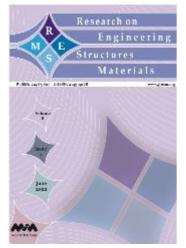


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Research Article

### Effects of immersion in the NaCl and H<sub>2</sub>SO<sub>4</sub> solutions on the corrosion rate, microstructure, and hardness of stainless steel 316L

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Article Info	Abstract
Article history:	Stainless steel is one of the materials with excellent corrosion resistance, so it is widely applied in various fields such as the chemical industry, aerospace, power
Received 20 Feb 2023 Accepted 16 May 2023	generation, and biomedicine. The environmental condition (neutral, acidic, salty, or basic) is the most important thing to be considered prior to choosing palpable stainless steel materials. In this study, the corrosion behaviors of stainless-steel type SS 316L under exposure immersions in salty and acidic environments using
Keywords:	sodium chloride (NaCl) and sulfuric acid (H <sub>2</sub> SO <sub>4</sub> ) solutions were investigated, by referring to a standard method of ASTM G31-72. Scanning electron microscopy
Corrosion rate; Stainless steel 316L; Corrosive medium; Microstructure; Material resistance	(SEM) results revealed that the specimen in NaCl solution exhibited a darker color, and relatively similar shape of cracks are observed. In contrast, the specimen immersed in $H_2SO_4$ was observed to have a large diagonal crack. The corrosion rate of SS 316L immersed in NaCl was lower with a value of 0.4260 mpy, compared to that of $H_2SO_4$ which was 2.5141 mpy. The Vickers hardness test showed that the hardness value of NaCl immersion was higher than that of $H_2SO_4$ , which was 6.34% and 1.94%, respectively. It can be concluded that the SS 316L is better used in a salty environment compared to acidic conditions, because the main function of SS 316L is to be able to withstand rust, one of which is caused by chloride compared to other types of stainless-steel.

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#### 1. Introduction

Austenitic stainless steel is a common material that has good corrosion resistance, excellent mechanical strength, low thermal expansion, lower process costs, and good reliability [1]. Most manufacturing industries use this material as the primary material for making their products. Stainless Steel AISI 316L (SS 316L) is one of the austenitic stainless steel materials with excellent corrosion resistance, so it is widely applied in various fields such as the chemical industry, aerospace, power generation, and biomedicine [2]. This material is also recommended as a biomaterial due to its low carbon content, which increases its corrosion resistance to chlorinated environments [3,4]. The passive protective layer on the surface of SS material provides high prevention against uniform corrosion in ordinary oxidizing environments but can induce localized forms of corrosion under special conditions [5–7]. The existence of corrosive substances can lead to local passivation degradation and dangerous pitting [8–10]. The exposure of austenitic stainless steel 316L to the high temperature followed by slow cooling in air, such as, during welding, is corresponding to their vulnerability to another type of localized intergranular corrosion. In addition, low-chromium regions are the preferred pathways for crack propagation under other localized corrosion attacks and tensile stresses [11,12].

This type of stainless steel is used in medical devices such as implants [13]. Moreover, the high corrosion resistance of SS 316L provides by a thin chromium oxide film that has high stability against corrosion [14,15]. However, this SS 316L material still has some shortcomings; for example, when SS 316L is used as one of the materials for shipbuilding. Marine vessels will come into contact with the marine environment, which contains sodium chloride (NaCl) salt. Therefore this material often experiences pitting-type corrosion due to exposure to seawater and through the deposition of salt in the air, which has high humidity and high chloride concentration [16]. SS 316L is also sensitive to chloride (Cl) compounds in the environment, affecting this material's corrosion resistance. The presence of chloride compounds in the environment can corrode SS 316L and affect its service life.

