Application of Pulsed Laser Fabrication in Localized Corrosion Research

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1. Introduction

Aluminum and its alloys have been known as light metals because they are used to reduce the weight of automobiles and components. Aluminum is the second most used and produced metal in the world nowadays. It is well known that one of the typical corrosion morphologies of aluminum alloys in chloride containing environments such as seawater is pitting corrosion. Many papers have been investigating pitting corrosion ((Ito et al., 1968), (Horibe et al., 1969), (Goto et al. 1970), (Blanc et al., 1997), (Kang et al., 2010)). Electrochemical techniques, such as model macro-pits (Itoi et al., 2003) and electrochemical noise analysis (Sakairi et al., 2005, 2006, 2007) have been applied to investigate pitting corrosion of aluminum alloys.

Details of the propagation of pitting corrosion (Fig. 1) are not fully understood, however, the aspect ratio of pits (pit depth/pit diameter) plays a very important role in the growth of corrosion pits (Toma et al., 1980). To understand this effect, an in-situ artificial pit fabrication technique with area selective dissolution measurements would be helpful. One such technique is pulsed laser fabrication, which uses focused pulsed Nd-YAG laser beam irradiation to remove material from the substrate, combined with anodizing. Some of the authors have reported on the electrochemical behavior of artificial pits fabricated on aluminum alloy ((Sakari et al., 2009), (Yanada et al., 2010)).

In this chapter, the results of the effect of aspect ratio on dissolution behavior of the artificial pits formed on aluminum alloys are explained, and the chapter also explains the rate of pit fabrication and how to activate only the bottom surface of the formed pits.

2. Artificial pit fabrication by pulsed laser

2.1 Experimental

Sheet specimens of 2024 (15 × 20 × 2.0 mm) and 1050 (15 × 20 × 1.1 mm) aluminum alloys were used. Table 1 shows the chemical composition of the used aluminum alloys. Specimens were cleaned in doubly distilled water and an ethanol ultrasonic bath, and then polished

Fig. 1. Schematic representation of propagation of pitting corrosion in chloride ion containing solutions.

	Si	Fe	Cu	Mn	Mg	Zn	other	Al
2024	0.06	0.20	4.78	0.59	1.43	0.07	0.26	balance
1050	0.08	0.27	0.03	-	-	-	0.05	balance

Table 1. Chemical compositions of used aluminum alloys.

chemically in 0.1 kmol m^{-3} NaOH for 1800 s. A protective film is required to investigate the electrochemical reactions at only the laser beam irradiated area, and porous type anodic oxide films were formed at a constant current density of 100 A m^{-2} in 0.22 kmol m^{-3} $\text{C}_2\text{H}_2\text{O}_4$ at 293 K for 1800 s. Anodized specimens were dipped in boiling doubly distilled water for 900 s (pore sealing) to improve the protective performance of the formed anodic oxide films.

Specimens with protective films were irradiated by a focused Nd-YAG laser beam (Septra Physics GCR-130, wave duration 8 ns, frequency 10 s^{-1} , wave length 532 nm) for $t_i = 0$ to 30 s while immersed in 0.5 kmol m^{-3} H_3BO_3 - 0.05 kmol m^{-3} $\text{Na}_2\text{B}_4\text{O}_7$ (Borate). The laser beam power was adjusted to 30 mW in front of the lens. Fig. 2 shows a schematic outline of the laser irradiation and electrochemical measurement apparatus used in this chapter.

Surface and pit size observations: Specimen surfaces after the experiments were examined by an optical microscope, confocal scanning laser microscope (CSLM; 1SA21, LASERTEC Co.), and a scanning electron microscope (SEM; TM 1000, Hitachi Co.). The formed artificial pit depths and geometry were measured with the analysis function of the CSLM and X-ray Computed Tomography (X-ray CT; ELE SCAN mini, NS-ELEX Co. Ltd.).

Fig. 2. Schematic representation of the laser irradiation and electrochemical measurement apparatus.

