Laser shock peening (LSP): Electrochemical and hydrodynamic investigation of corrosion protection pre-treatment for a copper surface in 3.5 % NaCl medium

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ABSTRACT

Besides electrochemical corrosion protection, laser shock peening provides extended fatigue life and mechanical properties for metals, as reported in the literature. However, most of the studies were performed under static conditions. The effectiveness of this method under flow conditions and prolonged immersion duration is addressed in this study. Copper discs were peened with Nd: YAG laser in the presence of ~2 mm thickness of water and polyvinyl chloride to absorb the intensity and provide residual stress without surface damage. Electrochemical and hydrodynamic analyses indicate that the peened surface showed higher corrosion resistance in 3.5 % NaCl compared to the unpeened.

1. Introduction

The material surface plays a major role in the electrochemical response and engineering applications. Therefore, surface modifications are conducted for a wide range of reasons including the improvement of physical, chemical, and mechanical properties. This follows the fact that the quality and performance of a metallic product can be directly correlated to its surface integrity achieved from different surface treatment processes carried out during production [1,2]. Several methods of surface modification techniques are in practice such as mechanical attrition, hot and cold rolling, high-pressure torsion, ball milling, mechanical shot peening, and laser shock peening to improve the surface microstructure, mechanical strength, corrosion resistance and tribological properties of the metallic material [3–6]. The surface treatment process focused on the present research is the laser shock peening (LSP) used for the improvement of corrosion protection and fatigue life of metals [7–9]. LSP is preferred to the rest of the processes because of its consistency in results and high-quality surface finish [10–12]. Application of laser light on material surfaces is an age-long process; however, LSP is a recently developed advanced surface modification technology

that has been very effective in improving wear resistance, hardness, and corrosion resistance of metallic materials. It is a cold working process where pulses hit the metal surface through a high power intensity laser; generating shock waves that propagate through the metal to a certain depth (~0 to 2 mm) depending on the power density applied [3]. The basic mechanism involved in this process is to create a compressive residual stress on the material which helps to delay premature failures. The compressive residual stress is seen to be responsible for the modification of the microstructure and improvement of the surface hardness and corrosion resistance. [13–15].

In addition to the improvement of electrochemical corrosion resistance and fatigue life, laser shock peening was found to be very useful in the enhancement of the metal resistance to stress corrosion cracking (SCC) and grain boundary refinement. Consequent to the grain boundary refinement, reduction in defects such as porosity and micro-cracks were also observed, these in turn reduce the oxygen diffusivity to the metal substrate when LSP treatment is applied on the surface [9,16,17]. Mahdavi et al. [8] investigated the effects of LSP on the electrochemical corrosion and microhardness properties of ANSI 316 L stainless steel alloy. Comparing the untreated and treated samples, it was found that

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surface hardness improved by 35 % while the corrosion protection performance increased by 100 % for the treated samples in 3.5 % NaCl solution. In another development, the corrosion resistance of a biodegradable metallic implant (magnesium–calcium alloys) treated by LSP showed an increase in corrosion resistance by more than a 100-fold in simulated body fluid electrolyte as reported by Guo et al. [18]. Similarly, Zhang et al. [19] reported a wide positive shift in the corrosion potential of nickel-titanium alloy from -440 mV to -80 mV (a difference of -360 mV) and reduction in corrosion current density from 1.41 to $0.67 \mu\text{A cm}^{-2}$ for the untreated and the LSP treated surfaces respectively, tested in a simulated body fluid. This result by Zhang group was accompanied by an increase in hardness value from 226 to 261 HV due to the LSP treatment.

Further investigation was carried out by Vijayalakshmi et al. [7] on the effects of laser shock peening (LSP) on titanium alloy (Grade 5—Ti6Al4V) to establish the best set of LSP parameters for the enhancement of surface characteristics of the alloy. The result shows an excellent mechanical property for the peened surface compared to the unpeened. The experiment optimized the parameters such as power density (3 GW/cm^2), wavelength (532 nm), and pulse overlap (60%) for titanium alloy. The rate of corrosion was found to be least at these sets of parameters due to low roughness and surface oxide layer which holds the fluctuation in current density with respect to potential. Liu et al. [16] demonstrated a repair method for fatigue damage using LSP in which they reported both improvement in fatigue life and healing effect on the copper film. Yujie et al. [20] showed that the distribution of grain size becomes more uniform and smaller near the top copper surface after LSP treatment. In addition, grain refinement was observed for copper along the normal (depth) direction [21,22]. Kim et al. [23] recently investigated the effect of laser shock peening on the fretting resistance of copper used for electrical components in automobiles. In their experiment surface hardness was improved from 55 to 120 HV for the unpeened and peened copper respectively. The wear loss of the base metal was decreased from $1.61 \times 10^{-3} \text{ mm}^3$ to $0.94 \times 10^{-3} \text{ mm}^3$. Thus, the laser shock-peened copper maintained a low electrical contact resistance throughout the fretting test. Electrical contacts in automobiles usually fail due to wear arising from the small amplitude of reciprocal motion and blockage by corrosion products in the presence of air and moisture. However, the electrochemical corrosion behaviour of the treated copper surface was not investigated in this experiment. Moreover, despite these improvements in the use of LSP on metal protection, all the previous LSP corrosion analyses were performed under static conditions.