Recently, the relationship between extreme environments and the pitting corrosion tendency of stainless steel materials has been the field of numerous investigations. Based on the available literature, pH, temperature, and types of the environment play a critical role in the corrosion susceptibility of stainless steel materials [17–19]. A combination of pickling and passivation in aqueous solutions of hydrochloric and nitric acids has been found to be the best choice for rust removal [20]. Several studies have reported the role of MnS inclusions in initiating pitting corrosion in sensitized austenitic SS. Another study compared the effect of different mechanical surface treatments on the pitting resistance of sensitized SS 304 in a 3.5 wt% of NaCl solution [18]. Immersion method has been popular to investigate the corrosion and durability property of stainless-steels. Lieth et al investigated effects of heat treatment on different water condition i.e., sea water, fresh water, and crude oil by immersion method. They found that the severe corrosion occurred in the sea water at high temperature [21]. Another study by Ji et al found that the higher the concentration of the mixed solution and the higher the test temperature, the greater the corrosion rate of 316L stainless steel wire and the lower the breaking strength of 316L stainless steel wire [22]. Adnan et al observed that heat treatment affects the corrosion rate. The annealed sample showed the lowest corrosion rate, while the quenched sample seemed to corrode more compared to the other samples [23]. Moreover, Reda and coworkers have reported some important studies regarding the effect of several environmental aspects on the microstructure of stainless-steel materials have been investigated. They found that hydrogen caused the corrosion and alter the microstructure of electroplated steel alloy 4130 by diffusion mechanism [24]. Next, they also found that heating and solution treatment caused a hardening effect that improved the mechanical and microstructure properties of aluminum-zinc alloy 7075 [25]. Similarly, the heating and solution treatment has been employed on aluminum-copper 2024 alloy, which found that the heating and solution treatment increases the mechanical properties of the steel by short-distance diffusion coupled with precipitation mechanisms [26].

Several studies have been carried out to investigate the resistance of stainless steel materials in solution treatments, including the corrosion rate and hardness of SS 304L in a 0.01M NaCl medium [27]. Improvement of corrosion properties of nanostructured SS 304L in 0.6 M NaCl medium and the effect of incorporation of molybdenum Disulfide (MoS<sub>2</sub>) in organic coatings on the corrosion resistance of stainless steel in 3.5% NaCl medium[28]. Furthermore, several other studies also discuss the resistance of 316L when immersed in an  $H_2SO_4$  medium, including the corrosion rate of SS 316L in an  $H_2SO_4$  medium containing the amino acid methionine [29] and the corrosion rate of SS 316L in an  $H_2SO_4$  medium with various temperature treatments [30].

According to the literature, it is reported that the environmental condition (neutral, acidic, salty, or basic) is the most important thing to be considered prior to choose the right stainless steel materials [8,20,31,32]. Nevertheless, the comparison study regarding the corrosion properties of stainless steel, specifically SS 316L, under different conditions is

extremely limited; thus, a comprehensive study that provides a comparison of different environmental conditions on the corrosion behavior of stainless steel materials is urgently needed. Therefore, in this study, the corrosion behaviors of SS 316L under exposure immersions in salty and acidic environments using NaCl and H<sub>2</sub>SO<sub>4</sub> solutions were investigated. The ultimate goal of this study is to investigate the effects of acidic and salty conditions on the corrosion behavior of SS 316L. The specimen of SS 316L was immersed in NaCl and H<sub>2</sub>SO<sub>4</sub> under the same condition, by referring to a standard method of ASTM G31-72. The corrosion behavior was determined using the corrosion rate and mass loss method. Microstructures of SS 316L of before/after exposure immersion experiment were analyzed using scanning electron microscopy (SEM). Meanwhile, the chemical composition of the specimen was analyzed using energy dispersive spectroscopy (EDS). The mechanical properties of SS 316L before/after treatment were assessed using the Vickers hardness test. Finally, the results of this study was expected to provide recommendations for industry in selecting the right condition for a durable and corrosion-resistant use of SS 316L.

#### 2. Materials and Method

#### 2.1. Materials

A standard stainless steel 316L (SS 316L) plate was supplied from Bozhong Metal Groups, China, which fulfilled the standard specification of SS 316L by ASTM A240 and JIS G4304. The standard composition and mechanical properties of SS 316L that was used in this study are summarized in Table 1 and standard mechanical properties of SS 316L are listed in Table 2. Sodium chloride (NaCl, 99%), sodium hydroxide (NaOH, 99%), and sulfuric acid (H<sub>2</sub>SO<sub>4</sub>, 98%) that were used for the immersion experiment were provided by Merck, Germany. Deionized (DI) water was always used in chemical solution preparation and all experimentation.

