Corrosion Probes for Fireside Monitoring in Coal-Fired Boilers

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ABSTRACT

Corrosion probes are being developed and combined with an existing measurement technology to provide a tool for assessing the extent of corrosion of metallic materials on the fireside in coal-fired boilers. The successful development of this technology will provide power plant operators the ability to (1) accurately monitor metal loss in critical regions of the boiler, such as waterwalls, superheaters, and reheaters; and (2) use corrosion rates as process variables. In the former, corrosion data could be used to schedule maintenance periods and in the later, processes can be altered to decrease corrosion rates.

The research approach involves laboratory research in simulated environments that will lead to field tests of corrosion probes in coal-fired boilers. Laboratory research[1] has already shown that electrochemically-measured corrosion rates for ash-covered metals are similar to actual mass loss corrosion rates. Electrochemical tests conducted using a potentiostat show the corrosion reaction of ash-covered probes at 500ºC to be electrochemical in nature. Corrosion rates measured are similar to those from an automated corrosion monitoring system.

Tests of corrosion probes made with mild steel, 304L stainless steel (SS), and 316L SS sensors showed that corrosion of the sensors in a very aggressive incinerator ash was controlled by the ash and not by the alloy content. Corrosion rates in nitrogen atmospheres tended to decrease slowly with time. The addition of oxygen-containing gases, oxygen and carbon dioxide to nitrogen caused a more rapid decrease in corrosion rate, while the addition of water vapor increased the corrosion rate.

INTRODUCTION

The research presented here shows that electrochemical corrosion rate (ECR) probes are viable for use in fossil fuel energy conversion systems. ECR probes will allow Power Plant owners to be proactive in terms of protecting their most valuable piece of equipment, the Power Plant. These probes will be used to continuously monitor corrosion rates of key Power Plant components and will be able to associate changes in corrosion behavior with other process changes. Corrosion rate will become a process variable for Power Plants. ECR probes will also be used to track corrosivity of environments and corrosion losses of key components in order to more efficiently schedule downtime for maintenance activities.

A number of research efforts have been aimed at developing high temperature corrosion probes for various industries. The majority of the research has been based on the use of electrochemical noise (EN)[2-8] techniques. Others have considered the use of electrochemical impedance spectroscopy (EIS)[4-6], linear polarization resistance (LPR)[7], zero resistance ammetry (ZRA)[5], and electrical resistance (ER)[5]. However, only a limited effort has been made to quantify[2] the operation of corrosion rate probes. For these probes to be accepted routinely in the power generation industries, it will be necessary to determine if they accurately measure corrosion rate and the changes in corrosion rate that occur in environments of interest, if the sensor materials have an optimum composition for the intended exposure, and if the sensitivity or accuracy of the sensor changes with exposure time in fireside environments. Once this is established, electrochemical corrosion rate probes can be used extensively, allowing corrosion rate to become a process variable for power plant operators.

Most electrochemical corrosion rate measurement techniques measure a resistance that is representative of the rate of the corrosion reaction. This is true of the LPR, EN, and EIS techniques. These resistances are related to corrosion rate by the Stern-Geary linear approximation to the Butler-Volmer equation,

$$R_p = R_n = \frac{\Delta E}{\Delta i_{applied}} = \frac{\beta_a \beta_c}{2.303(i_{corr})(\beta_a + \beta_c)} = \frac{B}{(i_{corr})} \tag{1}$$

where R_p is a polarization resistance obtained from the LPR and EIS techniques, R_n is a reaction resistance obtained from the EN technique, ΔE is the incremental change in potential measured due to the incremental change in applied current density, $\Delta i_{applied}$, B is the Stern-Geary constant, β_a and β_c are the anodic and cathodic Tafel constants, respectively, and i_{corr} is the corrosion current density from which a corrosion rate may be calculated using Faraday's Law. The Stern-Geary constant (determined by the Tafel constants) is the only variable that is normally not measured, but commonly assumed to have a value of 0.020 to 0.030 V/decade. Because B is related to Tafel constants, it can be measured using either standard electrochemical polarization techniques or the harmonic distortion analysis (HDA) technique that is used in this report.