Therefore, the present research focused on the electrochemical and hydrodynamic flow characterization of the corrosion resistance and protection persistence of laser peened copper in 3.5% NaCl medium. The present investigation will explore the effect of exposure time and hydrodynamic flow on the peened and unpeened surfaces. Following the fact that among a large number of adverse environments for corrosion such as salinity, dissolved oxygen, pH, temperature variation, etc. flow environment has one of the highest destructive impacts on material loss and degradation [24–28]. This is important as LSP is fast becoming adopted in the area of corrosion protection, though it does not stand as a replacement to other corrosion protection methods in the literature. This is because the application of each method is defined by the peculiarity of the environment and nature of the management process involved. Laser shock peening is mainly targeted at aerospace, automobile, and marine environment components. In such environments, the use of inhibitors, conducting polymers, and self-assembly monolayer (SAM) of corrosion inhibiting compounds can not be applied even when they are economically more affordable. Laser shock peening has an advantage, as it can be applied to almost all metallic material for different aggressive environments. Therefore, validating the performance of LSP on copper corrosion protection in the flow environment becomes imperative not only to fill the knowledge gap but also for quality corrosion protection performance of copper and its alloys. This will provide a qualitative and

quantitative assessment of this method for its proper application on metal corrosion protection especially where other methods such as corrosion inhibitors [29–31], may not be applicable. The present study will be compared to the corrosion protection efficiencies of the other methods in the literature. The electrochemical parameters that define metal stability in the corrosive media will be compared to ascertain the level of progress made. It is worthy to mention that the proposed LSP process is cost-effective especially at industrial scale since the same laser source can be used for a large number of products. It is easily applicable by simply shining the laser light on the metal surface and environmentally friendly since the biological hazard can only occur at exposure such as a direct incident of laser on eye or skin and this can be managed through the laid down protocols [32].

2. Experimental procedure

2.1. Metallic sample preparation (laser shock peening)

Laser shock peening (LSP) experiment was performed by directing a low energy Nd: YAG laser (Litron lasers, UK) on a 1 cm^2 polished Cu disc after covering with polyvinyl chloride (PVC) tape of $\sim 1 \text{ mm}$ thickness. The surface was submerged in a thin layer of flowing water ($\sim 2 \text{ mm}$) to further control the laser intensity [33,34]. Some authors preferred to use only flowing water without a covering or coating sub-layer thus they termed the peening process as laser shock peening without coating (LSPWC) [3,6]. The LSPWC allows a higher intensity of the laser light to hit the metal surface unlike the coated or covered one. This has a consequence of increasing the surface roughness and thermal effects, so we decided to use the covering method for the present work to minimize these problems. The Cu surface was thus covered with polyvinyl chloride tape in the present method as earlier stated. Therefore, the peening process was termed as laser shock peening with polyvinyl chloride (LSPVC). This term will be used throughout the discussion of this work.

The LSPVC optimization was carried out using the design of experiments (DoE) by tuning the laser energy and the overlap percentages as tabulated in Table 1. Also, the study investigated the sacrificial opaque layers of PVC tape at high, low, and medium power density and percentage overlap as used in the experiment. The energy consumption for the laser is varied with a pulse duration of 10 ns delivered with a spot diameter of 0.8 mm and with a wavelength of 1064 nm . The maximum available repetition rate of 10 Hz was used for the whole LSPVC process. The exhibited radiance density (laser beam brightness) was $345.04 \text{ mW. mm}^2.\text{Sr}^{-1}.\mu\text{m}$. The laser comprised of an M^2 value of 2 and a beam divergence of 0.5 mrad . The beam shape was of a top-hat profile [35, 36]. The electrical dryer was used to protect the lens from the water spilling during the LSPVC process. A 2D (XY) translation motorized stage (SVP Lasers, India) was employed to perform the LSPVC experiments in transverse and longitudinal directions.

2.2. Microstructural characterization

The phase structure and crystal orientation were investigated by X-ray diffraction (XRD) using a Brucker diffractometer with Cu-K α radiation ($\lambda = 1.54056 \text{ \AA}$) as a source. The scan was carried out at 2θ range from $30^\circ - 85^\circ$ at the scan rate of 5° per minute. The surface roughness, topography, and morphology were examined by atomic force microscopy (AFM, Nanosurfesycan 2, Switzerland) and scanning electron microscope (Zeiss SEM instruments). The water droplet contact angle (θ)

Table 1
Design of the experiments (DoE) for LSPVC optimization.

Sample No	Power density (Gw/cm^2)	Description	Overlapping (%)	Mask
1	4	Low	40	PVC
2	6	Medium	70	PVC
3	8	High	50	PVC

was measured on the peened and unpeened specimen using a goniometry instrument (Model Phoenix 300, SEO, Korea) at room temperature (25 °C). The droplet volume of 10 μ L distilled water was used to examine the surface wettability characteristics. Further, the mechanical properties of the compressed Cu surface were investigated at the nanoscale with the Nanoindentation instrument (Nano Test Vantage T61) from Micro Materials UK.

2.3. Electrochemical measurements

An Autolab Potentiostat (PGSTAT302) with three-electrode arrangements was used for electrochemical characterization. Platinum and saturated calomel electrodes (SCE) were used as counter and reference electrodes respectively, while the LSPVC peened Cu samples acted as the working electrodes. All the tests were carried out at room temperature using a 3.5 % NaCl electrolyte. Prior to all the electrochemical studies, the working electrode was introduced into the test solution and was allowed to attain a steady-state potential. Then a potential range of -0.3 to + 0.6 V vs. SCE was applied with scan rate at 1 mV/s. E_{corr} and I_{corr} were obtained from the potential (E) vs. $\log j$ curves by the Tafel extrapolation method. EIS studies were carried out at static conditions for short and extended immersion time, thereafter; the hydrodynamic effect was investigated through electrochemical experiments carried out under flow conditions using a rotating cage (RC) instrument [24]. The initial frequency of 100 kHz, final frequency 10 mHz, and AC sine wave amplitude perturbation of 10 mV was applied at OCP. The impedance spectra were recorded with the unpeened and LSPVC peened Cu at different laser overlap percentages. All electrochemical experiments were replicated three times and all the potential values were measured with respect to SCE.

3. Results and discussion

3.1. X-ray diffraction (XRD) analysis

Fig. 1 shows the XRD patterns of the LSPVC treated Cu surfaces at 40, 50, and 70 % overlap in comparison with the unpeened surface. The results showed that the unpeened Cu exhibited three peaks observed at $2\theta \sim 44^\circ$, $\sim 51^\circ$, and $\sim 75^\circ$ indicating the characteristic peaks of Cu metal before LSPVC treatment. After the treatment, similar peaks were obtained in all the treated samples with minor changes observed on the (111) and (220) planes. This fact showed that there was neither formation of surface oxide or surface damage nor alteration of Cu crystal

structure in the course of LSPVC treatment. A clear observation of the (111) plane showed little diminished intensity at 40 and 50 % overlap while 70 % overlap showed effect at the (220) plane which could be signs of surface modification. Although XRD which normally gives information from the bulk sample may not provide enough information on the surface changes. Thus further analysis was done with the help of scanning electron microscopy (SEM) as presented in the following discussion.