Elementals	Composition (wt%)	
C (Carbon)	≤ 0.03	
Si (Silicon)	≤ 1.00	
Mn (Manganese)	≤ 2.00	
P (Phosphorous)	≤ 0.045	
S (Sulfur)	≤ 0.045	
Cr (Chromium)	16.0 - 18.0	
Mo (Molybdenum)	2.0	
Ni (Nickel)	10.0	

Table 1. Standard additive composition in SS 316L [33,34].

Table 2. Standard mechanical properties of SS 316L [35,36].

Mechanical Properties	Unit	Value
Yield strength	MPa	≥ 200
Tensile strength	MPa	≥ 480
Elongation at break	%	≥ 50
Hardness	HV	≤ 180

#### 2.2. Corrosive Medium Immersion Test

The corrosive medium immersion test was conducted by adapting a standardized method of ASTM G31-72 [37]. This method is commonly used in the determination of laboratory corrosion testing for metals. In this experiment, NaCl and  $H_2SO_4$  were used as the immersion medium that simply represented the salty and acid environments, respectively.

Firstly, 100 mL of NaCl and H<sub>2</sub>SO<sub>4</sub> solutions with the same molar concentration (0.017 M) were prepared in a different compartment. After that, a rectangular shape of SS 316L was prepared with a dimension of 20 mm x 10 mm x 2 mm and hanged with a inert rope [38,39]. Fig. 1 depicts the schematic representation of corrosive medium immersion test. The initial weight of the plate was weighed using a precision analytical balance (Precisa Series 390, USA) with an accuracy of 4 significant figure. Then, the SS 316L plates were immersed in both solutions and were left for three days. After three days, the SS 316L samples were taken from the solution and washed with DI water to clean the spent solution [37,39]. The cleansed samples were dried in an oven for four hours until they were completely dried. The samples were put into a desiccator before being weighed, which is noted as the final weight after the corrosion test. Finally, the corrosion rate can be simply determined using Eq. 1 [40,41].

$$C_{R} = \frac{\left(W_{i} - W_{f}\right) \times K}{\rho \times A \times t}$$
<sup>(1)</sup>

Where  $C_R$  represents the corrosion rate (mils per year/mpy),  $W_i$  and  $W_f$  are the initial and the final weight (mg) of the sample, respectively. *K* is a constant in the corrosion rate equation (534), and  $\rho$  shows the density of the immersion solution (g/cm<sup>3</sup>). *A* is the total surface area of the sample (cm<sup>2</sup>), and t is the time of exposure (h).

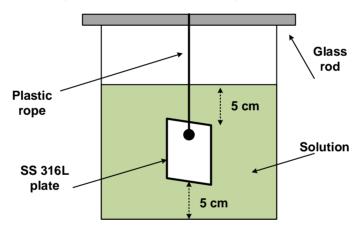


Fig. 1 Schematic illustration of corrosion immersion experiment

#### 2.3. Characterization

Scanning electron microscopy (SEM, Thermo-Fischer Scientific, USA) was performed to capture the surface morphology of the pristine SS 316L and samples after going through the corrosion immersion test. This instrument can provide a surface picture of the samples with some magnifications. To further understand the effects of corrosive medium immersion on the chemical composition of the SS 316L, an energy dispersive spectroscopy (EDS, Thermo-Fischer Scientific, USA) was used. This analysis can show the chemical entities of the samples, which corresponds to the SEM images [42–44]. Therefore, some new insights regarding the effects of corrosive medium immersion using NaCl and  $H_2SO_4$  on the microstructure of SS 316L can be achieved.