EXPERIMENTAL DETAILS

ECR PROBE CONSTRUCTION

Isothermal probes are exposed nominally to the same temperature as the environment. These can be made as single or multi-probes. Multi-probes allow for increased amount of data for each experiment, where different alloy/ash combinations can be exposed simultaneously in the same experiment. Air cooled probes contain a cooling chamber that allows the use of liquids or gases to adjust the temperature of the metallic sensors to a temperature of interest. Probes built and used to date were built for use in laboratory tube furnaces. Figure 1 shows an air-cooled probe. Probes intended for use in field tests will incorporate features of the laboratory probes but also will need to be more rugged to survive the actual industrial environment.

Air cooled probes and isothermal probes consist of three sensors of the same material and size with electrical leads welded to each sensor. Air-cooled probes also contain an air chamber, stainless steel tubes to carry air into and out of the chamber, a type K thermocouple at the outer surface of the probe, and a type K thermocouple within the air chamber. For all probes, the outer two sensors

Figure 1 – Photo of an air-cooled probe showing wire leads connected to sensors and thermocouples and two lengths of 1/8 in stainless steel tubing for air inlet and outlet

were used as the working and counter electrodes and the inner sensor was the reference electrode. The thermocouples are intended to give adjusted probe temperatures and a thermal gradient across the sensors. The air chamber is made by casting a cube of Styrofoam within the sensor adjacent to the back of the sensors and then removing that cube by dissolving it in acetone after the potting compound hardens. Ceramcast 586 has been used to pot all probes to date. Isothermal probes are identical to the air-cooled probe without the thermocouples, air chamber, and air tubes. Multi-probes are made by combining up to three isothermal probes together in series. Each probe has separate electrical leads to their respective sensors.

HARDWARE/SOFTWARE SYSTEMS

The hardware used to measure the majority of the corrosion rates from the ECR probes is the SmartCET® Corrosion Monitoring System. This system used three separate electrochemical techniques: linear polarization resistance (LPR), electrochemical noise (EN), and harmonic distortion analysis (HDA). All three techniques measure a

corrosion rate, but in addition, EN also measures a localized (pitting) corrosion factor that varies from 0 to 1, and HDA measures the Tafel (β_a and β_c) and Stern-Geary (B) constants. Corrosion rates and other variables are reported every 7 minutes and stored and displayed to a computer using FieldCET® software. All corrosion rates reported in this paper are taken from the LPR measurements modified using the measured B values from each experiment.

Electrochemical measurements, both similar to and different from those made with the SmartCET®, were made using a Princeton Applied Research (PAR) 273A potentiostat/galvanostat. Potentiodynamic polarization tests were conducted at 0.167 mV/s from -0.2 V vs open circuit potential (OCP) to +0.5 V vs OCP. The reference electrode was the inner sensor of the probe and was of the same material as the other two sensors. When pitting corrosion was suspected, the potential was reversed at +0.5 V vs OCP and scanned to -0.2V vs OCP. LPR measurements were made at 0.167 mV/s from -15 mV to +15 mV vs OCP and returning to –15 mV vs OCP. All electrochemical measurements were controlled using DC Corrware software.

RESEARCH APPARATUS

All experiments were conducted in a 3-zone tube furnace containing a 2 in diameter alumina tube. Each zone was controlled with a separate temperature controller. Set temperatures for each zone were determined for each test temperature using an external calibration thermocouple and taking measurements at 1 inch (2.5 cm) intervals. For example, settings of 490, 510, and 400ºC for zones 1, 2, and 3, respectively, gives an internal temperature of 500ºC with a flat profile over 12 inch (30.5 cm) of the 24 in heated zone. An alumina D-tube was inserted in the furnace tube to allow a platform for mass loss coupons. Measurement and control thermocouples were inserted in a 316 SS sheath that was inserted into the D-tube.

Gas flows were controlled using digital mass flow controllers that are controlled using Labview programs. Water vapor is added through an air-powered metering pump that pumped a specific quantity of water into a heated chamber through which the test gas flowed.

