SEM/EDXS morphology of 25% PVP sample. Analysis of SEM images indicate PVP agglomeration in certain regions of the base matrix, resulting in a reduction in distinct grain boundaries and an increase in porous areas across the matrix.





Figs. 12 and 13 show XRD profiles of 10 and 25% PVP samples, respectively. The plots analysis reveals the highest characteristic diffraction peaks for Al-Si based compound (AlZnS<sub>2</sub>), in both PVP samples, at  $2\theta = 39^{\circ}$  and  $2\theta = 108^{\circ}$ . These peaks correspond to the substantial integration of PVP within the MMC. Furthermore, a range of additional crystalline formations was observed across diverse phases, including Al(Mn), Ca(Zn), Mg(Si) and Mg(Zn), which provides evidence supporting the enhancement of corrosion resistance, strengthened adhesion and refined surface conditioning.



Figure 12: XRD of coated samples (10% PVP).



Figure 13: XRD of developed sample (25% PVP).

#### Conclusion

In conclusion, the study revealed notable findings across various experiments. HRB test demonstrated an average increase in hardness values from 28.1 HRB, for the control sample, to 38.5 HRB for the composite containing 25% PVP. CR values from for control and composite samples with 25% PVP were 12.06 and 6.13 mm/yr, respectively, which is a substantial decrease. Electrical resistivity and conductivity measurements showed intriguing deviations from expected trends: resistivity values shifted from 5.07 to 5.11  $\Omega$ m; and conductivity values changed from 0.1943 to 0.1971  $\Omega$ m<sup>-1</sup>, across different composite samples. Microstructural analysis confirmed uniform dispersion of PVP within the matrix, with SEM/EDXS and XRD analyses highlighting its integration within the MMC, and the presence of additional crystalline formations. Overall, these results underline PVP potential as an effective AMMC reinforcement, offering promising prospects for the development of lightweight and high-strength materials with reduced environmental impact and production costs.

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## **Competing interests**

The authors declare no competing interests.

## Authors' contributions

**A. D. Adeleye**: writing of manuscript and figures. **O. S. I. Fayomi**: research supervision. **J. O. Atiba**: writing of manuscript and tables.

## Abbreviations

Al<sub>2</sub>O<sub>3</sub>: aluminum oxide **AMMC**: aluminum metal matrix composites **ASTM**: American Society for Testing and Materials CMC: ceramic matrix composites **CR**: corrosion rate Ct: concentration Ecorr: corrosion potential **EDXS**: energy dispersive X-ray spectroscopy HCl: hydrogen chloride HRB: hardness Rockwell B J<sub>corr</sub>: corrosion current **MMC**: metal matrix composites **PMC**: polymer matrix composites **PV**: *Phaseolus vulgaris* (bean shell) **PVP**: *Phaseolus vulgaris* particulates **R**<sub>p</sub>: polarization resistance **SEM**: scanning electron microscopy SiC: silicon carbide

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