#### 2.4. Mechanical Strength Evaluation Using Vickers Hardness Test

Vickers hardness test method was used to evaluate the mechanical strength of the SS 316L before and after corrosion immersion tests. The basic concept of this test is to test the

hardness of the material with a stacker. The shape of this pounder is like a pyramid and is made of square diamonds. The angle produced by Vickers's eye is 136°[45]. To describe the hardness of the sample, Vickers hardness number (VHN) is used, as can be expressed in Eq. 2 [46,47].

$$VHN = \frac{1.854 \times P}{d^2} \tag{2}$$

Where *VHN* stands for Vickers hardness number (kg/ $\mu$ m<sup>2</sup>). *P* is the applied load (kg), and *d* is the average diagonal length ( $\mu$ m).

#### 3. Results and Discussion

# 3.1. Effects of Immersion in NaCl and $\rm H_2SO_4$ on The Mass Reduction and Corrosion Rate of SS 316L

Effects of different types of corrosive medium, which are salty and acidic conditions, on the SS 316L sample were investigated. Table 3 summarizes the data regarding the mass reduction and corrosion rate of SS 316L after being immersed in NaCl and H<sub>2</sub>SO<sub>4</sub> solution. According to Table 3, it can be seen that initially, the SS 316L samples had a mass of 2.9796 g and 2.9938 g before being immersed in NaCl and  $H_2SO_4$  solutions. The samples were left for 72 h to assess the corrosion process of SS 316L in salty and acidic conditions. After 72 h, it can be observed that the final weights of both samples are 2.9795 g for immersion in NaCl and 2.9933 g for  $H_2SO_4$ . Hence, the weight difference can be simply computed by subtracting the initial and final weights of the samples. It was found that the SS 316L exhibited 0.0001 g of weight loss due to immersion in the NaCl solution. Moreover, the SS 316L possessed a more dramatic weight loss of 0.0005 g when the sample was immersed in the H<sub>2</sub>SO<sub>4</sub> solution. The occurrence of weight loss can be due to the reaction between the dominant metal in the SS 316L, which is iron (Fe), and the corrosive solutions. The greater mass reduction represents the greater loss of Fe that can be extracted from the body of SS 316L [48]. The sample that was immersed in  $H_2SO_4$  exhibited greater mass loss. It is explainable that the corrosion reaction in this condition occurs in two ways, namely, the redox corrosion reaction between iron and oxygen and the direct acid leaching due to a large number of H<sup>+</sup> ions [49]. The reactions can be written as follows [48,50].

Redox reaction mechanisms;

Oxidation reaction:

$$Fe_{(s)} \to Fe_{(aq)}^{2+} + 2e^{-}$$
 (3)  
 $E^{o} = +0.44 V$ 

**Reduction reaction:** 

$$O_{2(aq)} + 2H_2O_{(aq)} + 4e^- \to 4OH_{aq}^- \qquad E^o = +0.40 V \tag{4}$$

Overall reaction:

$$2Fe_{(s)} + O_{2(aq)} + 2H_2O_{(aq)} \to 2Fe_{(aq)}^{2+} + 4OH_{(aq)}^{-} \qquad E^o = +0.84 V$$
(5)

Acid leaching reaction:

$$Fe_{(s)} + H_2SO_{4(aq)} \rightarrow 2H^+_{(aq)} + FeSO_{4(aq)}$$
<sup>(6)</sup>

On the other hand, the sample that was immersed in NaCl solution seems to have a lower mass loss compared to that of  $H_2SO_4$ . It is clearly due to the corrosion reaction that occurred in the NaCl system being just the redox mechanisms (see Eq. 3-5); thus, the mass loss due to corrosion is found lower. Corroboratively, the measured corrosion rate of the SS 316L in  $H_2SO_4$  is way much faster than that of NaCl, with the exact values of 2.5141 mpy and 0.4260 mpy, respectively. The corrosion rate in  $H_2SO_4$  is 5.9 times higher than that in NaCl. Fortunately, the corrosion rate for SS 316L is still acceptable when it falls in the range of 1 - 5 mpy, and when the corrosion rate is < 1 mpy, it is called exceptional [51,52]. Therefore, these results indicate that SS 316L is better used for equipment in a salty environment compared to the acidic one.

