

Corrosion Analysis of Friction Stir–Welded Brass–Brass and Copper–Copper Samples

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Abstract

Friction stir–welding has become widely practiced in the fabrication of light–weight structures requiring high strength–to–weight ratios and superior corrosion resistance. The friction stir–welding (FSW) process and tool parameters play a key role in determining the joint's characteristics. In this paper, Copper–copper and brass–brass plates were welded using friction stir welding machine at same rotational and transverse speeds. The corrosion resistance of a friction stir–welded copper–copper and brass–brass with parent copper and brass samples respectively were studied via polarization and weight loss method. The microstructures were examined to observe corrosion resistance. Lieca optical microscope used to obtain the microstructures of parent, welded and corroded samples.EDS was conducted to know elements present in the parent metal.

Key words: Friction stir welding, corrosion, polarization, Hardness, microstructure.

Introduction

Friction stir welding avoids formation of the solidification, cracking, porosity and improves the weld properties [1]. Due to great electrical and thermal conductivities brass and copper are widely used in industrial applications [2]. Welding of copper is usually difficult by conventional fusion welding technique because of its high thermal diffusivity and difficulty in fusion welding of brass is zinc which reduces the weldability of brass depending on proportion to the present of zinc in the brass [3]. In friction stir welding the metals are brought in contact with each other due to this frictional heat is generated and the metal reaches in to plastic state and by giving upset pressure weld joints were obtained [4]. Friction stir welding has represented the superior joint strength when joining dissimilar materials [5]. Weld zone of high strength precipitate alloys very high prone to inter granular corrosion on the grain boundaries [6]. Initial surface condition, feed speed and transverse speed have a significant effect on weld quality [7].Corrosion behavior can be investigated by

immersion test and when compared to base material the corrosion resistance of friction stir welds is greatly improved and the weld nugget has the highest corrosion resistance [8]. Corrosion features of weld regions opposed to parent alloy due to welding breakdown and dissolution of inter metallic particles, then increasing in inter metallic particles distributed throughout the matrix of weld regions increases the galvanic corrosion couples and decreasing the grain size in the weld regions [9]. Morphology is strongly affected by applying cooling rate during solidification [9].

Experimental Procedure

Copper and Brass alloy plates (length×breadth×thickness=60×40×6mm) were used in this present work. The chemical compositions of parent species were obtained from energy dispersive X-ray diffraction attached to FESEM (EDX INCA, OXFORD Instruments.) and presented in Table 1 and Table 2. The joints were fabricated by friction stir welding machine Biss-ITS. The tool bit used in this experiment made up of hot die steel– H13, designed as shown in figure 1 and has the hardness of 35.5 HRC. The weld parameters were used as shown in Table 3. Microstructure evaluation is carried out using Leica microscope (Leica Microsystems DMI5000M) and the images were taken in the magnification range of 20X to 50X. For corrosion study potentiodynamic polarization test was conducted as per the standard ASTM G 59–97 using potentiostat (IEC 61326, IVIUM technologies, Netherland) on friction stir welded copper–copper and brass–brass weld joints. The two welded samples (Copper–Copper & Brass–Brass) interface junction was compared with the parent metal in the corrosive environment of 0.2N HCl. Along with this, detailed corrosion study was carried out by immersion method (in 0.2N HCl) as well as salt spray (in 5% NaCl) methods. The parent and welded samples were polished by different grades of emery sheets to remove impurities present on the surface. The samples after corrosion were used as such to study the microstructure which is then compared with parent and welded samples respectively.

