

Effect of Thickness of Water Film on Corrosion Behavior of Carbon Steel in Marine Atmosphere

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The relative humidity, RH , in equilibrium with solutions for various concentrations of sea salt could be calculated by using available thermodynamic data [1].

To confirm the relationships between chemical composition of water film and RH in actual environment, amount of absorbed moisture was measured. Stainless steel sheets and QCM, Quartz Crystal Microbalance, [2] were covered with given amount of simulated sea salt, W_s , which contained NaCl and $\text{MgCl}_2 \cdot 6\text{H}_2\text{O}$; $m[\text{Mg}^{2+}]/m[\text{Na}^+] = 0.11$ [3], and were exposed in the constant humidity chamber for 2 ~ 4 h. The amount of absorbed moisture, W_{ab} , was determined as weight change during the exposure.

Relationships between the ratios of (W_{ab}/W_s) and RH are shown in Fig.1. The value of (W_{ab}/W_s) is also given as follows [1];

$$(W_{ab}/W_s) = \{(1000s/m_2) - 6sM_w\} / M_t \quad (1)$$

$$M_t = M_1 + sM_2 + 6sM_w$$

where $s = m[\text{Mg}^{2+}]/m[\text{Na}^+] = 0.11$, m_2 is concentration of MgCl_2 , and M_1 , M_2 and M_w are molecular weights of NaCl , MgCl_2 and water, respectively. This calculated value of (W_{ab}/W_s) is also plotted in the figure, and it agreed well with measured data, when $W_s \geq 10^{-2} \text{g/m}^2$. Thus, d is in proportion to W_s when $W_s \geq 10^{-2} \text{g/m}^2$.

Relationships between thickness of water film, d , and RH under various W_s conditions are shown in Fig.2. The values of d were calculated as follows [1];

$$d = (sW_s/M_r r_s) \cdot \{(M_1 m_1/m_2) + M_2 + (1000/m_2)\} \quad (2)$$

where r_s is specific gravity of the solution, and m_1 is concentration of NaCl .

Carbon steel coupons and QCM sensors [4] were sprayed with diluted sea water and dried to deposit given amount of sea salt on them. Carbon steel coupons were exposed in the constant humidity chamber for 1 month. As for QCM sensors, time variations of them were measured in the constant humidity chamber for 2~3 h, and corrosion rate, $\Delta L/\Delta t$, was determined as;

$$(\Delta L/\Delta t) = (1/r_{\text{Fe}}) \cdot (M_{\text{Fe}}/M_{(\text{OH})_x}) \cdot (\Delta m/\Delta t) \quad (3)$$

where M_{Fe} and $M_{(\text{OH})_x}$ are molecular weights of Fe and $(\text{OH})_x$, r_{Fe} is density of Fe , $r_{\text{Fe}} = 7.86 \text{g/cm}^3$, and $(\Delta m/\Delta t)$ is mass change measured by QCM sensor. In this work, the primary corrosion product of Fe was considered to be $\text{Fe}(\text{OH})_{2.5}$ [4].

Exposure test results are shown in Fig. 3 in terms of corrosion rate, $CR[\text{mm/y}]$, and thickness of water film, d . The value of CR increases with increasing d when $d < 56 \mu\text{m}$. It reaches up to the maximum of $CR = 0.28 \text{mm/y}$ at $d = 56 \mu\text{m}$, and it takes the constant value of $CR = 0.16 \text{mm/y}$ when $d \geq 170 \mu\text{m}$.

References

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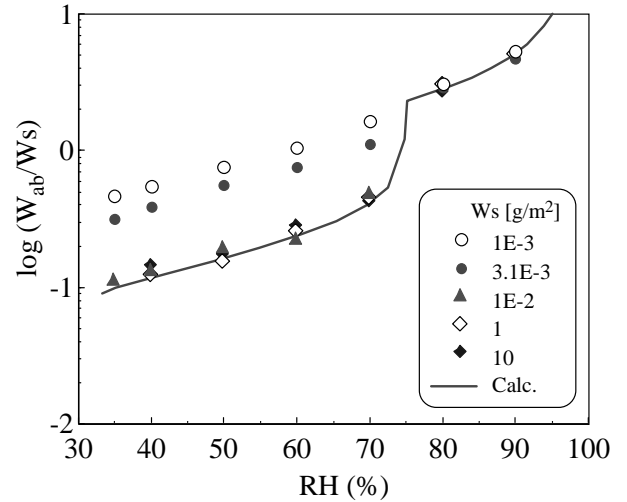


Fig. 1 Relationship between (W_{ab}/W_s) and RH .

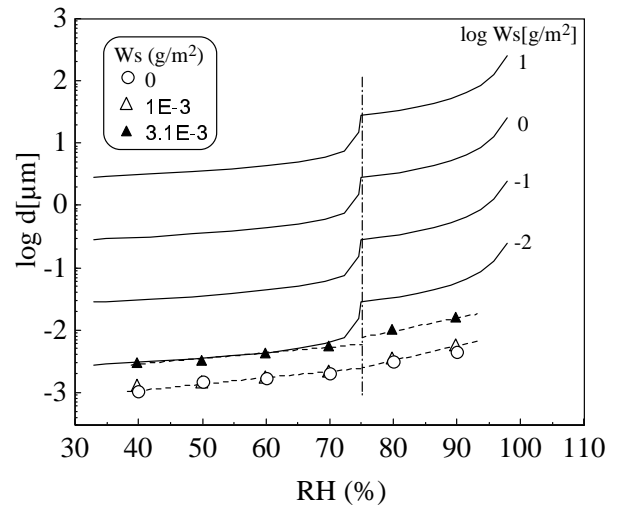


Fig.2 Relationship between d and RH under various W_s conditions.

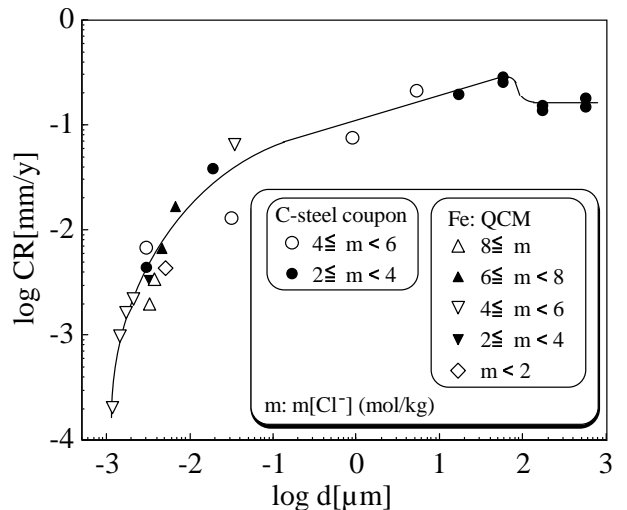


Fig.3 Effect of d on CR under various conditions.