2.2 Results and discussion

2.2.1 Kinetics of pit fabrication

Artificial pit depth and morphology changes with continuous laser beam irradiation time were investigated in Borate. Fig. 3 shows SEM surface images of laser beam irradiated 2024 aluminum alloy specimens at $t_i = 0.1$ to 150 s. The anodic oxide film and aluminum alloy substrate can be removed at the irradiated area, and the shape of the area where oxide film was removed is almost circular. The center of the oxide film removed area is bright initially and then becomes dark with t_i , and it becomes larger with t_i .

Fig. 3. SEM surface images of fabricated artificial pits on 2024 aluminum alloy at different laser beam irradiation times.

Figure 4 shows X-ray CT vertical sectioning images of fabricated artificial pits on 2024 aluminum alloy. Fig. 5 shows horizontal section images of a $t_i = 150$ s pit and a schematic representation of the section positions. From Fig. 4, the depth of a fabricated pit becomes deeper with longer t_i . From the horizontal sectional images in Fig. 5, the shape of fabricated artificial pits are almost completely circular from the top to the bottom.

Fig. 4. X-ray computed tomography (X-ray CT) vertical section images of fabricated artificial pits on 2024 aluminum alloy.

Fig. 5. X-ray CT horizontal section images of pits fabricated on 2024 aluminum alloy ($t_i = 150$ s) and schematic representation of section positions.

Fig. 6. CSLM depth profiles of fabricated artificial pits in 2024 aluminum alloy.

Figure 6 shows CSLM depth profile of fabricated pits with t_i . The CSLM depth profile also shows that the center area of the laser beam irradiated area is deeper than the other areas. The pit becomes deeper with t_i , however, the cross-sectional shape is independent of t_i .

Figure 7 shows changes in diameter (width) of artificial pits fabricated in both the 1050 and 2024 aluminum alloys with t_i . The pit diameter increases sharply at $t_i < 1$ s and the slope of the diameter change curve becomes flatter with longer t_i . The value of the diameter of 1050 aluminum alloy is about 20% larger than that of 2024 aluminum alloy. The laser beam used here has a Gaussian energy distribution and aluminum metal changes to gas or plasma only at the center of the irradiated area. However, the outer rim of the laser beam has sufficient energy to melt the aluminum substrate. This melted metal was ejected or flows by the effect of the evaporated gas or formed plasma (Fig. 8). If the irradiating conditions do not change during the experiments, then, after some time, the size of the melted area would not change with t_i .

Figure 9 shows the increases in depth of artificial pits fabricated on both the 1050 and 2024 aluminum alloys with t_i . The pit depth increases sharply at $t_i < 1$ s and the slope of the depth change curve becomes flatter with t_i . The specimen did not move during the laser beam irradiation, and therefore the distance between lens and irradiated surface (bottom of the pit) becomes longer with t_i . This distance change causes a decrease in the mean beam energy available for pit fabrication. This is a reason why the slope of the pit depth change curve becomes flatter with t_i . The pit formation rate of the 1050 aluminum alloy is about twice that of the 2024 aluminum alloy.

Fig. 7. Changes in the diameters of fabricated artificial pits on 1050 and 2024 aluminum alloys as a function of. irradiation time.

Fig. 8. Schematic representation of pit fabrication mechanism by continuous laser beam irradiation.

Fig. 9. Changes in the depths of fabricated artificial pits as a function of irradiation time.

These results shown here clearly substantiate that continuous focused pulse laser beam irradiation makes it possible to fabricate artificial pits on aluminum alloy in solutions, and the shape of the pits appear to be bell shaped. The detailed mechanism of pit fabrication is explained in section 2.2.2, but a possible mechanism is laser ablation.

Figure 10 shows changes in aspect ratio with t_i for both the 1050 and 2024 aluminum alloys. The aspect ratio of the formed pits on both aluminum alloys increases with t_i and the aspect ratio of both aluminum alloys at the same t_i are similar. This result shows that the proposed technique here makes it possible to fabricate artificial pits with different aspect ratios (0.13 - 2.3) on anodized aluminum in solutions.

Fig. 10. Changes in the aspect ratios of fabricated artificial pits as a function of irradiation time.

2.2.2 Pit fabrication mechanism

The detailed explanation of laser ablation to remove oxide film or metals is shown in the literature (Sakairi et al., 2007).