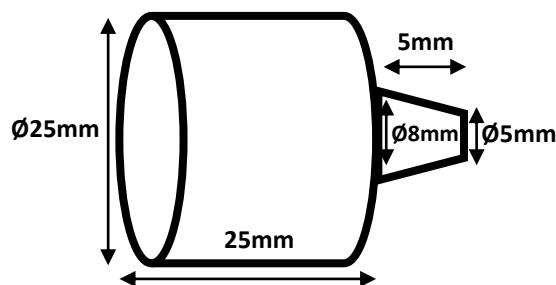


Figure 1. Tool bit dimension used in friction stir welding

Table:-1 Chemical composition of copper sample

Element	Cu	Si	Al	O	C
Wt%	126.83	0.55	0.61	1.35	8.67

Table:-2 Chemical composition of brass

Element	Cu	Zn	Fe	Al	O	C
Wt%	80.54	58.31	0.87	0.47	2.05	12.61

Table:-3 Parameters used for friction welding

Parameters	Upset Pressure (M Pa)	Speed of rotation (rpm)	Feed Speed (mm/min)	Burn off length (mm)
	25	950	80	30

Corrosion study

Potentiodynamic polarization curves (PR)

A typical three-electrode electrochemical corrosion cell was used in all the experiments. A saturated calomel electrode (SCE) was used as reference. All measured potentials were referred to this electrode. A platinum foil was used as the counter electrode. Then four samples parent copper, parent brass, Friction stir welded copper-copper and friction stir welded brass-brass samples were used as working electrodes. All experiments were carried out using ordinario without supporting neither electrolyte nor deaeration. A potentiostat (IEC 61326, IVIUM technologies, Netherland), controlled by a personal computer, equipped with a GPIB card and the commercial software was employed to obtain the potentiodynamic polarization curves. Before each experiment, the open circuit potential (OCP) was recorded for at least 30 min. Polarization curves was obtained potentiodynamically; the linear potential sweep was performed at a ± 248 mV potential window around the measured OCP, from the cathodic to the anodic side, at scan rates of 0.1 and 1 mV s⁻¹.

Salt Spray method (SS)

Salt spray testing was conducted in accordance with ASTM B117 at Assured Testing services, Ridgway, PA. This practice provides a controlled corrosive environment that has been utilized to produce relative corrosion resistance information for specimens of metals exposed in a test chamber. Prediction of performance in natural environments has seldom been correlated with salt spray results when used as stand-alone data. The apparatus for salt spray (fog) exposure consists of a fog chamber, a salt solution reservoir, a supply of suitably conditioned compressed air, one or more atomizing nozzles, specimen supports,

provision for heating the chamber, and necessary means of control. Drops of the solution that accumulate on the ceiling or cover of the chamber shall not be permitted to fall on the specimens being exposed. Drops of solution that fall from the specimens shall not be returned to the solution reservoir for re-spraying. All the four specimens were tested per material/process condition.

Immersion method (IM)

Polished specimens were initially weighed in an electronic balance. Weighed samples are immersed in 100 mL of the acid (0.2 N HCl). After 72 hrs, they are then taken out and then washed thoroughly with tap water, rinsed with distilled water, dried, stored in desiccators and reweighed. From the change in weight of specimens the corrosion rate was calculated using the following relationship,

$$\text{Corrosion Rate} = \frac{534 \times \Delta W}{(A \times T \times D)}$$

ΔW = Loss in weight in mg

A = surface area of the specimen (inch²)

T = Time in hrs

D = Density

Using the above expression we obtained corrosion rates of all four samples shown in Table 6 and Table 9.

Results and Discussion

Corrosion study analysis of Brass Samples

Potentiodynamic polarization method

Polarization resistance measurements are an accurate and rapid way to measure the general corrosion rate. Electrochemical polarization test methods are extremely pertinent for understanding and evaluating the corrosion resistance of materials and the effect of changes in the corrosive environment. They can establish criteria for anodic or cathodic protection and susceptibility to several forms of corrosion.

Figure 2 shows the potentiodynamic polarization curves of brass and friction stir welded brass-brass in 0.2N HCl. Kabasakaloglu et al. [10] have concluded that the corrosion process of brass in HCl media starts at -1.0 V vs SCE, with the subsequent formation of $\text{ZnO}/\text{Zn}(\text{OH})_2$ and zinc ions. According to the authors, dissolution of copper is prevented up to the CuCl formation potential. Peaks observed at very negative potentials, i.e., around

–1.3 V, have been attributed to the formation of ZnO/Zn(OH)₂. However, this peak at very negative potentials has not been much studied yet.