Table 3. The mass reduction and corrosion rate of SS 316L after being immersed in NaCl and  $H_2SO_4$  solutions.

Type of solution	Starting date	Finishing date	Duration (h)	Initial weight W <sub>i</sub> (g)	Final weight W <sub>f</sub> (g)	Weight difference ΔW (g)	Corrosion rate (mpy)
NaCl	2 September 2022	5 September 2022	72	2.9796	2.9795	0.0001	0.4266
$H_2SO_4$	2 September 2022	5 September 2022	72	2.9938	2.9933	0.0005	2.5176

# 3.2. Effects of Immersion in Corrosive Medium on The Surface Micromorphology and Chemical Composition of SS 316L

Fig. 2 shows the SEM images and EDS spectra of pristine SS 316L without immersion treatment. It can be seen that the surface micromorphology of the sample is quite uniform without any sign of corrosion. Meanwhile, The SEM test on the sample of SS 316L that was immersed in NaCl solution obtained a hollow structure in stainless steel as a result of immersion for three days, as can be seen in Fig. 3. After experiencing treatment, the material has defects in the microstructure of the material, which can further affect the service life of the material's properties. This also causes a reduction in the mass of the material because cracks and degradation occurred on the surface. This was also reported in a study, they explained that there were long depressions in the diagonal direction after being immersed in the corrosive solution [53]. Moreover, another finding can be observed in Fig. 2 that when stainless steel material is immersed in NaCl solution, some crystal constituents were formed on the surface of the material. The circle image shown is the form of crystallization that occurs when immersing SS 316L in NaCl solution. The type of corrosion occurred in immersion into NaCl solution is uniform corrosion [54].

Moreover, SEM results also show that the corrosion that occurs on SS 316L when immersed in  $H_2SO_4$  solution possesses worse corrosion, with uniform defects observed on the surface of SS 316L as can be seen in Fig. 4. Corrosion occurs almost all over the surface of the specimen, and there are the same depressions and cracks, indicating that the type of corrosion that occurs is a type of uniform corrosion [55–57]. Almost the entire top surface of stainless steel formed new crystals and black spots. However, what distinguishes between the immersion of the two solutions is the erosion of the material when immersed in  $H_2SO_4$  so that in general the hardness value of the specimen immersed in  $H_2SO_4$  is much lower than the specimen immersed in  $H_2SO_4$  solution than in SS 316L specimens immersed in  $H_2SO_4$  solution than in SS 316L specimens immersed in  $H_2SO_4$  is much lower than the specimen immersed in  $H_2SO_4$  solution than in SS 316L specimens immersed in  $H_2SO_4$  is much lower than the specimen immersed in  $H_2SO_4$  solution for the specimen immersed in  $H_2SO_4$  is much lower than the specimen immersed in  $H_2SO_4$  is much lower than the specimen immersed in  $H_2SO_4$  is much lower than the specimen immersed in  $H_2SO_4$  is much lower than the specimen immersed in  $H_2SO_4$  is much lower than the specimen immersed in  $H_2SO_4$  is much lower than the specimen immersed in  $H_2SO_4$  is much lower than the specimen immersed in  $H_2SO_4$  solution for  $H_2SO_4$  is much lower than the specimen immersed in  $H_2SO_4$  solution  $H_2SO_4$  is much lower than the specimen immersed in  $H_2S$ 

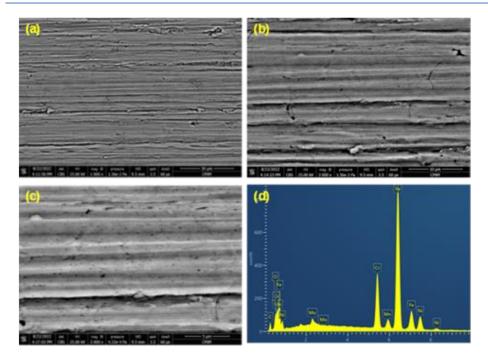


Fig. 2 SEM images of the pristine SS 316L under magnifications of (a) 1000x, (b) 3000x, and (c) 5000x. (d) EDS spectra of pristine SS 316L