Anodic oxide films formed on aluminum alloys are almost completely transparent at the laser frequency of 532nm used here. As continuous irradiation, oxide films are removed after several irradiation pulses by the laser beam. These situations indicate that almost all of the irradiated laser light energy reaches the metal-oxide interface or metal surface. It is not certain that the reflectivity of high energy density light is the same as low energy density beams, however, the reported reflectivity value of 0.82 at 532 nm (Waver, 1991-1992) is used to estimate the adsorbed power density here. The adsorbed power density in this experimental condition, with the wave duration 8 ns, frequency 10 s^{-1} , irradiated diameter 300 μm , and $P = 3.0\text{ mJ}$ (30mW/10 Hz) becomes about 10^{12} W/m^2 . According to the literature (Ready, 1971), the approximate expression of the minimum laser power density for ablation of aluminum ($r = 2700\text{ kg m}^{-3}$, $L = 10778\text{ kJ kg}^{-1}$, $k = 1.0 \times 10^{-4}\text{ m}^2\text{ s}^{-1}$) is about $0.7 \times 10^{12}\text{ W/m}^2$. The value of the adsorbed power density in the present investigation is larger than that of ablation of aluminum. This suggests that laser ablation takes place beneath the area where the laser was irradiated. The ablation of metal produces pressure at the film/substrate or solution/substrate interfaces immediately after the irradiation. The pressure of laser ablation is simply calculated by using the laser power density for ablation, the specific thermal capacity of the aluminum, the initial, and the vaporization temperatures of the aluminum (Scruby, 1990). The estimated value is about 10^8 Pa . The deformation pressures of micro-filters made of porous type anodic oxide films with 45 μm thickness is about $2 \times 10^8\text{ Pa}$, and the pressures are proportional to the film thickness (Hoshino et al., 1997). The pressure estimated here is almost same as the deformation pressure of the thick porous type anodic oxide film. It may be concluded that the anodic oxide film and aluminum substrate can be destroyed and removed by the high pressure at the interface produced by the laser ablation of the aluminum substrate itself.

3. Corrosion behavior in formed artificial pits

3.1 Experimental

Borate with 0 to 0.01 kmol m^{-3} NaCl was used as electrolyte for the corrosion tests. An Ag/AgCl sat. KCl electrode was used as the reference electrode, and Pt plate (2x2 cm) was used as the counter electrode.

Polarization curves of chemically polished 2024 alloy specimens (un-anodized) were measured to determine the optimum applied potential and Cl^- concentration for investigation of the effect of the aspect ratio on the current transient in the artificial pits. In this experiment, the potential was swept at the constant rate of 0.83 mV/s from the rest potential to the anodic potential direction.

Two different types of electrochemical corrosion tests were carried out after fabrication of artificial pits with different aspect ratios on 2024 alloy, namely with the current transients at constant potential and with the rest potential changing.

Current transients: Artificial pits with two different depths formed by $t_i = 1\text{ s}$ and 120 s were formed in Borate with 0.01 kmol m^{-3} NaCl, then a constant potential of -300 mV was applied. The current was measured to establish that no further dissolution or passivation was occurring in the pits, and after that one more pulse of laser light was applied to activate the bottom of the pits. The current transients after the activation were measured with a digital oscilloscope (Yokogawa Electric Co., DL708E).

Rest potential: The artificial pits with two different depths formed at $t_i = 1$ s to 120 s were formed in Borate with 0.001 to 0.01 kmol m^{-3} NaCl, then the bottom of the pits were activated by one more pulse of laser light. After the activation, the rest potential was measured with the digital oscilloscope.

One of the authors have reported on the effect of an aperture on the oxide film removed area (Sakairi et al., 1998). Using an aperture makes it possible to narrow the irradiated area at the focus point. Therefore, to activate only the bottom of the pits, an aperture placed in front of the lens was used. In the case of specimens with deep pits, $t_i = 10$ s to 120 s, the distance between lens and pit bottom was adjusted to focus the point of the used lens by using an XYZ stage (Fig. 1).

After the experiments, the specimen surfaces were examined by an optical microscope.