Morales et al. [11, 12], among others, have demonstrated that the dissolution of zinc takes place with anodic polarization even in the passive region. Actually, they showed that the passive layer formed on Cu–Zn alloys in chloride-buffered media consists of a film whose composition is ZnO.xH₂O + Cu₂O + CuO/Cu(OH)₂ + CuCl [13–15]. During the anodic polarization, the dissolution of the passive film occurs above a certain potential, known as the breakdown potential, E_b, which shifts negatively as the chloride concentration increases. Those authors have concluded that brass is less resistant to corrosion than copper. The breakdown of the passivity has been attributed to the formation of soluble species, i.e., chloro- and/or oxycomplexes.

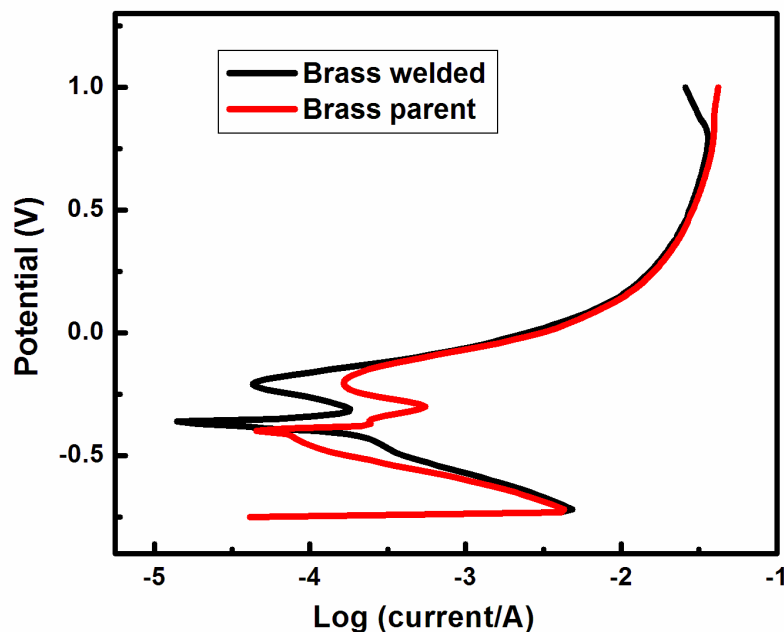
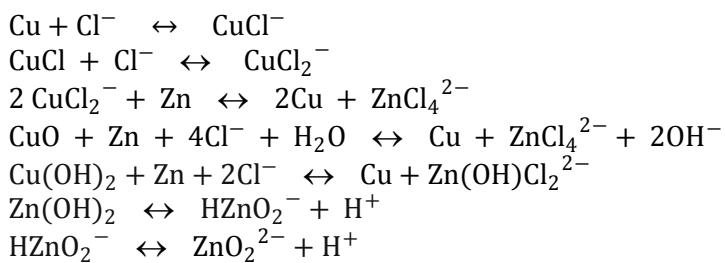
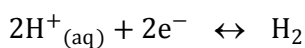


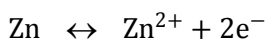
Figure 2. potentiodynamic polarization curves of brass and friction stir welded brass-brass in 0.2N HCl

In the present paper, a comparative study is performed on the voltammetric behavior of brass and friction stir welded brass–brass specimens. Polarization resistance can be related to the rate of general corrosion for metals at or near their corrosion potential, E_{corr} .