The chemical composition of pristine SS 316L and the immersed SS 316L samples in NaCl and  $H_2SO_4$  was analyzed using EDS. Table 4 summarizes the elemental composition of the samples. It is found that the pristine SS 316L consists of 68.1, 16.5, 10.3, 2.9, 1.2, and 1.0% of Fe, Cr, Ni, O, S, and Mn, respectively, which is in similar manner with previous studies [33,34]. While for the sample that immersed in NaCl exhibited compositions of 66.9, 16.4, 10.3, 4.1, 1.2, and 1.1 wt% for Fe, Cr, Ni, O, S, and Mn, respectively. While the sample immersed in H<sub>2</sub>SO<sub>4</sub> solution exhibited compositions of 65.7, 15.6, 9.2, 6.5, 1.7, and 1.3 wt%, for Fe, Cr, Ni, O, S, and Mn, respectively. Based on the results, it is found that the immersion process reduced the Fe content and increased the oxygen content. This process followed the corrosion reaction mechanisms as discussed perviously. It is also clearly observed that H<sub>2</sub>SO<sub>4</sub> caused worse corrosion on the main metals that constructed the SS 316L, such as Fe, Cr, and Ni, compared to that of in NaCl solution. It can be due to a strong electrochemical reaction between  $H_2SO_4$  and the metals [58,59], as previously discussed in section 3.1. It is also evident by the SEM images that a sample of SS 316L in H<sub>2</sub>SO<sub>4</sub> exhibited more defects in its microstructure compared to that of NaCl. The reduction content of constituent metals can alter the mechanical strength of the SS 316L sample [60]. Therefore, it can be concluded that H<sub>2</sub>SO<sub>4</sub> caused more serious corrosion activity on SS 316L compared to NaCl.

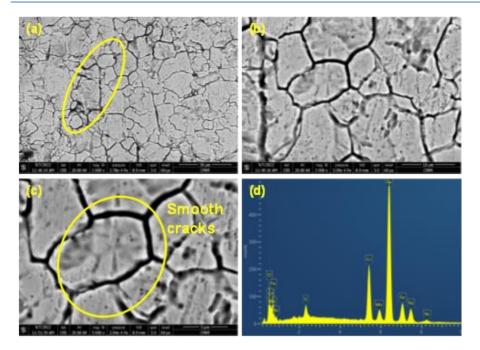


Fig. 3 SEM images of SS 316L sample after immersed in NaCl solution under magnifications of (a) 1000x, (b) 3000x, and (c) 5000x. EDS spectra of SS 316L sample after immersed in NaCl solution

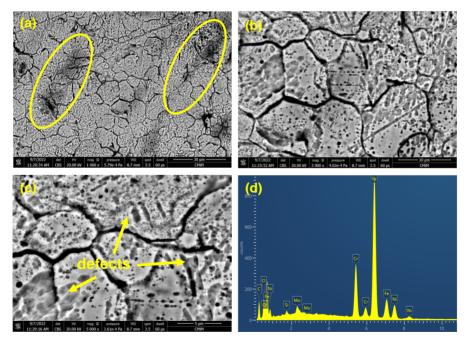


Fig. 4 SEM images of SS 316L sample after immersed in  $H_2SO_4$  solution under magnifications of (a) 1000x, (b) 3000x, and (c) 5000x. EDS spectra of SS 316L sample after immersed in  $H_2SO_4$  solution.

Elementals	Composition (wt%)			
	Pristine SS 316L	SS 316L (NaCl)	SS 316L (H <sub>2</sub> SO <sub>4</sub> )	
Iron (Fe)	68.1	66.9	65.7	
Chromium (Cr)	16.5	16.4	15.6	
Nickel (Ni)	10.3	10.3	9.2	
Oxygen (0)	2.9	4.1	6.5	
Sulfur (S)	1.2	1.2	1.7	
Manganese (Mn)	1.0	1.1	1.3	

Table 4. Chemical compositions of the SS 316L samples after being immersed in NaCl and  $H_2SO_4$  solutions.