RESULTS AND DISCUSSION

ECR PROBE RESPONSE

A 500 h exposure test for an incinerator ash-coated 304 SS ECR probe in a $N_2/O_2/CO_2$ gaseous environment was conducted to show the effect of gas composition changes on probe response. The probe was heated to temperature in N_2 and allowed to remain in N_2 for about 165 hrs. During this time the corrosion rate remained high at 4-5 mm/y, decreasing slightly. After 165 hours, the gas composition was changed to $N_2/O_2/CO_2$ which caused the corrosion rate to begin a decrease to about 1 mm/y at 230 h. Here the gas was again switched to N_2 which again caused the corrosion rate to increase to 4-5 mm/y. After peaking at 4.8 mm/y, the corrosion rate started decreasing. The addition of CO_2 at 250 h continued the decrease in corrosion rate but with a change in slope. The addition of O_2 at 295 h again continued the decrease in corrosion rate with yet another change in slope. This experiment shows that that the ECR probe is responsive to environment changes and that both O_2 and CO_2 cause the corrosion rate of the ash-covered 304 SS to decrease as soon as those gases are present. Corrosion rates in N_2 decreased only after an incubation period.

EFFECT OF ASH COMPOSITION

Two different types of ash were used in this research. The first was an ash from the Lee #1 Municpal Incinerator supplied by Covanta Energy, Inc. The second was a coal ash described as AEP TIDD Ash derived from Illinois #8 coal, and supplied by EERC, the University of North Dakota Energy and Environmental Research Center. The composition of the two types of ash are: **Illinois No. 8 Coal Ash**: Al_2O_3 – 7.6%, CaO – 28.3%,K_2O – trace, TiO_2 – trace, MgO – 15.6%, Na_2O – trace, P_2O_5 – trace, Fe_2O_3 – 6.4, SiO_2 – 23.7, SO_3 – 15%**. Incinerator Ash**: Al – 3.63%, Mn – 631 ppm, As – 107 ppm, Ca – 9.97%, K – 3.69%, Pb – 2.7%, Ti – 0.514%, Mg – 0.734%, Na – 3.53%, Cr – 482 ppm, Ag – < 20 ppm, P – 0.213%, Cd – 883 ppm, Se – <50 ppm, Cl **–** 6.71%, Left on Ignition, LOI – 26.6%, C – 0.135%, N_2 – 0.171%, Fe – 5%, Si – 8%, S – 6.47
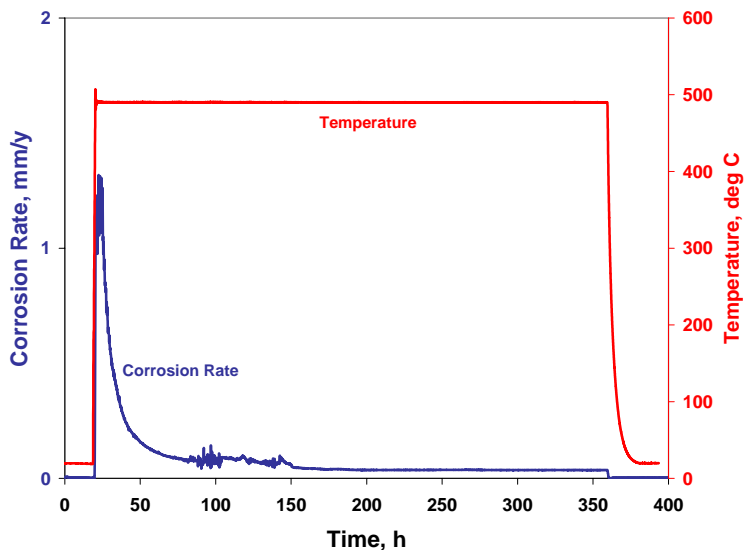
Figure 2 – Corrosion rate response of an incinerator- ash-covered 304 SS ECR Probe at 500ºC.

Incinerator ash was used for the majority of the experiments discussed in this paper. This ash shows high concentrations of corrosion-causing elements such as S, Cl, Pb, and K, all of which are able to form low melting point compounds and eutectic mixtures. The gas mixture consisted of 68 vol% N_2, 15 vol% H_2O, 9 vol% O_2, and 8 vol% CO_2 which is typical of waste combustion environments. Typical test periods were 100 to 500 hours and temperatures ranged from 450 to 600ºC. Figure 2 shows the typical response of a 304 SS ECR probe coated with incinerator ash. An expansion of the beginning of the experiment would show a small amount of corrosion when the ash/methanol slurry was applied to the probe before heat-up. Also visible would be the fact that the corrosion rate increase lags behind the temperature increase until approximately 400ºC. Corrosion rate was typically unstable for a period of time after the addition of the experiment environment. After that point the corrosion rate decreased over a period of 50 to100 h.