3.2. Scanning electron microscopy (SEM) analysis

The images of the Cu samples were taken after laser treatment using the scanning electron microscope. Fig. 2 shows the micrographs of the unpeened Cu surface and the peened at 40, 50, and 70 % laser overlap. These show clearly that there were surface impacts which were minimal at 40 %, high at 50 %, and moderate at 70 %, see Fig. 2b, c, and d respectively. As observed, the 40 % overlap was minimally impacted by the laser light due to a reduced power density (4 Gw/cm²) with pulse overlap. 50 % overlap shows high impact as depicted by the more pronounced surface dimples and compact shrinking attributed to the doubling of the laser power density to 8 Gw/cm² at this overlap, whereas 70 % overlap peened at 6 Gw/cm² shows less formation of dimples compare to 50 % overlap but depicts more local impact leading to surface shrinking and initiation of valleys. The optimization of these parameters was shown in Table 1. The observed effects clearly show that power density has more impact on the surface changes than pulse overlap during the LSP process [8]. Remember that the Power density has to do with the laser energy while the pulse overlap is the number of packets of the laser light hitting the surface at a particular point in time. The combination of the effects of these two parameters gives the overall peening impact. In Fig. 2, all the peened surfaces showed impacts as a result of the compression of the top surface leading to the intertwining structures and inducement of compressive residual stress which in turn led to shrinking and dimples on the peened samples relative to the applied parameters. The viewed effects of the SEM analysis were clearly represented in Fig. 3, which shows the schematic representation of the residual stress and impact depth of the unpeened and LSPVC peened surface at 40, 50, and 70 % overlap. This is to further show that the 40 % overlap has low impact depth with shallow dimples, 50 % showed multiple effects and more dimples while 70 % shows fewer impacts but valleys of local and nonuniform laser effects [13,37].

3.3. Contact angle analysis

The contact angle analysis of the unpeened and LSPVC peened Cu surface at 40, 50, and 70 % overlap measured at room temperature under atmospheric pressure is presented in Fig. 4. Contact angle measurement describes the surface energy and the wettability characteristics of a material. A given surface at a particular temperature and pressure has a unique equilibrium contact angle. If the water contact angle is smaller than 90°, the solid surface is considered hydrophilic and if the water contact angle is larger than 90°, the solid surface is considered hydrophobic. Fig. 4 shows that the contact angle value obtained for unpeened is 36.1°, whereas the peened surface shows 63.8° for the 40 %, 90.3° for 50 %, and 86.7° for the 70 % overlap. This indicates that the contact angles of the Cu surface increased with the increase in the laser overlap up to 50 % and dropped at 70 %. Despite the increase, it was only the 50 % overlap that effectively transformed from a hydrophilic surface to a hydrophobic one. At 70 % overlap, there was a fall in the contact angle value due to reduced hydrophobicity may be a result of local valleys which increased the roughness. Since roughness enhances the wettability and surface chemistry, it, therefore, means that 90.3 % contact angle of the peened surface shows that the surface roughness was not excessively increased by the laser treatment. The relationship between surface roughness, measured contact angle and the Young contact angle was explained using Wenzel correlation for a

Fig. 1. XRD patterns of the unpeened and LSPVC peened Cu surface at 40, 50 and 70 % overlap.

Fig. 2. SEM micrographs of the unpeened and LSPVC peened Cu surfaces (a) Unpeened bare Cu and the peened at (b) 40 %, (b) 50 %, and (c) 70 % overlap.

Fig. 3. Schematic representation of the Unpeened Cu surface compared to the LSPVC peened showing dimples, compact top layer, and the residual stress impact of the peened at 40, 50, and 70 % overlap.

homogeneous surface in the following Eqn;

$$\cos(\theta_m) = r \cos(\theta_Y) \quad (1)$$

Where θ_m is the measured contact angle, θ_Y is the Young contact angle and r is the roughness ratio. The roughness ratio is defined as the ratio between the actual and projected solid surface area [38–40].

3.4. Analysis of mechanical properties

Fig. 5 shows the nanoindentation data of the unpeened and LSPVC peened Cu surface at 40, 50, and 70 % overlap. Fig. 5a is the hysteresis of

the loading and unloading curves. The area bounded by loading and unloading curves defines the degree of resistance to plastic deformation. An increase in the area correlates to an increase in plastic deformation and vice versa [41]. In the present research, a 50 % overlap showed the highest resistance to plastic deformation as indicated by the small area under the curve. Fig. 5b shows the varying hardness values with laser overlap. The hardness of the unpeened bare Cu was 1.188 GPa, surprisingly higher than that of 40 % peened which was found to be 1.025 GPa. This could be attributed to the generation of nanoparticles and contaminants at the metal surface at the beginning of the LSP process when the laser intensity is low [8]. This shows that at low laser overlap

Fig. 4. Contact angle data of the unpeened and LSPVC peened Cu surface at 40, 50, and 70 % overlap measured at room temperature under atmospheric pressure.

and power density, the surface becomes poorly modified yielding surface products or pollutants of less hardness. At 50 % overlap hardness increased to 2.597 GPa while at 70 % it was 1.522 GPa, indicating the presence of compressive residual stress on both samples. This shows that the optimum hardness was achieved at 50 % where the power density had a good match with the overlap, below or above which there could be

fluctuations in hardness due to mismatch of laser energy with overlap.