## 3.3. Effects of Immersion in Corrosive Medium on The Hardness Level of SS 316L

Effects of immersion in NaCl and H<sub>2</sub>SO<sub>4</sub> solutions on the mechanical strength of SS 316L were assessed through the Vickers hardness test. Fig. 5 shows the results of Vickers hardness numbers (VHN) for three samples, namely pristine SS 316L, SS 316L (H2SO4), and SS 316L (NaCl). It can be seen that the pristine SS 316L exhibited a VHN of 159.3  $\pm$  1.32 kg/µm<sup>2</sup>. The immersed products showed VHN numbers of 169.4  $\pm$  0.52 kg/µm<sup>2</sup> for SS 316L (NaCl) and 162.4  $\pm$  0.47 kg/µm<sup>2</sup> for SS 316L (H<sub>2</sub>SO<sub>4</sub>). It is found that the immersed samples have higher VHN values compared to the pristine SS 316L. It is explainable by the formation of a brittle surface of SS 316L due to the corrosion reactions on the surface. This corroded surface attributes more brittle property due to the oxidation, thus, the immersed samples attributed higher VHN values [61,62]. The HV value increased by 6.34% for materials soaked in NaCl solution, and the value obtained from materials soaked in H<sub>2</sub>SO<sub>4</sub>.

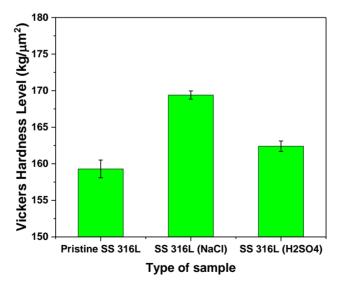


Fig. 5 Results of the hardness of the tested SS 316L samples.

The defect-free corrosion surface can be attributed to enhancing the mechanical strength of the SS 316L sample in NaCl. The generated crystals on the surface of SS 316L can also provide higher mechanical strength [59,63]. The hardness value of SS 316L increased by 6.34% for those immersed in NaCl solution. Therefore, SS 316L itself is widely used for the manufacture of equipment that is in direct contact with salt solutions because the main function of SS 316L is to be able to withstand rust, one of which is caused by chloride [64].

#### 3.4. Effect of pH Variations

Effect of pH variations on the corrosion rate and hardness of the immersed SS 316L was investigated and the results are depicted on Fig. 6. Based on Fig. 6 it can be seen that the corrosion rate of SS 316L decreased significantly with a rise of pH. pH of solution was controlled using NaOH and H<sub>2</sub>SO<sub>4</sub>. Sample of SS 316L at lowest pH of 1 experienced the highest corrosion rate of 2.5563 mpy, while at pH of 7 and above the corrosion rate remained constant at 0.5317 mpy. pH is a measure of the acidity or basicity of a solution and plays a crucial role in the corrosion of stainless steel. The corrosion of stainless steel is an electrochemical process that involves the transfer of electrons between the metal and its environment. At low pH levels, the concentration of hydrogen ions (H<sup>+</sup>) in the solution increases, leading to an increase in the corrosion rate of stainless steel [65]. Conversely, at higher pH levels, the concentration of H+ ions decrease, resulting in a decrease in the corrosion rate [66]. Therefore, pH variations can have a significant impact on the corrosion rate of stainless steel [1]. Similar results have been found that corrosion rate of stainless steels is affected by pH variations. For instance, a study by Samosir in 2017 demonstrated that the corrosion rate of stainless steel increased with a decrease in pH below 4 [67]. On the other hand, maintaining a neutral or higher pH can help reduce the corrosion rate of stainless steel [68]. Therefore, it is crucial to maintain optimal pH levels to minimize the corrosion rate of stainless steel and prolong its lifespan.