The figure 2 clarifies typical Tafel behavior of brass where in active zinc metal in acid solutions, when dissolved in presence of oxygen, both hydrogen evolution and oxygen reduction reactions will be possible. However, in view of the fact that, the saturated solubility of oxygen in pure water at 25°C is only about $10^{-3} \text{ mol dm}^{-3}$ [16] and decreases slightly with increasing the concentration of dissolved salts. In addition, the concentration of H_3O^+ in acid solutions, at $\text{pH} \approx 0$, is high, and since this ion has a high rate of diffusion, consequently, the contribution of the hydrogen evolution reaction at the cathodic process will overcome the oxygen reduction reaction. Therefore, the corrosion of zinc in acid solution proceeds via two partial reactions [17]. The partial cathodic reaction involves evolution of hydrogen gas.



The partial anodic reaction involves the oxidation of Zn and formation of soluble Zn^{2+}



The results of the polarization curve are shown in Table 4 in presence of 0.2N HCl at room temperature. It is observed that the parent brass metal shows high corrosion rate of 0.2549 mm per year while friction stir welded brass imparts 0.1049 mm per year and it has better corrosion resistance. It is also to be noted that the corrosion current is also higher for the parent metal compared to the welded junction. This result indicate that in the friction stir welded brass sample, the weld joint act as a cathode and give better resistance to corrosion when compared to parent brass sample.

Table 4:– Results of the polarization curve of brass samples

Sample	i_{corr} (Corrosion current density in mA/cm^2)	Corrosion rate in mm/year	Corrosion rate in mills/year	Area of the sample in cm^2	Polarisation Resistance R_p
Brass	7.79E-5	0.2549	10.035	1	423.5
FSW Brass–Brass	3.207E-5	0.1049	4.129	1	443.6

Salt Spray method and Immersion Method

The results of salt spray method shown in Table 5 and immersion method in Table 6 are completely in agreement with the polarization data showing that the friction stir welded brass sample shows low corrosion rate compared to parent brass sample

Table 5:- Salt Spray Results

Sample	Medium	$\Delta W(\text{mg})$	$\text{CR} = 534 \Delta W / \text{DAT}$
Brass Sample	5% NaCl	11 mg	27.87 mpy
FSW brass-brass	5% NaCl	3.7 mg	9.299 mpy

Table 6:- Immersion Method Results

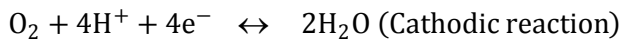
Sample	Medium	$\Delta W(\text{mg})$	$\text{CR} = 534 \Delta W / \text{DAT}$
Brass Sample	0.2 N HCL	130.1 mg	27.47 mpy
FSW brass-brass	0.2 N HCL	79.4 mg	16.63 mpy

Corrosion study analysis of Copper Samples

Figure 3 shows the potentiodynamic polarization curves of copper samples in 0.2N HCl. The electrochemical potentiodynamic polarization parameters, i.e. corrosion potential (E_{corr}) and the corrosion current density (i_{corr}), obtained from the intersection of the anodic and cathodic Tafel lines with the corresponding corrosion potential are given in Table 7. It has been shown that in the Tafel extrapolation method, use of both the anodic and cathodic Tafel regions is undoubtedly preferred over the use of only one Tafel region [18]. The corrosion rate can also be determined by Tafel extrapolation of either the cathodic or anodic polarization curve alone. If only one polarization curve alone is used, it is generally the cathodic curve which usually produces a longer and better defined Tafel region. Anodic polarization may sometimes produce concentration effects, due to passivation and dissolution, as well as roughening of the surface which can lead to deviations from Tafel behavior [19].

The cathodic parts of the polarization curves show limiting current corresponds to the oxygen reduction reaction. This indicates that the cathodic process is controlled by diffusion of oxygen gas from the bulk solution to the metal surface. This behavior is well known since copper can hardly be corroded in the deoxygenated dilute hydrochloric acid [20], as copper cannot displace hydrogen from acid solutions according to theories of chemical thermodynamics. However, in aerated hydrochloric acid, dissolved oxygen is reduced on copper surface and this will enable some corrosion to take place [21]. Cathodic reduction of oxygen can be expressed either by two consecutive $2e^-$ steps involving a

reduction to hydrogen peroxide first followed by a further reduction to water or by a direct 4e⁻ transfer step [22] as shown by equation.