Moreover, residual stress was further studied in detail using the load-depth curves. It is well known that the residual stresses significantly affect load-depth curves. Residual stress can be evaluated using indentation tests if the value of the hardness and the loading curve is higher than initial values after LSPVC processing; it indicates that compressive residual stress is present on the surface. If the hardness is lesser, then tensile residual stresses might be present [33,42,43]. The impact of residual stress on the load-depth curves was examined at different positions where the indentation depths were held constant (Fig. 5c). For example, at a constant depth of 700 nm, it was obvious that the curves rose according to their compressive residual stress with the 50 % overlap showing the highest residual stress. This is in line with the report in the literature that the compressive stress can constrain the indentation plasticity, thus leading to a higher indentation load than that for the stress-free state where less load was required for the same indentation depth [44]. On the other hand, holding the load constant shows similar trends for the effects of residual stress on the indentation depth Fig. 5d. Apart from the 40 % overlap, it is clear from Fig. 5d that the indentation depths shifted to the left in accordance with their residual stress. 50 % shows the least indentation depth penetration. The relatively lower depth of penetration in the laser peened sample could be attributed to the enhanced hardness and compressive residual stress inducement. Therefore, it can be concluded from this experiment that residual stress does not increase indefinitely with the increase in the overlap but there exists an optimal value where the overlap together with the applied power density makes a good match as stated earlier, above or below this optimal value the laser light might not impact directly on the residual stress but can be converted to surface melting. Therefore, in general,

Fig. 5. Nanoindentation data of the unpeened and LSPVC peened Cu surface at 40, 50 and 70 % pulse overlap (a) hysteresis of the loading and unloading curves (b) variation of hardness values with laser overlap, residual stress impacts on (c) loading curves at constant depths and (d) the indentation depth at constant loads.

LSPVC induces compressive residual stresses and there is an optimal set of operating parameters for this condition to improve hardness and surface quality.

3.5. Potentiodynamic polarization

Prior to potentiodynamic polarization and other electrochemical measurements, the stability and free dissolution pattern of the unpeened and the LSPVC peened Cu surfaces was first monitored by potential measurement in the absence of current known as the open circuit potential (OCP). Fig. 6 shows the OCP, potentiodynamic polarization, and the corrosion rate curves for the unpeened and peened Cu surface at 40, 50, and 70 % overlap; immersed in 3.5 % NaCl solution at room temperature. Fig. 6a shows the variations in the OCP of the unpeened and peened samples with pulse laser overlap. The measurements were recorded for 1000s since all the peened samples showed stability with less than 0.001 V variation beyond 500 s. LSPVC 50 % showed potential decay from -0.145 V to a final value of -0.162 V vs. SCE. The other peened samples, show minimal changes with time, probably due to reduced roughness. Power density tends to increase surface hardness with a consequent increase in roughness [7]. But the advantage of the improved hardness outweighs that of the roughness in the long run for the metal protection. LSPVC 40 and 70 % showed minor potential decay whereas the unpeened Cu surface showed potential decay from -0.29 to a final value of -0.32 V vs. SCE, typical E_{corr} of bare Cu surface in 3.5 % NaCl [29,45]. In all the plots, the free potential (E_{corr}) of Cu tends positive with peening overlap with 50 % overlap showing the highest

positive value.

Fig. 6b shows the potentiodynamic polarization curves for the unpeened Cu in comparison with the peened surfaces at 40, 50, and 70 % overlap. The corrosion potentials obviously shift to more positive values from -0.323 to -0.162 mV at 50 % overlap (equivalent to 49.9 % increase). The corrosion current density reduced from 6.31 to 0.35 $\mu\text{A cm}^{-2}$, indicating that the best performance of the modifications is at this overlap applied at a power density of 8 Gw/cm². However, looking at the curve due to LSPVC 50 %, the E_{corr} is in the active dissolution region of the anodic part, which shows extensive Tafel behaviour indicating a high rate of corrosion at the onset before exhibiting a reduction in corrosion current. This could be the evidence of the potential decay witnessed in the OCP analysis, probably due to the roughened part of the surface by high power density [7], compared to the less impacted surface at 40 and 70 % overlap with the lower power density of 4 and 6 Gw/cm² respectively. However, the power density effect on the LSPVC 50 % could not overturn the overall performance in the reduction of the corrosion current density as the surface is at the same time hardened by the compressive residual stress arising from the higher laser power. Fig. 6c shows clearly the extent of performance of the entire modification process when the best performing sample (LSPVC 50 %) was compared to the unpeened surface. This can be ascribed to the compact high resistant surface due to the shrinking effect and grain refinement of the high powered laser light from LSP processing, [9]. The performance of the LSPVC modification at different overlaps was further analyzed quantitatively and presented in Table 2. The trend of the parameters in Table 2 shows that the inhibition efficiency and the free corrosion

Fig. 6. (a) Potential vs. time (OCP), (b) potentiodynamic polarization curves for unpeened and peened Cu surfaces at 40, 50, and 70 % overlap immersed in 3.5 % NaCl solution at room temperature, (c) the direct comparison of the best performing (LSPVC 50 %) with the unpeened sample and (d) the corrosion rate at 0 % (unpeened), 40, 50 and 70 % overlap.

Table 2

Parameters related to the Potentiodynamic Polarization study of Cu immersed in 3.5 % NaCl for the unpeened and peened surface at 40, 50 and, 70 % overlap at room temperature.