Coal ash was used for one experiment that ran for about 150 h. This ash notably lacks most of the corrosion-causing elements that are present in the incinerator ash. The gas mixture used, 69 vol% N_2, 15 vol% CO_2, 10 vol% H_2O, 5 vol% O_2, and 1 vol% SO_2 was typical of coal combustion environments. Figure 3 shows the typical response of a 304 SS ECR probe coated with coal ash. An expansion of the beginning of the experiment would show a small amount of corrosion when the ash/methanol slurry was applied to the probe before heat-up. Also visible would be the fact that the corrosion rate increase lags behind the temperature increase until approximately 400ºC. In coal ash, however, the corrosion rate is much lower at the beginning of the experiment than that shown in Figure 2. Another interesting phenomenon is that when the $N_2/CO_2/O_2/SO_2$ environment was added, the corrosion rate did not increase. The corrosion rate did finally start increasing 24 h later when water vapor was added to the environment. Corrosion rates did not decrease with time as shown in Figure 2 for the incinerator-ash-covered 304 SS ECR probe.
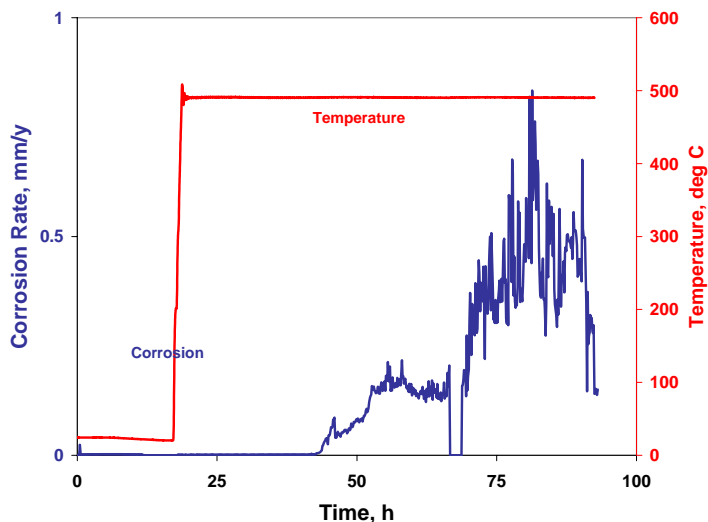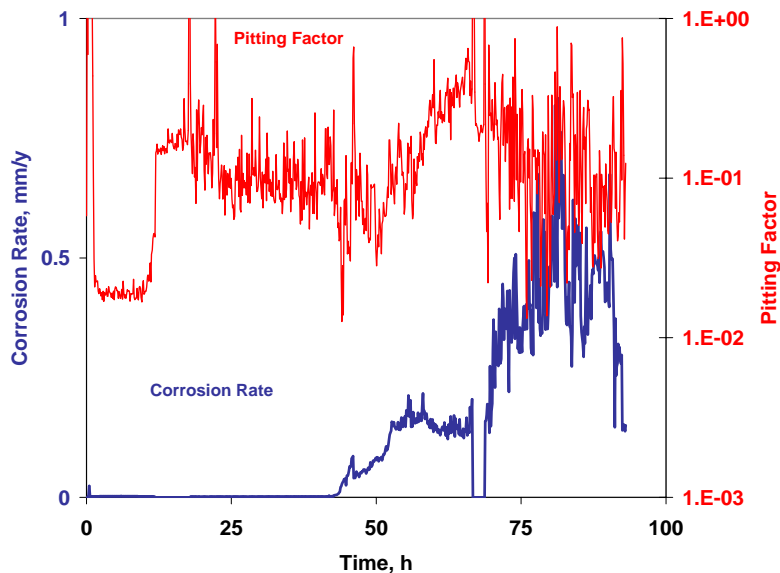

Figure 3 – Corrosion rate response of a coal-ash-covered 304 SS ECR probe at 500ºC.