The partial anodic reaction involves the oxidation of Cu and formation of soluble Cu²⁺
 $\text{Cu} \leftrightarrow \text{Cu}^{2+} + 2\text{e}^- \text{ (Anodic reaction)}$

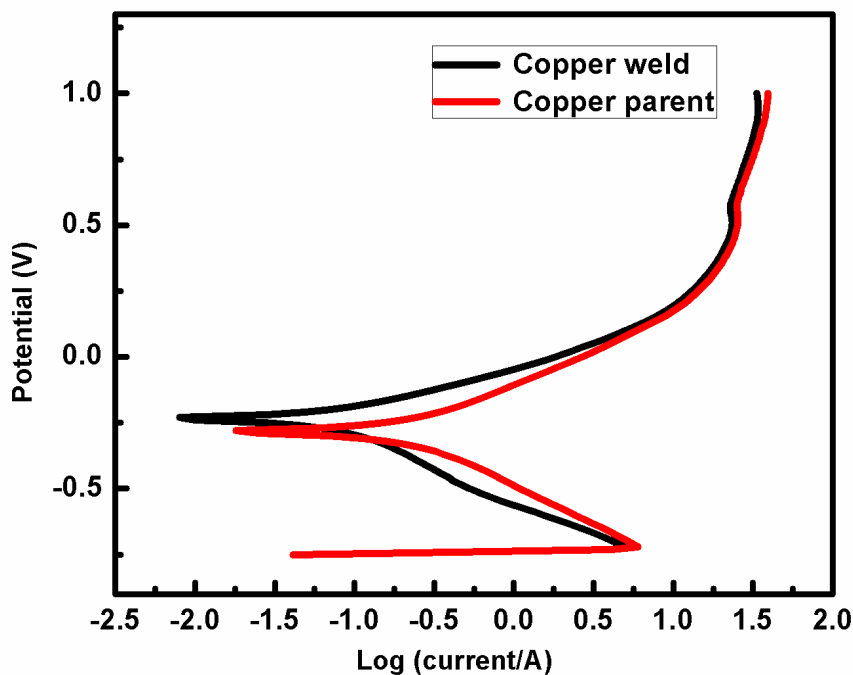


Figure 3. Potentiodynamic polarization curves of copper and friction stir welded copper-copper in 0.2N HCl

The results of the polarization curve are shown in Table 7 in presence of 0.2N HCl at room temperature. It is observed that the parent copper metal shows high corrosion rate of 0.01183 mm per year while friction stir welded copper imparts 0.0001228 mm per year and it has better corrosion resistance. It is also to be noted that the corrosion current is also higher for the parent metal compared to the welded junction. This result indicate that in the friction stir welded copper sample, the weld joint act as a cathode and give better resistance to corrosion when compared to parent copper sample.

Table 7:- Results of the polarization curve of copper samples

Sample	i_{corr} (Corrosion current in mA/Cm ²)	Corrosion rate in mm/year	Corrosion rate in mills/year	Area of the sample in cm ²	Polarisation Resistance R_p
Copper	3.615 E-6	0.01183	0.46	1	8477
FSW copper-copper	3.754E-8	0.0001228	0.0048	1	8.331E5

Salt Spray method and Immersion Method

The results of salt spray method shown in Table 8 and immersion method in Table 9 are completely in agreement with the polarization data showing that the friction stir welded copper sample shows low corrosion rate compared to parent copper sample

Table 8:- Salt Spray Results

Sample	Medium	ΔW (mg)	CR=534 ΔW /DAT
Copper sample	5% NaCl	87	295.53mpy
Friction stir welded Copper-Copper	5% NaCl	93	176.02mpy

Table 9:- Immersion Method Results

Sample	Medium	ΔW (mg)	CR=534 ΔW /DAT
Copper Sample	0.2 N HCL	570.0	216.19 mpy
Friction stir welded Copper-Copper	0.2 N HCL	587.8	92.81 mpy

In both the cases the weld metal (core zone) region, the metal behaves more corrosion resistance compared to parent material.