LSPVC Overlap (%)	E_{corr} (V _{SCE})	I_{corr} (μA cm ⁻²)	E_{corr} (mV/decade)	β_c (mV/decade)	Corrosion rate (mmpy)	η (%)
Unpeened	-0.323	6.31	129.5	140.3	0.123	–
40	-0.304	2.40	94.5	136.2	0.114	61.97
50	-0.162	0.35	54.6	79.7	0.008	94.45
70	-0.313	1.54	106.7	120.7	0.105	75.59

potential (E_{corr}) increased in the positive direction while the corrosion current density and the corrosion rate decreased with an increase in laser percentage overlap. The trend further showed that the corrosion current density decreased with the increase in laser overlap with a minor deviation at LSPVC 70 %. The 50 % peened has the lowest corrosion current density of 0.35 μA cm⁻². The corrosion rate was reduced from 0.123 to 0.008 mmpy with corrosion protection efficiency of 94.45 % as calculated from the I_{corr} value for LSPVC 50 %. The trend in all the parameters in Table 2 indicates the effective protection of Cu in 3.5 % NaCl by the LSPVC process. The potentiodynamic data in Table 2 were obtained from Tafel extrapolation of the just discussed E vs. $\log j$ curves of Fig. 6b. We can see clearly that the copper electrode immersed in a 3.5 % NaCl solution exhibits a typical Tafel behaviour in both the anodic and cathodic parts [46,47]. The cathodic parts were not clearly limited by oxygen diffusion but still display a Tafel behaviour with the corrosion current increasing proportionally with the potential (see Fig. 6b), typical behaviour of copper in NaCl system [48,49]. It is known from the literature that the extrapolation of both anodic and cathodic sides gives more accurate values of the potentiodynamic parameters thus was applied in this study [9,46]. The extrapolation of the pair of Tafel regions in the curves and the corrosion rates were accurately determined using the Ec-lab corrosion software model 10.40. The condition is that the Tafel region must be linear for at least one decade of current after a potential scan of about ±300 mV, and is taken within 50–100 mV above and below the E_{corr} [50]. The current value at the E_{corr} intersection is then the corrosion current (I_{corr}), directly called the corrosion current density when the working electrode (WE) is a standard 1 cm² surface area. Other values of the WE area must be used to divide the measured equilibrium corrosion current to obtain the corrosion current density (I_{corr}) according to the following equation [50].

$$I_{corr} = I_{eq}/A_{WE} \quad (2)$$

Where I_{eq} and A_{WE} are the measured equilibrium corrosion current and the WE area respectively.

From the corrosion current density, the protection efficiency (η %) of the peened surfaces were determined against the unpeened, according to the inhibitory efficiency formula, Eq. 3.

$$\eta \% = \frac{I_{u_{corr}} - I_{p_{corr}}}{I_{u_{corr}}} \times 100 \quad (3)$$

Where $I_{u_{corr}}$ and $I_{p_{corr}}$ are the corrosion current densities of the unpeened and the peened surfaces respectively.

The electrochemical results are further depicted in the plot of the corrosion rate versus percentage overlap of the different modifications compared to the zero modified surface (unpeened Cu) in Fig. 6d. This shows that there was a tendency to reduce the corrosion rate in all the laser peened samples in comparison to the unpeened Cu, whereas the highest performance in corrosion rate reduction was found at LSPVC 50 %. The results not only confirm that the LSP modification is optimum at LSPVC 50 % but also showed that the LSP process is an effective method for Cu corrosion mitigation [7,9,19,51].

3.6. EIS analysis

The electrochemical impedance plots for peened and unpeened Cu surface immersed in 3.5 % NaCl solution at room temperature under static conditions are presented in Fig. 7. Fig. 7a shows the Nyquist plots for the unpeened and peened Cu surfaces at different modifications of 40, 50, and 70 % pulse overlap. The diminished impedance capacitive loop for the unpeened is shown in the inset of Fig. 7a. This shows that there was a large improvement in the corrosion resistance of all the laser modified surfaces. The peened samples show an increased diameter of the impedance capacitive loops due to enhanced corrosion performance. The Nyquist plots exhibit a single capacitive loop implying that the corrosion of the peened samples is mainly charge transfer controlled (i.e. charge transfer is the slowest process) from the surface to the electrolyte through the shrunk compact Cu surface [52,53]. The resistance increased with an increase in the laser percentage overlap and the applied power density. LSPVC 50 % showed the highest corrosion resistance, due to the high impact of laser power density which optimized at 8 Gw/cm² with a 50 % overlap. This is in line with the improved hardness and compressive residual stress as observed from the indentation results (Fig. 5) [7]. At LSPVC 70 % there was a decrease in the corrosion resistance which could be attributed to a mismatch of power density and the percentage overlap as the surface behaviour is dependent on the interplay of the overlap and power density inputs as shown in Table 1. This could also be attributed to an increase in local roughness as witness in this sample by the aid of SEM [8]. Fig. 7b and c are the Bode modulus and phase angle plots which are also in line with the Nyquist plot data, the LSPVC 50 % shows the highest impedance modulus and phase angle value of all the modifications and the unpeened Cu surface.

Fig. 8 showed the equivalent circuits fit for the quantitative analysis of the systems which could be grouped into two, Fig. 8a was used to fit all the plots except those with Warburg impedance (W) (see plots of unpeened bare Cu, inset of Fig. 7a and 0 h – 228 h of Fig. 9b) which were fitted into the circuit of Fig. 8b. These circuits depict the electrical components of the physical parameters of the working electrode and the solution system. The solution medium shows the solution resistance R_s to the movement of ions and species in solution. The surface of the electrode exhibited high charge transfer resistance (R_{ct}) due to LSPVC peening. R_f represents the film resistance developed on the Cu surface after peening process and during immersion as a barrier accumulated between the metal and the aggressive medium interface. The major effect of the peening process is not the development of film but to induce residual stress which makes the surface compact and thus the R_{ct} was improved tremendously [7,8]. The overall resistance of the system is expressed as the polarization resistance (R_p) which is the combination of the R_f and R_{ct} . Further the data showed two non-ideal capacitances represented as constant phase element (CPE1) arising from the film/solution interface and (CPE2) for the double layer capacitance due to surface inhomogeneity between the metal and the surrounding film. The Y_{o1} and Y_{o2} are the magnitudes of the constant phase elements corresponding to the CPE1 and the CPE2 for the film and the metal surface respectively. The dispersion coefficients n_1 and n_2 showed value Table 3 less than unity as an indication for the film imperfection due to roughness and porosity, see Table 3, [54,55].