The localized corrosion or pitting factor is an indication of the probability of pitting and it is measured with the EN technique. The pitting factor (PF) ranges from 0 to 1 but is best divided into logarithmic decades of 0.001 to 0.01 (no probability of localized corrosion), 0.01 to 0.1 (slight probability of localized corrosion), and 0.1 to 1 (high probability of localized corrosion). In all of the experiments conducted on ECR probes covered with incinerator ash,

Figure 4 – Corrosion rate and pitting factor as a function of time for a coal-ash-covered 304 SS ECR probe at 500ºC.

there was never an indication of the possibility of localized corrosion. This is despite the fact that the incinerator ash contained high levels of chlorides. Results were different for the coal-ash-covered ECR probe. Figure 4 shows the pitting factor to be in the highest range suggesting a high probability of pitting.

ELECTROCHEMISTRY OF THE CORROSION PROCESSES ON ASH-COVERED ECR PROBES

Electrochemical measurements were conducted using the PAR 273A potentiostat to verify the accuracy of the SmartCET® measurements, to show that other types of equipment can be used reliably to make similar measurements, and to attempt to understand the electrochemical nature of the corrosion reactions on the ash covered probes. These measurements were taken by interrupting and disconnecting the SmartCET® system, allowing the PAR 273A potentiostat to be connected. The potentiodynamic polarization behavior of an incinerator-ash-covered 304 SS ECR probe resulting from this measurement is shown in Figure 5. This curve is typical of an actively corroding metal and suggests that all or part of the corrosion reactions are electrochemical in nature. The linear part of the curve near the corrosion potential in Figure 5 was further analyzed to derive the polarization resistance, R_p. R_p was then converted to a corrosion rate for comparison to the SmartCET® LPR corrosion rate. Standard LPR tests were also conducted (but not shown) and the data converted to a corrosion rate. All of these measurements are compared in Table 1.

The measurements in Table 1 show a good agreement between the PAR and the SmartCET LPR corrosion rates. The potentiodynamic (PD) corrosion rates are actually close to the SmartCET LPR corrosion rates, but not quite as close as the PAR LPR corrosion rates. This is to be expected because of the different parameters used to make the PAR LPR and PAR PD measurements.

Electrochemical tests on a coal-ash-covered 304 SS ECR probe, shown in Figure 6, show a more complicated potentiodynamic polarization behavior. Part of
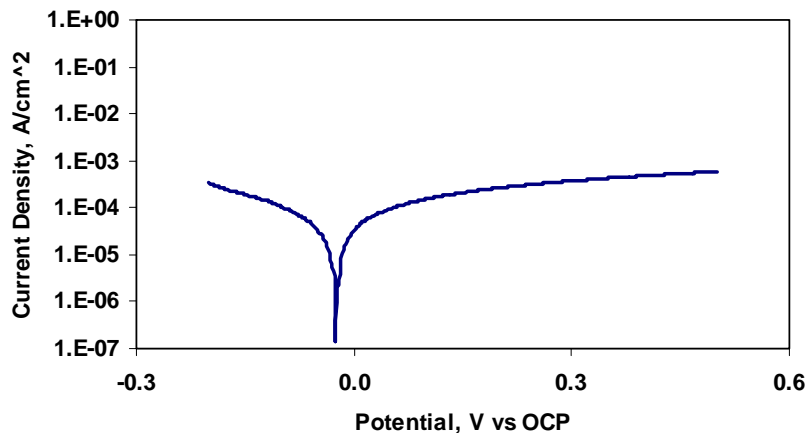

Figure 5 – Potentiodynamic polarization curve of an incinerator-ash-covered 304 SS ECR probe at 500ºC.

the reason for this is that the potential was reversed at 0.5 V vs OCP in order to determine if pitting was occurring. Another reason is that the more resistive ash caused the electrochemical behavior to be noisy, requiring smoothing to produce the final curve in Figure 6.

The polarization curve in Figure 6 suggests active-passive behavior, pitting breakdown (a pitting potential, E_p), and a repassivation potential (E_{rp}). If this proves to be reproducible in future experiments, it supports the pitting factor information reported by the SmartCET® unit. Microscopic examination of the probe will be done to verify if localized corrosion occurred.