3.2 Results and discussion

3.2.1 Polarization results

The potentiodynamic polarization curves of chemically polished 2024 aluminum alloys were measured in Borate with different concentrations of NaCl (Fig. 11) to determine the Cl^- concentration and applied potential for electrochemical corrosion tests of the artificial pits. The rest potential shifts to lower potentials with higher NaCl concentrations. At the low Cl^- concentration of 0.001 kmol m^{-3} , the relationship between current density and potential shows no changes due to the added Cl^- . No current fluctuations are observed reveals suggesting that no localized corrosion is taking place. However, at higher Cl^- concentrations ($>0.002 \text{ kmol m}^{-2}$), the current shows sudden increases at the start of the polarization with further current fluctuations. This result at the higher Cl^- concentrations shows that when

Fig. 11. Potentiodynamic anodic polarization curves of chemically polished 2024 aluminum alloy in 0.5 kmol m^{-3} H_3BO_3 - 0.05 kmol m^{-3} $\text{Na}_2\text{B}_4\text{O}_7$ with 0 to 0.01 kmol m^{-3} NaCl.

aluminum substrate becomes exposed to the solution by laser beam irradiation, pitting corrosion tends to occur even at the open circuit condition. From these results, Borate with 0.01 kmol m^{-3} NaCl and an applied potential of -0.3 V was chosen for the electrochemical measurements of the artificial pits formed on the 2025 aluminum alloy. To investigate the effect of Cl^- concentration on pit propagation at open circuit condition (no potential applied), Borate with 0.001 and 0.01 kmol m^{-3} NaCl were chosen.

3.2.2 Current measurements

Figure 12 shows changes in the current of the pits formed on 2024 aluminum alloy with $t_i = 1 \text{ s}$ and 120 s after activation by 1 pulse of laser beam irradiation at -0.3 V in Borate with 0.01 kmol m^{-3} NaCl. After the laser beam irradiation, the current increased instantaneously through a maximum, then decreases with time in both pit conditions. After the test, white corrosion products can be seen on the specimen surfaces at the pit (Fig. 13).

Figure 14 shows changes in rest potential during and after pit formation on 2024 aluminum alloy in Borate with 0.01 kmol m^{-3} NaCl. Results with specimens without pits are also shown in the figure to evaluate the protectiveness of the anodic oxide film. The rest potential in both irradiated specimens show negative values during pit fabrication, and then there are increases after the pit fabrication. Fluctuations which relate to localized corrosion events are also observed in the rest potential changes. There are no very large potential fluctuations in the results for the anodized specimens here, indicating that anodic oxide film has good corrosion resistance for long times and that the measured potential fluctuations are related to events inside the formed pits.

Figure 15 shows changes in the rest potential during and after pit formation on 2024 aluminum alloy in Borate with $0.001 \text{ kmol m}^{-3}$ NaCl. The changes in the rest potential at each t_i are very similar to those in Fig. 14, with no significant fluctuations observed. This means that the formed pits are repassivated after some time of pit formation, because of the low Cl^- concentration.

Fig. 12. Changes in the current of the pit formed on 2024 aluminum alloy after activation by one pulse of laser beam irradiation at -0.3 V in $0.5 \text{ kmol m}^{-3} \text{ H}_3\text{BO}_3 - 0.05 \text{ kmol m}^{-3} \text{ Na}_2\text{B}_4\text{O}_7$ with 0.01 kmol m^{-3} NaCl.

Fig. 13. Optical images of specimen surfaces after the test in Fig. 12.

Figure 16 shows rest potentials 2400 s after the finish of the pit fabrication, E_{2400} , in Fig. 15 as a function of t_i . The E_{2400} decreases with increasing t_i , suggesting pit depth or aspect ratio influence on the protective thin oxide film formation or differences in repassivation kinetics. Figure 17 shows different stages of rest potential changes by polarization curves at each step of the pit formation process. During the pit formation, the aluminum substrate is frequently activated causing increases in anodic currents and decreases in the rest potential. After the pit formation, the aluminum substrate is not further activated by the laser irradiation, and there is repassivation or further localized corrosion progresses in higher concentrations Cl^- solutions. If the surface repassivates, the anodic current decreases causing rest potential increases.