The quantitative values of the charge transfer resistance (R_{ct}) and other electrochemical impedance parameters related to the unpeened and the different peened samples are presented in Table 3 which were generated from electrochemical circuit fit carried out using Zsimpwin 3.21. It is clear from Table 3 that LSPVC 50 % shows the highest R_{ct} value, 17,020 Ω. cm² compared to 957 Ω. cm² of the unpeened, however, there was increasing trend in the R_{ct} with laser overlap. The value of film resistance (R_f) in Table 3 is relatively small probably because the major resistance portion is arising from surface compressive residual stress which contributed more to the improvement of R_{ct} of Cu through

Fig. 7. The electrochemical impedance (EIS) plots for peened and unpeened Cu surface immersed in 3.5 % NaCl solution at room temperature under static conditions: (a) the Nyquist, (b) Bode modulus, and (c) Bode phase angle at 0 % (unpeened), 40 %, 50 %, and 70 % overlap.

the shrunk surface, thus, minimal film formation was observed [8]. Overall film resistance of the LSPVC peened samples is greatly enhanced compared to the unpeened specimen [7]. The Y_{01} and Y_{02} showed decrease in trend with increase in laser overlap see Table 3, confirming the improvement of the surface resistance to water penetration due to the surface compactness. In addition, the dispersion coefficient (n) increased with increase in overlap percentage as well, this is in line with improved corrosion protection behaviour of the Cu surface in the aggressive medium by the LSPVC process. The chi square (χ^2) values in Table 3 further validates the minimal error margin in the electrochemical circuit fit since the values are only little above 1×10^{-3} . All these amount to the protection efficiency (η %) of 94.38 %, from EIS, calculated using Eqn. 3 and is in good agreement with the potentiodynamic result in the previous discussion.

$$\eta \% = \frac{R_{pct} - R_{uct}}{R_{uct}} \times 100 \quad (4)$$

Where, R_{pct} and R_{uct} are the charge transfer resistances of the peened and unpeened surfaces respectively.

Therefore, to further investigate the efficacy of this method on Cu, electrochemical impedance spectroscopy was carried out to determine the effect of extended immersion time and flow in the corrosive environment of the same electrolyte (3.5 % NaCl), the results are presented in Fig. 9. This experiment was performed with best performing sample (LSPVC 50 %) in comparison with unpeened Cu. The Nyquist plots of the LSPVC 50 % peened and the unpeened Cu surface are shown in Fig. 9a and b respectively at different immersion durations from 0 h to 304 h (12 days plus 6 h). The overall impedance results showed a reduction in

R_{ct} for both systems (peened and unpeened) during long immersion duration compared to the previous results of Fig. 7. This is definitely due to prolonged corrosion attack in the presence of chloride ion (Cl^-). The increasing tendency of the R_{ct} with immersion time in Fig. 9a and 9b could be attributed to concentration polarization resulting from accumulated corrosion products since by extending the immersion duration the resistance became lower than the original value but regained some resistance as time progressed further. It is interesting to note that despite the exposure for long time duration the LSPVC 50 % peened still showed evidence of better performance compared to the unpeened which showed drastic fall in the R_{ct} values with time as clearly shown by Fig. 9c. The results conclude that the LSPVC surface modification is capable of providing long-time protection for Cu metal in the corrosive chloride medium. Further, the samples were exposed to a flow environment of the same corrosive electrolyte (3.5 % NaCl) to investigate the hydrodynamic effect on the performance of the peened samples in comparison to the unpeened. This is very crucial since the corrosion attack is often more aggravated by hydrodynamic flow than the static systems. Fig. 9d presents the representative results of the hydrodynamic experiments at different flow rates of 120, 240, and 360 rpm. The results revealed the stable nature of the LSPVC 50 % compared to the unstable bare unpeened Cu surface. This was indicated by the gradual reduction in the R_{ct} by the peened at all the rotation speeds (120, 240 and 360 rpm) as a result of flow effect whereas in the case of the unpeened there was a drastic rise in the R_{ct} at 240 rpm, probably due to formation of corrosion products and its sudden fall at 360 rpm as the corrosion products probably get transported away from the surface as a result of flow. In chloride medium Cu system, CuCl corrosion product is insoluble and can

Fig. 8. Equivalent circuits for the electrochemical parameters fitted for (a) the curves without Warburg impedance and (b) curves showing Warburg impedance (W).

be easily adsorbed on the surface. However, the adhesion of CuCl (formed as a result of the reaction in Eq. 5) to the copper surface is not sufficient to protect the copper from further corrosion. In the second part of the reaction in Eq. 6, the CuCl is transformed into a soluble copper (I) chloride complex [56], which can easily diffuse into the bulk of the solution especially by the aid of the flow medium. All the electrochemical experiments, therefore, prove the better performance of the peened surface compared to the unpeened.

3.7. SEM analysis before and after extended immersion duration

The surface morphology and the extent of corrosion damage before and after the extended immersion duration for the peened (LSPVC 50 %) and unpeened samples were viewed by the aid of scanning electron microscope (SEM) and the results are presented in Fig. 10. Fig. 10a and b are the unpeened bare Cu before and after long-duration immersion respectively. The bare Cu after corrosion showed a high rate of degradation and surface damage with deposits of corrosion products. This could be due to the direct interaction of the unpeened Cu surface with the aggressive chloride medium for an extended immersion period which made the corrosion impacts more evident as observed in Fig. 10b. Whereas, in the case of the LSPVC 50 % peened Cu, Fig. 10c and d, represent the surface before and after long-duration exposure to chloride

medium respectively. The peened surface before immersion (Fig. 10c) showed a different morphology compared to the bare unpeened Cu (Fig. 10a), due to the peening effect. Fig. 10c thus showed surface shrinking, dimples, and intertwining of the physical structures therein, due to laser shock effect. This resulted in the stabilization of the peened Cu surface (Fig. 10d) as it showed virtually no corrosion damage unlike the unpeened. Fig. 10e and f are the digital images of the unpeened and the peened respectively, after long-duration exposure to the chloride medium showing a physical view of the same surfaces previously imaged with SEM in Fig. 10b and d. This clearly showed the real-life damage due to aggressive chloride medium attack. The damage on the unpeened surface is so severe with an indication of a localized micro gully due to the concentration of metal loss in the absence of peening treatment as captured by both SEM and camera imaging. In the case of the LSPVC 50 % peened surface after immersion for 304 h (12 days 6 h), the surface was intact with minimal localized pit damage. These analyses are in agreement with the SEM and conclude that the LSPVC treatment by the laser light has modified the surface into an unpenetrable one by Cl^- through the shock shrinking process which leaves it as compressed permanent stress.