Microstructure Analysis

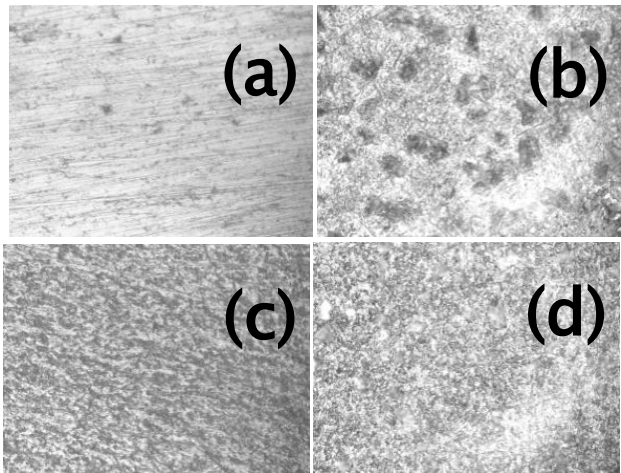


Figure 4. Microstructure of (a) brass parent (b) brass parent corroded (c) brass–brass weld and (d) brass–brass weld corroded samples.

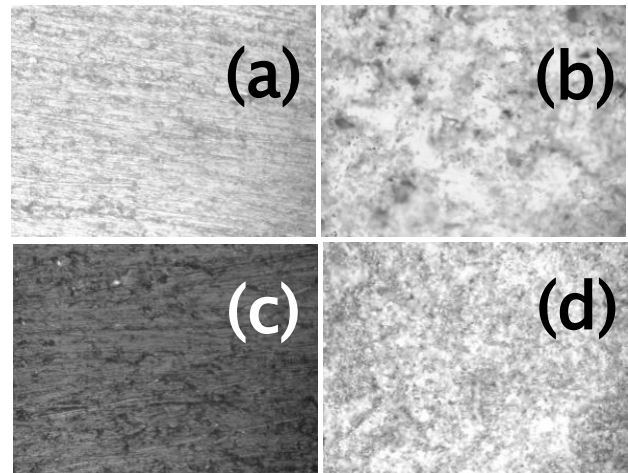


Figure 5. Microstructure of (a) copper parent (b) copper parent corroded (c) copper–copper weld and (d) copper–copper weld corroded samples.

The microstructural characterizations of the 950 rpm FSW conditions of both brass–brass and copper–copper samples were presented in Figure 4 (c) & Figure 5 (c) respectively. One can clearly observe joint has no defects or any other imperfections (lack of penetration, porosity etc.).

Comparison of Figure 4 (a) & 4 (c) (brass before FSW & after FSW) and Figure 5 (a) & 5 (c) (Copper before FSW & after FSW) clearly indicates the sample after FSW shows uniform grain size distribution.

After the corrosion studies were conducted for brass samples, Figure 4 (b) (parent brass after corrosion) shows high dominating pits indicating high corrosion damage whereas FSW brass–brass sample (Figure 4 (d)) shows comparatively very less number of pits after corrosion indicating a low corrosion rate.

Copper samples on observed after the corrosion studies shows, Figure 5 (b) (parent copper after corrosion) shows large number of pits indicating high corrosion damage whereas FSW copper–copper sample (Figure 5 (d)) shows comparatively very less number of pits after corrosion indicating a low corrosion rate.

Table 10:– Hardness studies

Sample	Copper	Copper weld	Brass	Brass weld
Hardness (HRB)	73.75	76	30	53.5

Vickers Hardness values of parent and welded samples were shown in table 10 and it is observed hardness increased in welded samples as compared with parent samples due to high temperature generation at the time of FSW and decreasing of grain size as shown in Figure 4 (c) and 5 (c).