QUANTITATIVE NATURE OF ECR PROBES

One of the key features of corrosion rate probes is that they produce a reading that is equivalent to the actual corrosion rate of the structure. This is the meaning of "Quantitative Nature." If this is not the case, then it is essential that the signal be proportional in a regular manner to the corrosion rate. This would make the measurements semi-quantitative and would require a calibration curve or factor to determine the actual corrosion rate. The approach to understanding this feature of ECR probes is to use mass loss coupons embedded in ash and exposed at the same time 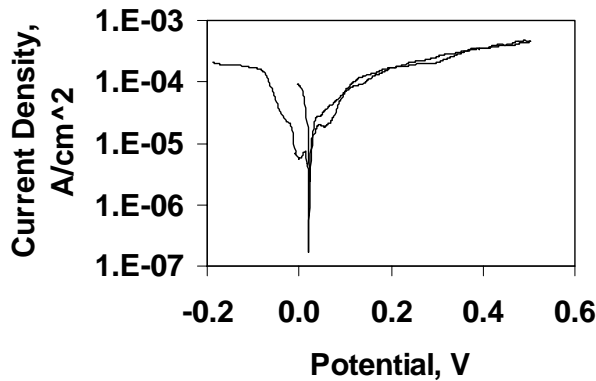and to the same conditions as the ECR probe. Figure 7 shows a picture of the apparatus that holds the ash embedded mass loss coupons in the experiment.

Table 1 – Comparison of SmartCET LPR, PAR LPR, and PAR PD (potentiodynamic) corrosion rates (mm/y)

SmartCET LPR	PAR LPR	PAR PD
0.0028	0.0008	na
1.83	1.63	na
1.79	1.65	1.52
0.66	0.60	0.45
0.44	0.41	0.30
1.44	1.28	0.91
0.15	0.14	0.14

Mass loss coupons were removed from the experiment, cleaned of all ash and corrosion deposits and weighed to determine weight loss due to corrosion. Corrosion penetration rates were then calculated using weight loss, exposed area, and exposure time. Electrochemical corrosion rates for the ECR probes were calculated either by: (1) converting corrosion rates to weight loss for each 7 min measurement period, summing the weight losses, and then converting to corrosion penetration rates as above, or (2) calculating an average corrosion rate over the exposure time period. Both techniques give nearly identical answers. Research[1] conducted previously has shown a close agreement between some alloy/environment combinations and

Figure 6 – Potentiodynamic polarization behavior of a coal-ash-covered 304 SS ECR probe at 500ºC.

poor agreement with others. Additional experiments are being conducted in order to resolve these differences

Another technique, optical profilometry, will be used in future tests to attempt further verification of the quantitative nature of ECR probes. Optical profilometry can measure and digitize the surface of materials. If there is a reference point, then it will be possible to measure and calculate the volume of a material lost due to corrosion. This can then be converted into a mass loss and then, as above, a corrosion rate. For the ECR probes, the reference point(s) are two pieces of alumina embedded at the top and bottom of the sensors and the surface to be analyzed is typically the center sensor, the unpolarized reference electrode. The reference electrode sensor is chosen because it most closely resembles the mass loss coupons.
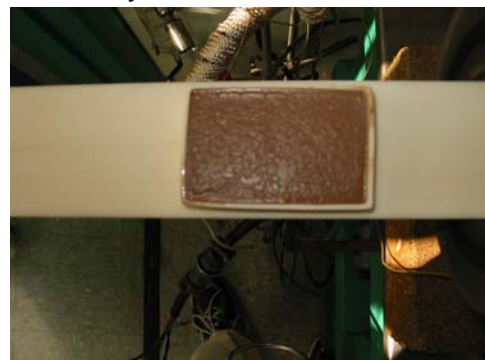

Figure 7 – Ceramic boat containing two ash-embedded mass loss coupons.

The counter and working electrode sensors are polarized every 7 minutes during the LPR and HDA measurements. Measurements taken before exposure are subtracted to provide a corrected corrosion volume loss.

SUMMARY/CONCLUSIONS

- ECR probes are responsive to changes in temperature and gas composition. Ash composition does affect this response.
- ECR probes appear to be quantitative based on initial measurements.
- Corrosion reactions occurring in both the incinerator and coal ashes appear to be at least partially electrochemical.
- Both EN and potentiodynamic polarization measurements predict that localized or pitting corrosion occurs for the coal-ash-covered 304 SS ECR probe.

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