3.8. SEM analysis before and after flow corrosion

Fig. 11 shows the SEM micrographs of the unpeened (Fig. 11a–d) and the peened (Fig. 11e–h) before and after flow corrosion at the rotation speeds of 120, 240 and 360 rpm. The SEM results are in line with the

Fig. 9. The EIS data for the selected LSPVC 50 % and the unpeened Cu surface immersed in 3.5 % NaCl solution at room temperature, (a, b) Nyquist plots for 50 % peened and unpeened Cu respectively, (c) change in the charge transfer resistance (R_{ct}) with immersion time, under static conditions at extended immersion duration, and (d) comparison of R_{ct} with rotation speed at 120, 240 and 360 rpm for the experiments under hydrodynamic conditions.

Table 3

EIS Parameters of Cu in 3.5 % NaCl obtained for the unpeened and peened surface at 40, 50 and, 70 % overlap at room temperature.

LSPVC Overlap (%)	R_s ($\Omega \cdot \text{cm}^2$)	R_f ($\Omega \cdot \text{cm}^2$)	CPE ₁		R_{ct} ($\Omega \cdot \text{cm}^2$)	CPE ₂		$W \times 10^{-4}$ ($\Omega \cdot \text{cm}^2$)	$\chi^2 \times 10^{-3}$	η (%)
			$Y_{01} \times 10^{-4}$ ($\Omega^{-1} \text{cm}^{-2} \text{s}^n$)	n_1		$Y_{02} \times 10^{-4}$ ($\Omega^{-1} \text{cm}^{-2} \text{s}^n$)	n_2			
Unpeened	20.47	17.56	5.3	0.74	957	4.8	0.69	2.27	41.5	–
40	17.13	125.70	3.4	0.79	6675	2.4	0.80	–	20.1	85.66
50	29.32	204.78	2.6	0.88	17,020	1.4	0.85	–	20.0	94.38
70	27.15	151.40	2.2	0.84	9387	3.1	0.80	–	24.0	89.81

previous electrochemical analysis. Fig. 11a shows the image of the unpeened bare polished Cu before flow corrosion, whereas Fig. 11b–d show the images of the bare Cu at 120, 240, and 360 rpm after corrosion showing a progressive trend in the surface damage from micro holes at 120 rpm to crack initiation at 240 rpm and deep cracks at 360 rpm. In the case of the peened Cu surface, observable pitting corrosion occurred only at the 360 rpm (Fig. 11h), prior to that, the surface remained intact with minor tribological effects at 120 and 240 rpm (Fig. 11f and g).

3.9. AFM analysis for peened and unpeened Cu after corrosion

Fig. 12 shows the AFM topographic analysis of the peened and the unpeened after flow corrosion with their respective roughness profiles. The peened surface (Fig. 12a) showed a more even topology with reduced roughness (R_t) and arithmetic deviation of the roughness profile (R_a) in the range of 0.116 and 0.038 μm respectively, (see Fig. 12b). Whereas in the case of the unpeened (Fig. 12c), the surface was uneven

and heavily damaged after corrosion with higher R_t and R_a in the range of 0.969 and 0.121 μm respectively, (see Fig. 12d). Overall, the research showed that the laser-peened Cu surface showed higher resistance to aggressive corrosive 3.5 % NaCl medium than the unpeened Cu surface with the LSPVC 50 % proven as the best preforming peening parameter both under static and flow conditions.

3.10. Comparison of the present method with existing literature

At the end of the data acquisition and analysis, to understand the overall progress made, the results obtained in the present research work was compared to some of the prominent Cu corrosion protection methods in neutral chloride medium reported in the literature. Table 4 shows the corrosion current densities, protection efficiencies (η), and corrosion rates obtained by using different materials and methods such as the direct dissolution of corrosion inhibitors in the corrosive electrolyte, pre-treatment before immersion, self-assembly monolayer

Fig. 10. SEM micrographs of the surface morphology and post corrosion impact (a, b) bare polished Cu (unpeened), (c, d) LSPVC 50 % peened surface, before and after extended immersion duration respectively, in 3.5 % NaCl solution at room temperature under static conditions and (e, f) digital images of the same unpeened and peened surfaces respectively, taken at the end of the immersion experiment.

Fig. 11. SEM micrographs: (a - d) unpeened and (e - h) LSPVC 50 % peened Cu before and after flow corrosion in 3.5 % NaCl solution at room temperature at the rotation speeds of 120, 240 and 360 rpm.

(SAM), metal plating and different types of coatings for Cu protection. In Table 4, the highest protection efficiency (η) (98.7 %) and lowest corrosion current density ($0.06 \mu\text{A cm}^{-2}$) was reported for Cu in 3 % NaCl medium for benzotriazole (BTA) inhibitor via pre-treatment method [57]. By dissolving in the artificial seawater the BTA yielded 95.0 % efficiency [58]. Taking the concentration of the aggressive electrolyte into consideration, in 3.5 % NaCl, the highest efficiency and

lowest corrosion current density of 94.7 % and $0.38 \mu\text{A cm}^{-2}$ respectively, was reported for dipotassium 4,4'-{benzene-1,4-diyl-bis-[(1E)-3-oxoprop-1-ene-1,3-diyl]}-bis-[2-(2H-benzotriazol-2-yl)-phenol ate] (C1) dissolved in the electrolyte [60]. Several other methods achieved lower efficiencies compared to BTA and C1. However, with LSPVC process the performance in 3.5 % NaCl has improved to a value close to that of C1, which is encouraging since BTA

Fig. 12. AFM analysis of the surface topology (a, c) and roughness profile (b, d) of LSPVC 50 % peened and unpeened bare Cu surface after extended immersion in 3.5 % NaCl solution.