Fig. 6 also depicts the effect of pH variations on the hardness of SS 316L samples. It can be seen that the hardness of SS 316L decreased in a pH lower than 7 and higher than 7. The highest hardness value was found at neutral condition (pH=7) and the lowest hardness value was at acidic condition (pH=1). Hardness is an important characteristic of stainless steel, as it determines the material's resistance to wear and deformation. In a study conducted by Saefuloh et al. in 2020, the hardness of stainless steel 304 was measured in corrosive media, including sea water and 3.5% NaCl [65]. The results showed that the hardness of stainless steel decreased as the corrosion rate increased, indicating a negative correlation between hardness and corrosion rate. Additionally, maintaining a high level of hardness in stainless steel is crucial to ensure its durability and longevity in various applications. The pH level of the environment in which stainless steel is used can have a significant impact on its hardness. Research by Garcia-Cabezon et al. in 2022 found that precipitation hardening stainless steel (PH SS) manufactured by selective laser melting had varying levels of hardness depending on the pH level of the environment [69]. Similarly, Saefuloh's research in 2020 showed that the corrosion rate of stainless-steel duplex was affected by the pH level of the environment, but had no significant effect on the level of hardness [65]. However, research by Zai et al. in 2020 found that the high strength and hardness induced by precipitation hardening in PH stainless steels limits their workability [70]. Hence, it is important to balance pH control and hardness in stainless steel applications to ensure optimal performance. The hardness of stainless steel is a critical factor in its performance, and can be influenced by variations in pH levels. While maintaining a high level of hardness is important, it is also crucial to consider the effects of pH control on the material's corrosion resistance and workability. Further research is needed to fully understand the relationship between pH variations and the hardness of stainless steel, and to develop effective strategies for balancing these factors in various applications.

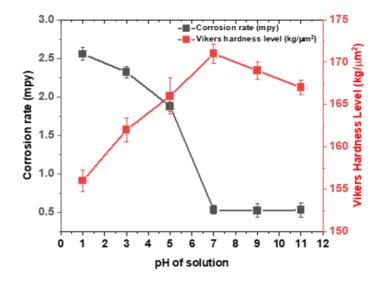


Fig. 6 Effect of pH of solutions on the corrosion rate and hardness of SS 316L

#### 4. Conclusion

In this study, the corrosion behaviors of stainless-steel type SS 316L under exposure immersions in salty and acidic environments using sodium chloride (NaCl) and sulfuric acid (H<sub>2</sub>SO<sub>4</sub>) solutions were investigated, by referring to a standard method of ASTM G31-72. Scanning electron microscopy (SEM) results revealed that the specimen in NaCl solution exhibited a darker color, and relatively similar shape of cracks are observed. In contrast, the specimen immersed in H<sub>2</sub>SO<sub>4</sub> was observed to have a large diagonal crack. It was found that the corrosion resistance of SS 316L immersed in NaCl solution is better because the corrosion rate value is < 1 mpy, which was 0.4260 mpy. This value is in the "excellent" category compared to the corrosion rate value of SS 316L immersed in H<sub>2</sub>SO<sub>4</sub> solution, which was 2.5141 mpy in the "excellent" category. This indicates that SS 316L is better placed in a salty environment than in acidic conditions. In the Vickers test, it was found that the value of each material increased not too significantly. The HV value increased by 6.34% for materials soaked in NaCl solution, and the value obtained from materials soaked in H<sub>2</sub>SO<sub>4</sub> solution increased by 1.94%. Moreover, the SS 316L sample from the NaCl solution has a slightly higher VHN than that of H<sub>2</sub>SO<sub>4</sub>. The defect-free corrosion surface can be attributed to enhancing the mechanical strength of the SS 316L sample in NaCl. The generated crystals on the surface of SS 316L can also provide higher mechanical strength. These results indicate that NaCl adds strength to the surface structure of the material compared to the  $H_2SO_4$  solution. Therefore, SS 316L itself is widely used for the manufacture of equipment that is in direct contact with salt solutions because the main function of SS 316L is to be able to withstand rust, one of which is caused by chloride. Finally, it can be concluded that SS 316L used in the marine industry will perform better in terms of resistance and hardness compared to the other industries that often intersect with acidic solutions. It is crucial to maintain optimal pH levels to minimize the corrosion rate of stainless steel and prolong its lifespan. Further research is needed to fully understand the relationship between pH variations and the hardness of stainless steel, and to develop effective strategies for balancing these factors in various applications.

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