Fig. 14. Changes in rest potential during and after pit formation in $0.5 \text{ kmol m}^{-3} \text{ H}_3\text{BO}_3 - 0.05 \text{ kmol m}^{-3} \text{ Na}_2\text{B}_4\text{O}_7$ with $0.01 \text{ kmol m}^{-3} \text{ NaCl}$. The rest potential of specimens without pits is also shown in the figure.

Fig. 15. Changes in rest potential during and after pit formation on 2024 aluminum alloy in $0.5 \text{ kmol m}^{-3} \text{ H}_3\text{BO}_3$ - $0.05 \text{ kmol m}^{-3} \text{ Na}_2\text{B}_4\text{O}_7$ with $0.001 \text{ kmol m}^{-3} \text{ NaCl}$

To investigate the effect of pit aspect ratio on the repassivation kinetics, the bottom of the pit was re-activated by one pulse of laser beam irradiation. The re-activation was carried out 2400 s after completion of the pit formation. Fig. 18 shows the changes in the rest potential after this re-activation of 2024 aluminum alloy in Borate with $0.001 \text{ kmol m}^{-3} \text{ NaCl}$. The potential changes to the negative direction in all specimens, shows a minimum value, and then shifts to the positive direction. This potential shift to the positive direction suggests repassivation of the re-activated surface at the bottom of the pit.

Figure 19 shows optical images after the re-activation tests. No corrosion products are observed in either the $t_i = 1 \text{ s}$ or 120 s specimens. This result suggest that repassivation took place because of the low concentration of Cl^- , in good agreement with the potential changes in Fig. 18.

The lowest rest potential after the re-activation as a function of aspect ratio is shown in Fig. 20. Low aspect ratio samples show the lowest reached potential, while higher aspect ratio specimens show very similar values.

To clarify the effect of the aspect ratio on the repassivation kinetics, a repassivation ratio concept is introduced. The repassivation ratio, r_p , is explained as follows

$$r_p = \frac{\Delta E_p}{\Delta E_{ac}}$$

$$\Delta E_{ac} = E_{2400} - E_{ac}$$

$$\Delta E_p = E_p - E_{ac}$$

where E_{2400} is the rest potential 2400 s after pit formation, E_{ac} is the lowest rest potential after re-activation, and E_p is the average value of the rest potential around 0.01 s or 10 s after the re-activation. Therefore, a high value of r_p indicates that repassivation has progressed.

Changes in r_p at $t_c = 0.01 \text{ s}$ (Fig. 21) and 10 s (Fig. 22) with the aspect ratio of the pit in Borate with $0.001 \text{ kmol m}^{-3} \text{ NaCl}$ were established. The r_p of the 1050 aluminum alloy is also shown

Fig. 16. Rest potential at 2400 s after the finish of the pit fabrication in Fig. 15 as a function of the aspect ratio of the formed pits.

Fig. 17. Schematic representation of rest potential changes by polarization curves at each step of the pit formation process.

Fig. 18. Changes in rest potential after the re-activation of 2024 aluminum alloy in $0.5 \text{ kmol m}^{-3} \text{ H}_3\text{BO}_3$ - $0.05 \text{ kmol m}^{-3} \text{ Na}_2\text{B}_4\text{O}_7$ with $0.001 \text{ kmol m}^{-3} \text{ NaCl}$. Re-activation was carried out 2400 s after the pit formation.

Fig. 19. Optical images after the re-activation tests in Fig. 18.

in the figures. It is clearly shown that r_p at 0.01 s decreases with the aspect ratio of the pit (Fig. 21) while at 10 s it increases with aspect ratio (Fig. 22). At both t_c , the r_p of the 2024 aluminum alloy is higher than that of the 1050 aluminum alloy. These results suggest that the repassivation rate of the 2024 aluminum alloy is faster than that of the 1050 aluminum alloy in low NaCl containing solutions.