Table 4

Comparison chart for the performance of the previous state of the art corrosion inhibitors based on polarization tests for copper in chloride medium with the present research.

Inhibitor	Electrolyte	Method of testing	I_{corr} ($\mu\text{A cm}^{-2}$)	η (%)	CR (mmpy)	Authors	Ref.
Benzotriazole (BTA)	3 % NaCl	Pre-treated	0.06	98.7	–	Notoya et al.	[57]
Benzotriazole (BTA)	Artificial sea water	Dissolved in electrolyte	–	95.0	–	Bokati et al.	[58]
4-Methyl-5-vinylthiazole	3 % NaCl	Dissolved in electrolyte	0.42	94.5	–	Mo et al.	[59]
Dipotassium 4,4'-{benzene-1,4-diyl-bis-[(1E)-3-oxoprop-1-ene-1,3-diyl]}-bis-[2-(2H-benzotriazol-2-yl)-phenol ate] (C1)	3.5 % NaCl	Dissolved in electrolyte	0.38	94.7	–	Jing et al.	[60]
2-mercapto-4-amino-5-nitroso-6-hydroxy pyrimidine (MAP)	3.5 % NaCl	Dissolved in electrolyte	1.25	88.9	–	Khaled	[61]
Gemini surfactant	3.5 % NaCl	Dissolved in electrolyte	0.25	94.5	0.071	Cao et al.	[62]
Olive leaf extract	3 % NaCl	Dissolved in electrolyte	0.60	86.5	0.007	Rahal et al.	[63]
Superhydrophobicelectroactive epoxy	3.5 % NaCl	Conducting polymer coated	4.33	86.8	–	He et al.	[64]
Chitosan	synthetic seawater	Biopolymer in solution	13.70	85.5	–	El Mouaden et al.	[65]
Nicotinamide	3.5 % NaCl	Drug mat. in electrolyte	9.00	80.8	–	Hippolyte et al.	[66]
2,4,6-Trimercapto-1,3,5-triazine	0.3 % NaCl	SAM coated	0.88	92.4	–	Chen et al.	[67]
1-(3-Aminopropyl) imidazole	3 % NaCl	SAM coated	0.26	93.1	–	Durainatarajana et al.	[68]
Heptadecafluoro-1,1,2,2-tetradecyl	3.5 % NaCl	Electro-plating	0.40	–	0.005	Fenghua Su	[69]
Nickael film	3.5 % NaCl	Metal coating	3.00	–	0.035	Zhen Yanga	[70]
CoWO ₄ + TX-100	3.5 % NaCl	Dissolved in electrolyte	0.78	90.0	–	Chukwuike et al.	[30]
Capped NiWO ₄	3.5 % NaCl	Dissolved in electrolyte	0.35	94.3	0.008	Chukwuike et al.	[29]
Laser residual stress	3.5 % NaCl	LSP	0.35	94.5	0.008	Present	–

was carried out in 3 % NaCl. In this work, the corrosion current density of Cu was brought down progressively from $6.31 \mu\text{A cm}^{-2}$ to $0.35 \mu\text{A cm}^{-2}$ with a protection efficiency of 94.45 % by using LSPVC peening process. These results are obtained under static conditions. In the present experiment; we went further to validate the achieved performance of static medium by subjecting the peened surface to a more aggressive flow environment and prolonged immersion tests. The overall results showed that the LSPVC process has the potential of effective corrosion protection of Cu in the chloride medium.

4. Conclusions

Laser shock peening was employed for Cu surface modification. The results showed improved corrosion resistance under static and flow corrosive medium of 3.5 % NaCl at the open atmosphere and room temperature. Thus, the research highlights the following key points:

There was a progressive change in the mechanical properties of Cu with laser light treatment at various overlaps (40, 50, and 70 %). This lead to increased hardness and a decrease in indentation depth penetration as a consequence of comprehensive residual stress from the laser light which improved the compact nature of the Cu surface.

Residual stress does not increase indefinitely with the increase in the laser overlap but there exists an optimal value above or below which the laser light might not impact directly on the residual stress. At a low value the impact energy might be too weak for effective modification while above certain value the energy becomes too high, thus converts to surface damage at the same time the sacrificial layer becomes consumed. In the present research, a 50 % overlap showed the most effective residual stress accumulation without surface damage.

It is possible to conclude that with moderate overlap (50 %) at an appropriate power density (8 Gw/cm^2) in presence of PVC minimal surface roughness can be achieved with LSPVC modification. This resulted in high resistance to corrosion to the tune of $17,020 \Omega \cdot \text{cm}^2$ in 3.5 % NaCl compared to $958 \Omega \cdot \text{cm}^2$ of the unpeened Cu surface. Reduction in the corrosion current density was found from 6.31 to $0.35 \mu\text{A cm}^{-2}$ equivalent to protection efficiency of 94.45 %.

The mechanism of this operation involves the inducement of residual stress to the Cu surface. Shrinking and intertwining of the top-surface structures leading to a more compact surface with minor roughness. This resulted in the increased top-surface hardness and hydrophobicity of Cu leading to a new surface with high corrosion resistance.

Data availability

The raw/processed data required to reproduce these findings cannot be shared at this time as the data also forms part of an on-going study.

Author statement

Author Agreement Statement, We the undersigned declare that this manuscript is original, has not been published before and is not currently being considered for publication elsewhere. We confirm that the manuscript has been read and approved by all named authors and that there are no other persons who satisfied the criteria for authorship but are not listed. We further confirm that the order of authors listed in the manuscript has been approved by all of us. We understand that the Corresponding Author is the sole contact for the Editorial process. He/she is responsible for communicating with the other authors about progress, submissions of revisions and final approval of proofs Signed by all authors as follows:

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Declaration of Competing Interest

The authors declare that there is no conflict of interest regarding the

publication of this paper.

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Appendix A. Supplementary data

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