Conclusion

The brass–brass and copper–copper samples were successfully welded with friction stir welding without failures or imperfections in the joints. On comparison with parent samples the Vickers hardness values showed an increment. Corrosion studies shows that the FSW samples are less prone to corrosion than parent samples. Moreover it is also observed from corrosion studies that the brass samples are more corroded when compared with copper samples and the reason for this can be attributed as dealloying in brass.

References

- [1] D. Raghu Rama, Muruganandan and L K Kuruswamy, *Int. J. Chem. Tech. Res*, 6, 4, pp 2577–2582, 2014.
- [2] Dhananjayulu Avula, R K Raj Singh, D K Dwivedi, N K Mehta, *World acad. Sci. Engg. Tech.*, 5, pp 02–22, 2011.
- [3] Ashok Kumar, R Shukla and A Venkatachalam, *Rasayan J. Chem.*, 6, 1, pp 12–14, 2013.
- [4] M S Moghaddam, R Parvizi, M Haddad–Sabzevar, A Davoodi, *Materials and Design*, 32, pp 2749–2755, 2011.
- [5] I Bhamji, R J Moat, M Preuss, P L Threadgill, A C Addison, M J Peel, *Sci. Tech. Weld. Join.* 17, 4, pp 314–320, 2012.
- [6] G Cam, S Misticoglu, M Pakdil, *Weld. J.*, 88, pp 228–232, 2009.
- [7] R A Bell, J C Lippold, D R Adolphson, *Weld. Res.* pp 325–332, 1984.
- [8] Hai-long Qin, Hua Zhang, Da-tong Sun and Quin-Yu Zhuang, *Int. J. Min. Met. Mat.*, 22, 6, pp 627–638, 2015.
- [9] F Gharavi, K A Matori, R Yunus, N K Othman and F Fadaeifard, *J. Mat Res Tech.*, 4, 3, pp 314–322, 2015.
- [10] Kabasakaloglu M, Kryak T, Sendil O, Asan A, *Appl Surf Sci.*, 193, 1–4, pp 167–174, 2002.
- [11] Morales J, Fernandez GT, Esparza P, Gonzalez S, Salvarezza RC, Arvia AJ, *Corros. Sci* 37, 2, pp211–225, 1995.
- [12] Morales J, Fernandez GT, Gonzalez S, Esparza P, Salvarezza RC, Arvia AJ, *Corros Sci* 40, 2–3, pp 177–190, 1998.
- [13] Milosev I, MikicKT, Gaberscek M, *Electrochim Acta* 52, 2, pp 415–426, 2006.
- [14] Morales J, Esparza P, Fernandez GT, Gonzalez S, Gracia JE, Caceres J, Salvarezza RC, Arvia AJ, *Corros Sci.*, 37, 2, pp 231–239, 1995.
- [15] Sinapi F, Deroubaix S, Pirlot C, Delhalle J, Mekhalif Z, *Electrochim Acta*, 49, 17–18 pp 2987–2996, 2004.
- [16] E.E. Foad El-Sherbini., S.M. Abdel Wahaab., M. Deyab. *Mater. Chem. and Phys.*, 89,2–3, pp183–191, 2005.
- [17] P. Jinturkar, Y.C. Guan, K.N. Han, *Corrosion*, 54, 2, pp 106–114, 1984.
- [18] R. Caban, T.W. Chapman., *J. Electrochem. Soc.*, 124, 9, pp 1371–1379, 1977.
- [19] L. L. Shreir, R. A. Jarman, G. T. Burstein, *CORROSION*, Vol 1, Metal/Environment
- [20] ZView2 help, Scribner Associates, 2000.
- [21] H.Ma, Sh.Chen, B. Yin, Sh. Zhao, X. Liu, *J. Corros. Sci.*, 45, 5, pp 867–882, 2003.
- [22] A.H. Moreira, A.V. Benedetti, P.L. Calot, P.T.A. Sumodja., *Electrochim. Acta.*, 38, 7, pp 981–987, 1993.
- [23] *Reactions*, Butterworth–Heinemann, Oxford, 3rd edition, 2000.