Figure 23 shows a schematic representation of the situation after activation at the bottom of a pit, (a) transfer of oxygen and (b) selection of anodic and cathodic sites and transfer of hydrogen. With 2024 aluminum alloy, a number of copper rich intermetallics are present in the substrate. In the pit as formed by laser irradiation, these intermetallics may be exposed to the solution and act as cathodic reaction sites during immersion corrosion tests.

In the pit here, the main cathodic reaction is oxygen reduction because the solution pH is close to neutral and no nitrogen or argon gas was bubbled into the solution. The exposed area of the intermetallics in the pit wall may also increase with aspect ratio and cause the increase in cathodic partial current, i_c , in Fig. 17. Here, as the anodic partial current, the dissolution of aluminum is increased by the re-activation, and there are no large rest potential changes. This also means an acceleration in the rate of passivation in low NaCl containing solutions. This may be concluded to be the reason why the r_p of the 2024 aluminum alloy is larger than that of the 1050 aluminum alloy in Figs 21 and 22.

Fig. 20. Lowest rest potentials after the re-activation as a function of aspect ratio in $0.5 \text{ kmol m}^{-3} \text{ H}_3\text{BO}_3$ - $0.05 \text{ kmol m}^{-3} \text{ Na}_2\text{B}_4\text{O}_7$ with $0.001 \text{ kmol m}^{-3} \text{ NaCl}$.

Fig. 21. Changes in repassivation ratios at 0.01 s with different aspect ratios of pits in $0.5 \text{ kmol m}^{-3} \text{ H}_3\text{BO}_3$ - $0.05 \text{ kmol m}^{-3} \text{ Na}_2\text{B}_4\text{O}_7$ with $0.001 \text{ kmol m}^{-3} \text{ NaCl}$.

Fig. 22. Changes in repassivation ratios at 10 s with different aspect ratios of pits in $0.5 \text{ kmol m}^{-3} \text{ H}_3\text{BO}_3 - 0.05 \text{ kmol m}^{-3} \text{ Na}_2\text{B}_4\text{O}_7$ with $0.001 \text{ kmol m}^{-3} \text{ NaCl}$.

After some time, dissolved oxygen in the solution inside the pit may be consumed and oxygen diffuse from the bulk solution to the pit. In the high aspect ratio pit, the distribution of oxygen concentrations becomes dominant dividing the cathodic reaction (near the pit mouth) and anodic reaction areas (near the pit bottom) in the pit (Fig. 23 (b)). This means that the dissolution rate of aluminum at the bottom also increases. The dissolved Al^{3+} reacts with water to form H^+ and lowers the pH locally. The higher aspect ratio makes it difficult to dilute the H^+ ions at the pit bottom, and this is a possible reason why the r_p at 0.01 s decreases with increasing aspect ratio. At $t_c = 10 \text{ s}$, the pH at the bottom of the pit may increase because of buffer reactions of the Borate and diffusion of H^+ ion into the bulk solution. The cathodic reaction rate of the high aspect ratio pit is still faster than that of the low aspect ratio pit. This fast cathodic reaction may make it easier to achieve repassivation at the bottom of the pit.

4. Summary

In this chapter, the application of a new in-situ artificial micro-pit formation method with an area selective electrochemical measurement technique was explained. The technique showed here uses focused pulsed Nd-YAG laser irradiation and anodizing. This technique was applied to investigate the effect of the geometry (aspect ratio) of artificially formed pits on the localized corrosion behavior of the formed artificial pits in aluminum alloys. The following conclusions may be drawn.

1. By controlling the laser irradiation time it becomes possible to form artificial micro-pits with different aspect ratios. An aspect ratio of about 2 is obtained by 120 s of laser irradiation.
2. The pit formation rate of the 2024 Al alloy is about four times slower than that of 1050 Al alloy.
3. The rest potential of the pits at 2400 s after completion of pit formation, E_{2400} , becomes lower with increasing aspect ratio.
4. The repassivation ratio at 0.01 s after activation becomes lower with increasing aspect ratio.

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The book starts with basic overview of physical phenomena on laser-matter interaction. Then it is followed by presentation of a number of laser applications in the nano-particles and thin films production, materials examination for industry, biological applications (in-vitro fertilization, tissue ablation) and long-range detection issues by LIDARs.

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